

Technical University of Denmark



Green's function surface-integral method for nonlocal response of plasmonic nanowires in arbitrary dielectric environments

Yan, Wei; Mortensen, N. Asger; Wubs, Martijn

Published in:
Physical Review B

Link to article, DOI:
[10.1103/PhysRevB.88.155414](https://doi.org/10.1103/PhysRevB.88.155414)

Publication date:
2013

Document Version
Publisher's PDF, also known as Version of record

[Link back to DTU Orbit](#)

Citation (APA):

Yan, W., Mortensen, N. A., & Wubs, M. (2013). Green's function surface-integral method for nonlocal response of plasmonic nanowires in arbitrary dielectric environments. *Physical Review B*, 88(15), 155414. DOI: 10.1103/PhysRevB.88.155414

DTU Library

Technical Information Center of Denmark

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Green's function surface-integral method for nonlocal response of plasmonic nanowires in arbitrary dielectric environments

Wei Yan, N. Asger Mortensen, and Martijn Wubs*

*DTU Fotonik, Department of Photonics Engineering, Technical University of Denmark, DK-2800 Kongens Lyngby, Denmark
and Center for Nanostructured Graphene, Technical University of Denmark, DK-2800 Kongens Lyngby, Denmark*

(Received 26 July 2013; revised manuscript received 23 September 2013; published 14 October 2013)

We develop a nonlocal-response generalization to the Green's function surface-integral method (GSIM), also known as the boundary-element method. This numerically efficient method can accurately describe the linear hydrodynamic nonlocal response of arbitrarily shaped plasmonic nanowires in arbitrary dielectric backgrounds. All previous general-purpose methods for nonlocal response are bulk methods. We also expand the possible geometries to which the usual local-response GSIM can be applied, by showing how to regularize singularities that occur in the surface integrals when the nanoparticles touch a dielectric substrate. The same regularization works for nonlocal response. Furthermore, an effective theory is developed to explain the numerically observed nonlocal effects. The nonlocal frequency blueshift of a cylindrical nanowire in an inhomogeneous background generally increases as the nanowire radius and the longitudinal wave number become smaller, or when the effective background permittivity or the mode inhomogeneity increase. The inhomogeneity can be expressed in terms of an effective angular momentum of the surface-plasmon mode. We compare local and nonlocal response of freestanding nanowires, and of nanowires close to and on top of planar dielectric substrates. Especially for the latter geometry, considerable differences in extinction cross sections are found for local as compared to nonlocal response, similar to what is found for plasmonic dimer structures.

DOI: [10.1103/PhysRevB.88.155414](https://doi.org/10.1103/PhysRevB.88.155414)

PACS number(s): 42.70.Qs, 78.20.Bh, 71.45.Gm, 71.45.Lr

I. INTRODUCTION

Plasmonic (metallic) structures support a surface-plasmon (SP) resonance, i.e., coherent free-electron oscillations at the structure boundary.¹ With the SP resonance, electric fields can be localized to the deep subwavelength scale, and accordingly be enhanced dramatically. This leads to numerous applications, including signal transfer in nanoscale photonic circuits, few-molecule biosensing, and nonlinear phenomena.^{1,2}

For individual plasmonic nanostructures of size larger than typically 10 nm, it is accurate to describe metals with a local bulk refractive index,¹ as evidenced by numerous experiments. With recent progress in nanofabrication techniques, the sizes of individual plasmonic nanostructures can be controlled down to the deep nanoscale, below 10 nm, and their relative distances even below a single nanometer.³⁻⁹ This brings us into a regime where the foundation for the local bulk theory is challenged, since nonlocal response and the quantum wave nature of free electrons start to play a role.⁵⁻¹⁰

Here we adopt a direct and simple generalization of the local theory, namely the hydrodynamic Drude model (HDM). Besides the usual electromagnetic waves it also describes longitudinal waves in the free-electron plasma, which makes the response nonlocal.¹¹⁻¹⁵ The nonlocal effects that we study here are a consequence of the fact that light interacts with moving charges, and manifest themselves only in nanoplasmonic structures. In the HDM, it is predicted that the nonlocal response blueshifts the resonance peak, modifies the field enhancement, gives rise to new resonances above the plasma frequency, and drives the second-harmonic generation of the plasmonic structure.¹⁵⁻²⁹

The advantage of the HDM is that calculations are feasible for larger and more complex-shaped nanoplasmonic structures than for the higher level density-functional theory (DFT), for

example.^{10,30-33} The numerical method that we develop below expands the realm of structures that can be analyzed with the HDM. This allows a precise comparison with many experiments on nanometer-sized or nanometer-spaced particles, and with other models. It enables one to identify whether phenomena in nanoplasmonic experiments that go beyond the usual local-response approximation are hydrodynamic in nature, or even go beyond the HDM.

Before discussing the HDM and our numerical method in more detail, it is useful to discuss its range of validity. In DFT calculations for gold spheres of radius 0.74 nm, core-plasmon resonances (rather than single-particle excitations) are observed around the surface plasmon peak.³² For a twice larger radius these core plasmons are already much less pronounced than the surface-plasmon peak. The HDM does not describe core plasmons, and we choose to apply it only to particles of size 2 nm and larger. Apart from the hydrodynamical description we will furthermore assume that the static background density of the electron gas is a step function at the metal surface. We thereby neglect Friedel oscillations, electron spill-out, and quantum tunneling. The assumption is better for silver and gold^{23,29} than for sodium for example, where DFT calculations in a jellium model explain the redshift of plasmonic resonances as due to electron spill-out.^{10,30,34} Quantum tunneling only becomes important for plasmonic dimer structures with separations less than half a nanometer.^{8,35}

For a few regularly shaped freestanding structures, such as a slab, cylinder, and sphere, the linearized hydrodynamic scattering problem can be solved analytically, for example using Mie theory or transformation optics techniques.^{15-20,23,25-27} For realistic complex-shaped structures on substrates on the other hand, the hydrodynamic Drude response must be calculated numerically. Within the framework of the local-response bulk

theory, the numerical simulations of the optical properties of plasmonic structures are mature since well-developed methods exist, such as the finite-difference time-domain method (FDTD),^{36,37} finite-element method (FEM),³⁸ and the Green's function surface-integral method (GSIM), which is also known as the boundary-element method (BEM).^{39–45} By contrast, few accurate numerical methods exist for the hydrodynamical response.

Recently, the FEM was generalized to calculate the hydrodynamic Drude response of arbitrary-shaped plasmonic structures.^{22,24,29,46} The method was applied to nanowire dimers that show huge field enhancement,²² to corrugated surfaces used for surface-enhanced Raman spectroscopy,²⁴ and to calculate extinction properties of V grooves.⁴⁶ Very recently, the method was extended to calculate nonlocal effects in the waveguiding properties of plasmonic nanowires.^{47,48}

The FEM is a volume method, and nonlocal FEM in principle can handle both nanowires and three-dimensional structures. Yet it becomes numerically challenging for larger structures, especially for three-dimensional ones. This motivated us to develop a numerically efficient method. Already for local response it can sometimes be advantageous to turn to surface methods instead, where surfaces rather than scattering volumes need to be discretized. We started the present work anticipating that this advantage will only be greater for the nonlocal HDM, where a new length scale appears, namely the wavelength of the longitudinal waves. Since numerical meshes should be chosen considerably smaller than all length scales in the physical problem, in the HDM the meshing grid should be in the subwavelength scale of the longitudinal waves, which is below 1 nm.^{22,24} This suggests a larger relative advantage of surface methods for the hydrodynamic Drude theory.

Here we generalize the known Green's function surface-integral (equation) method^{39–45} for local-response theories to include nonlocal response as described by the hydrodynamic Drude model. Moreover, we generalize the applicability of the usual local-response GSIM to an experimentally relevant class of geometries, namely where nanostructures rest on dielectric interfaces. These “touching geometries” may give rise to additional singularities in the surface integrals. We show how to regularize these singularities. The regularization procedure is the same in the local and nonlocal GSIM, and enables a convergent numerical implementation of the method.

Here we apply our nonlocal GSIM to structures with a translational symmetry (2D structure with 3D illumination). In particular, we investigate the effects of the nonlocal response on plasmonic nanowires, first for nanowires in a free-space background, and then for nanowires above or resting on a dielectric substrate. In all our fully converged numerical calculations, the numerical grid size on the surface is in the subwavelength scale of the SP wave. We develop an approximate analytical theory for nonlocal blueshifts, and show its accuracy by comparison with our full GSIM numerics. We then use this theory to demonstrate how the strength of the nonlocal effects is determined by (i) the nanowire size r_0 , (ii) the longitudinal wave number k^L , (iii) the environmental permittivity ϵ_b , and (iv) the angular momentum of the SP mode l .

The remaining part of the article is organized as follows. Section II introduces the plasmonic nanowire structures under

study and their environment. In Sec. III, the hydrodynamic Drude model is introduced to describe the nonlocal response. We generalize the GSIM to describe nonlocal response of nanowire structures in Sec. IV. The numerical implementation of the nonlocal GSIM is discussed in Sec. V, which also includes our new singularity regularization procedure that allows the GSIM to be applied to nanostructures that touch a dielectric interface. In Sec. VI, we develop approximate analytical expressions for nonlocal blueshifts for nanowires in inhomogeneous dielectric environments. Testing the approximate theory is the red thread in our subsequent GSIM numerical simulations in Sec. VII, for nanowires without, above, and on dielectric substrates. We summarize, conclude, and discuss our method and results in Sec. VIII. Some detailed derivations are relegated to Appendices A–C.

II. NANOWIRE SYSTEM

We consider a nonmagnetic nanowire system, invariant in the \hat{z} direction and with arbitrary cross section in the \hat{x}, \hat{y} plane; see the sketch in Fig. 1. The system is divided into two regions: the plasmonic scatter region denoted as A , and the dielectric background denoted as B . Region A consists of an arbitrary number of isolated plasmonic nanowires A_i ($i = 1, 2, \dots, N$). The individual nanowires are each a homogenous medium as described by the hydrodynamic Drude model. The dielectric function of the background is ϵ_b , which we allow to be space dependent, and is assumed to be nonmagnetic. The boundary between A_i and B is called S_i . The outward-normal and tangential unit vectors at S_i are denoted as \hat{n}_i and \hat{l}_i obeying $\hat{n}_i \times \hat{l}_i = \hat{z}$.

When exciting the system electromagnetically, for example with an electric current source $\mathbf{J}_b \exp(-i\omega t)$ in the region B , then the translation invariance suggests decomposing \mathbf{J}_b into Fourier components along the wires,

$$\mathbf{J}_b(\boldsymbol{\rho}, z) = \int dk_z \tilde{\mathbf{J}}_b(\boldsymbol{\rho}, k_z) \exp(ik_z z), \quad (1)$$

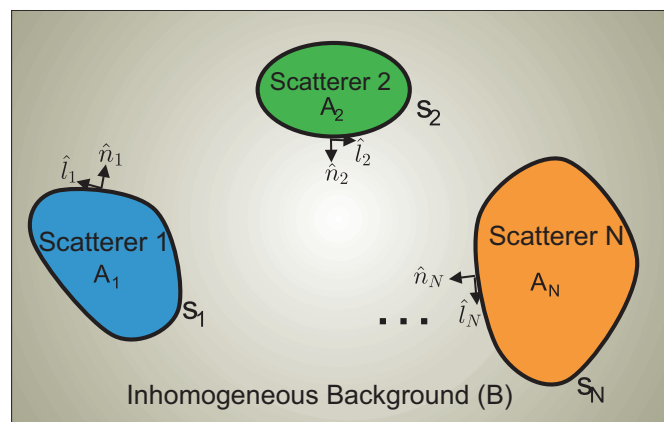


FIG. 1. (Color online) Illustration of the cross section of a nanowire system. N isolated nanowires denoted as A_i ($i = 1, 2, \dots, N$) placed in an arbitrary inhomogeneous background denoted as B . The surface between the background and each nanowire scatterer is denoted as S_i . The \hat{n}_i and \hat{l}_i denote the unit vectors normal and tangential to S_i , respectively.

where $\boldsymbol{\rho}$ represents (x, y) . The interaction between \mathbf{J}_b and the system is equivalent to a linear superposition of the subinteractions between the $\tilde{\mathbf{J}}_b$ and the system. Each subinteraction is a 2D problem in the $\hat{x}\text{-}\hat{y}$ plane with⁴⁹

$$(\nabla_{\boldsymbol{\rho}} + ik_z \hat{z}) \times \mathbf{E}(\boldsymbol{\rho}) = i\omega\mu_0 \mathbf{H}(\boldsymbol{\rho}), \quad (2a)$$

$$(\nabla_{\boldsymbol{\rho}} + ik_z \hat{z}) \times \mathbf{H}(\boldsymbol{\rho}) = -i\omega \mathbf{D}(\boldsymbol{\rho}) + \tilde{\mathbf{J}}_b(\boldsymbol{\rho}, k_z), \quad (2b)$$

with $\nabla_{\boldsymbol{\rho}}$ defined as $\hat{x}\partial_x + \hat{y}\partial_y$.

III. HYDRODYNAMIC DRUDE MODEL

Plasmonic nanowires are of special interest owing to their ability to support SP resonances. We use the hydrodynamic Drude model (HDM) to describe the dynamics of the free-electron gas.^{6,11–15} In the HDM, the electrons are collectively described by a density $n(\mathbf{r}, t)$ and velocity $\mathbf{v}(\mathbf{r}, t)$. The equation of motion is

$$m_e \left[\frac{\partial \mathbf{v}}{\partial t} + \mathbf{v} \cdot \nabla \mathbf{v} \right] = -m_e \gamma \mathbf{v} - \frac{\nabla p_{\text{deg}}}{n} + e(\mathbf{E} + \mathbf{v} \times \mathbf{B}), \quad (3)$$

where p_{deg} is the pressure from the ground-state energy of the degenerate quantum Fermi gas, and γ is the damping frequency. We describe the dynamics of the electron gas in the Thomas-Fermi approximation, where the pressure is given by $p_{\text{deg}} = \hbar^2 (3\pi^2)^{2/3} n^{5/3} / (5m_e)$. We thereby neglect exchange interactions and further many-body effects.³¹ Using the charge-conservation equation $-\partial n / \partial t = \nabla \cdot n\mathbf{v}$, we linearize Eq. (3), and obtain the constitutive relation of the free-electron gas

$$\frac{\beta^2}{\omega^2 + i\omega\gamma} \nabla \nabla \cdot \mathbf{P}_f(\mathbf{r}) + \mathbf{P}_f(\mathbf{r}) = -\epsilon_0 \frac{\omega_p^2}{\omega^2 + i\omega\gamma} \mathbf{E}(\mathbf{r}), \quad (4)$$

where ω_p represents the plasma frequency, and $\beta = \sqrt{3/5} v_F$ for $\omega \gg \gamma$ with v_F the Fermi velocity. The operator $\nabla \nabla \cdot$ in Eq. (4) makes the relation between the electric field and the polarization field a nonlocal one. Besides the free electrons, there are bound electrons, which constitute another mechanism to polarize the metal with light. The constitutive relation of the bound electrons is

$$\mathbf{P}_d(\mathbf{r}) = \epsilon_0 \chi_{\text{bd}}(\mathbf{r}, \omega) \mathbf{E}(\mathbf{r}), \quad (5)$$

a local relation, in contrast to Eq. (4). The total polarization field \mathbf{P} is $\mathbf{P}_d + \mathbf{P}_f$.

For infinite homogeneous systems (bulk metals), the polarization field $\mathbf{P}_{f,d}$ can be uniquely decomposed into its transverse part $\mathbf{P}_{f,d}^T$ with $\nabla \cdot \mathbf{P}_{f,d}^T = 0$ and its longitudinal part $\mathbf{P}_{f,d}^L$ with $\nabla \times \mathbf{P}_{f,d}^L = 0$, and similarly \mathbf{E} can be decomposed into its transverse and longitudinal parts \mathbf{E}^T and \mathbf{E}^L , respectively. When inserting these decompositions into Eqs. (4) and (5), independent equations for transverse and longitudinal fields can be obtained. In particular, the equation for the transverse fields is $\mathbf{P}^T = [-\epsilon_0 \omega_p^2 / (\omega^2 + i\omega\gamma) + \epsilon_0 \chi_{\text{bd}}] \mathbf{E}^T$, and the equations for the longitudinal fields are $-\beta^2 / (\omega^2 + i\omega\gamma) \nabla^2 \mathbf{P}_f^L + \mathbf{P}_f^L = -\epsilon_0 \omega_p^2 / (\omega^2 + i\omega\gamma) \mathbf{E}^L$, and $\mathbf{P}_d^L = \epsilon_0 \chi_{\text{bd}} \mathbf{E}^L$. By going to k space, two independent solutions of the dispersion relations can thus be found, two types of waves corresponding to the

transverse and the longitudinal dielectric functions

$$\epsilon_m^T(\omega) = \epsilon_{\text{bd}}(\omega) - \frac{\omega_p^2}{\omega^2 + i\omega\gamma}, \quad (6a)$$

$$\epsilon_m^L(\omega) = \epsilon_{\text{bd}}(\omega) - \frac{\omega_p^2}{\omega^2 + i\omega\gamma - \beta^2 k^2}, \quad (6b)$$

with $\epsilon_{\text{bd}} = 1 + \chi_{\text{bd}}$. The dispersion of the transverse waves is $k(\omega) = \omega \sqrt{\epsilon_m^T(\omega)} / c$, while the dispersion of the longitudinal waves is determined by $\epsilon_m^L(\omega, k) = 0$. Since they are independent solutions, the two types of waves do not interact with each other in infinitely extended metals.

Both types of waves also exist in finite homogeneous plasmonic structures, where they also propagate independently, except at boundaries. Boundary conditions dictate the generation of mixed excitations: External light, a transverse wave, not only excites transverse but also longitudinal waves in the metal.^{15,50} The transfer-matrix method for nonlocal response of metal-dielectric multilayer structures illustrates this point quite well.^{25,51} In our Green's function method below, we will also make use of this crucial fact that the transverse and longitudinal waves propagate independently within the homogeneous metal, but are not generated independently and at boundaries must occur in the right mixture so as to satisfy the boundary conditions.

IV. GREEN'S FUNCTION SURFACE INTEGRALS

A. Surface integrals for local response

We first give the known surface integrals for the local-response theory,^{40,43–45} before introducing in Sec. IV B the surface integrals for nonlocal response. In Ref. 43 it was stressed and shown that one of the advantages of the GSIM is that backgrounds such as infinite substrates can be taken into account in terms of their Green's functions. The surface integrals here are valid for arbitrary spatially inhomogeneous backgrounds. We also allow light propagation in the direction along the nanowires ($k_z \neq 0$), thereby generalizing the results of Ref. 43 where light propagation in more than two dimensions is not considered. Additionally, we allow the possibility that the inhomogeneous background responds nonlocally, as discussed in Sec. IV B. Finally, it was not clear until now how to apply the GSIM to nanostructures that touch a substrate. We solve the associated mathematical difficulties in Sec. V, which enables us to present converged numerical results of the GSIM for touching geometries in Sec. VII.

Surface integrals inside nanowires. First we give the surface-integral equations for the metal wires with the cross section A_i and the boundary S_i . In the local description, the transverse fields \mathbf{E}_i^T and \mathbf{H}_i^T are coupled and the constitutive relation reads $\mathbf{D}_i^T = \epsilon_0 \epsilon_m^T \mathbf{E}_i^T$. The \hat{x}, \hat{y} components of the fields can be expressed in terms of their \hat{z} components.⁴⁹ For positions $\boldsymbol{\rho} \in A_i$, the field components E_{zi}^T and H_{zi}^T satisfy the scalar surface integrals

$$E_{zi}^T(\boldsymbol{\rho}) = - \oint_{S_i} d\rho' [e_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zi}^T(\boldsymbol{\rho}') + e_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zi,n}^T(\boldsymbol{\rho}')], \quad (7a)$$

$$H_{zi}^T(\boldsymbol{\rho}) = - \oint_{S_i} d\rho' [m_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zi}^T(\boldsymbol{\rho}') + m_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zi,n}^T(\boldsymbol{\rho}')], \quad (7b)$$

with the integration kernels

$$e_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') = m_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') = \hat{n}_i(\boldsymbol{\rho}') \cdot \nabla_{\boldsymbol{\rho}'} g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}'), \quad (8a)$$

$$e_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') = m_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') = -g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}'). \quad (8b)$$

Here, the scalar Green's function $g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}')$ satisfies $[\nabla_{\boldsymbol{\rho}}^2 + (k_{\rho i}^T)^2]g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') = -\delta(\boldsymbol{\rho} - \boldsymbol{\rho}')$ and has the solution $iH_0^{(1)}(k_{\rho i}^T|\boldsymbol{\rho} - \boldsymbol{\rho}'|)/4$ with $H_0^{(1)}$ being the zeroth-order Hankel function of the first kind; the subscript “ n ” in $E_{zi,n}^T$ and $H_{zi,n}^T$ stands for the directional derivative normal to the surface; e.g., $E_{zi,n}^T = \hat{n} \cdot \nabla_{\boldsymbol{\rho}} E_{zi}^T$. The derivation of the surface integrals (7) is given in Appendix A 1.

Surface integrals outside of nanowires. Having discussed the surface integrals for the metal wires, we now turn to the background, which we allow to have an arbitrary spatially varying dielectric function $\epsilon_b(\boldsymbol{\rho}, \omega)$. This inhomogeneity makes the surface integrals more complicated than for the nanowires that we assumed homogeneous. For example, instead of scalar Green's functions the surface integrals will feature tensor components of dyadic Green's functions. As derived in Appendix A 2, the surface integrals for the \hat{z} components of the electric and magnetic fields are

$$E_{zb}(\boldsymbol{\rho}) = E_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \oint_S d\boldsymbol{\rho}' [e_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}')E_{zb}(\boldsymbol{\rho}') + e_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}')E_{zb,n}(\boldsymbol{\rho}')] + \oint_S d\boldsymbol{\rho}' [f_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}')H_{zb}(\boldsymbol{\rho}') + f_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}')H_{zb,n}(\boldsymbol{\rho}')], \quad (9a)$$

$$H_{zb}(\boldsymbol{\rho}) = H_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \oint_S d\boldsymbol{\rho}' [m_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}')H_{zb}(\boldsymbol{\rho}') + m_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}')H_{zb,n}(\boldsymbol{\rho}')] + \oint_S d\boldsymbol{\rho}' [h_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}')E_{zb}(\boldsymbol{\rho}') + h_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}')E_{zb,n}(\boldsymbol{\rho}')]. \quad (9b)$$

The E_{zb}^{inc} and H_{zb}^{inc} represent the \hat{z} components of the incident electric and magnetic fields. Note that the integrations in Eq. (9) are over all metal-dielectric surfaces with $S = \sum_i S_i$. Again we wrote the integration kernels in short-hand notation. They are scalar functions, given in terms of components of the background dyadic electric and magnetic Green's functions \mathbf{G}_e and \mathbf{G}_m (defined in Appendix A 2) and their spatial derivatives; i.e.,

$$(e,m)_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') = [(ik_z\hat{z} - \nabla_{\boldsymbol{\rho}'} \times \mathbf{G}_{e,m}^t(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{lz} + \frac{ik_z}{k_{\rho b}(\boldsymbol{\rho}')^2} [\hat{l}(\boldsymbol{\rho}') \cdot \nabla_{\boldsymbol{\rho}'} [(ik_z\hat{z} - \nabla_{\boldsymbol{\rho}'} \times \mathbf{G}_{e,m}^t(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}], \quad (10a)$$

$$(e,m)_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') = -\frac{k_b(\boldsymbol{\rho}')^2}{k_{\rho b}(\boldsymbol{\rho}')^2} [\mathbf{G}_{e,m}(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}, \quad (10b)$$

$$f_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') = -i\omega\mu_0[\mathbf{G}_e(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zl} + \frac{\omega\mu_0k_z}{k_{\rho b}(\boldsymbol{\rho}')^2} [\hat{l}(\boldsymbol{\rho}') \cdot \nabla_{\boldsymbol{\rho}'} \mathbf{G}_e(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}, \quad (10c)$$

$$f_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i\omega\mu_0}{k_{\rho b}(\boldsymbol{\rho}')^2} [(ik_z\hat{z} - \nabla_{\boldsymbol{\rho}'} \times \mathbf{G}_e^t(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}, \quad (10d)$$

$$h_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') = i\omega\epsilon_b(\boldsymbol{\rho}')[\mathbf{G}_m(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zl} - \frac{\omega\epsilon_b(\boldsymbol{\rho}')k_z}{k_{\rho b}(\boldsymbol{\rho}')^2} [\hat{l}(\boldsymbol{\rho}') \cdot \nabla_{\boldsymbol{\rho}'} \mathbf{G}_m(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}, \quad (10e)$$

$$h_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') = -\frac{i\omega\epsilon_b(\boldsymbol{\rho}')}{k_{\rho b}(\boldsymbol{\rho}')^2} [(ik_z\hat{z} - \nabla_{\boldsymbol{\rho}'} \times \mathbf{G}_m^t(\boldsymbol{\rho}, \boldsymbol{\rho}')]_{zz}, \quad (10f)$$

where $k_b = \omega\sqrt{\epsilon_b}/c$ and $k_{\rho b}^2 = k_b^2 - k_z^2$. The superscript “ t ” in \mathbf{G}^t represents the transpose operation. The $[\mathbf{G}]_{lz}$ is the tensor component $[\mathbf{G}]_{lz} = \hat{l}(\boldsymbol{\rho}') \cdot \{\mathbf{G}(\boldsymbol{\rho}, \boldsymbol{\rho}')\} \cdot \hat{z}$, and $[\mathbf{G}]_{zl}$ is analogously defined by $[\mathbf{G}]_{zl} = \hat{z} \cdot \{\mathbf{G}(\boldsymbol{\rho}, \boldsymbol{\rho}')\} \cdot \hat{l}(\boldsymbol{\rho}')$.

In the special case of a spatially homogenous dielectric background, the surface integrals for the background become similar to those for the homogenous plasmonic scatterer in Eq. (9). In particular, the coefficients $f_b^{0,1}$ and $h_b^{0,1}$ vanish, while $e_b^{0,1}$ and $m_b^{0,1}$ assume the same forms as $e_i^{0,1}$ and $m_i^{0,1}$ in Eq. (8), just with g_i^T replaced by the background scalar Green's function $g_b = iH_0^{(1)}(k_{\rho b}|\boldsymbol{\rho} - \boldsymbol{\rho}'|)/4$.

Returning to the general case of inhomogeneous dielectric backgrounds, one can split the dyadic Green's function into $\mathbf{G}_{e,m} = \mathbf{G}_{e,m}^0 + \mathbf{G}_{e,m}^s$, where $\mathbf{G}_{e,m}^0$ represents the dyadic Green's function for a homogeneous background, and $\mathbf{G}_{e,m}^s$ represents the scattering contribution owing to the inhomogeneity in the background.⁵² The nonvanishing scattering contribution $\mathbf{G}_{e,m}^s$ gives rise to nonzero values for $f_b^{0,1}$ and

$h_b^{0,1}$, and makes the other kernels more complicated. This is illustrated in Appendix B for the experimentally important example of a dielectric background consisting of a dielectric slab in air, i.e., a substrate layer that can support the plasmonic nanowires.

Summary of local-response GSIM. We have now in Eq. (7) described the fields inside the metal wires as the surface integrals over the fields on the interior of their surfaces, and similarly Eq. (9) gives the fields in the dielectric background in terms of the fields on the surface exterior to these metallic nanowires. For a unique solution of the fields in all of space we need to specify boundary conditions that relate the fields on both sides of the interfaces. In the local-response approximation that we consider in this subsection, these are just the usual Maxwell boundary conditions, namely that the tangential electric and magnetic fields be continuous across the boundaries. We stress that in the above we arrived at a powerful generalization of the existing local-response GSIM, by allowing the background dielectric function $\epsilon_b(\mathbf{r})$ to have

an arbitrary spatial dependence. The procedure is now to first solve for the fields on the surfaces, and after that to use these solutions in combination with the surface integrals to uniquely determine the fields in all of space. When solving for the fields on the surfaces, singularities in the integration kernels need to be dealt with. This is detailed in Sec. V, where it is also shown how to regularize additional singularities in case the surface touches a dielectric interface.

B. Surface integrals for nonlocal response

We now turn to the nonlocal-response theory and its associated surface integrals and boundary conditions. As was mentioned in Sec. III, in the hydrodynamic Drude model additional longitudinal waves exist in the metal, besides the usual transverse waves. These longitudinal and transverse waves propagate independently in the homogeneous metallic nanowires, except at their boundaries. The key insight leading to our Green's function surface-integral method for the hydrodynamic model is then that for these longitudinal waves an additional surface integral can be formulated, independently of the other two, as presented below.

Additional surface integral. The longitudinal field \mathbf{E}_i^L by definition is rotation-free and in the plasmonic nanowire A_i can thus be expressed in terms of a potential ϕ_i via

$$\mathbf{E}_i^L = -(\nabla_\rho + ik_z \hat{z})\phi_i, \quad (11)$$

where ϕ_i satisfies the scalar wave equation

$$(\nabla_\rho^2 + k_{\rho i}^L)\phi_i(\rho) = 0, \quad (12)$$

with $k_{\rho i}^L = k_i^L - k_z^2$ and $\beta k_i^L = (\omega^2 + i\omega\gamma - \omega_p^2/\epsilon_\infty)^{1/2}$. The scalar Green's function g_i^L associated with Eq. (12) is defined as the solution of

$$(\nabla_\rho^2 + k_{\rho i}^L)g_i^L(\rho, \rho') = -\delta(\rho - \rho'). \quad (13)$$

Directly analogously to the Green's functions that we introduced before, the solution is given by

$$g_i^L(\rho, \rho') = \frac{i}{4} H_0^{(1)}(k_{\rho i}^L |\rho - \rho'|). \quad (14)$$

The main physical difference is that the longitudinal wave vectors $k_{\rho i}^L$ are typically much larger than the transverse wave vectors $k_{\rho i}^T$ of the metal and $k_{\rho b}$ of the dielectric background. Analogously to the derivation of Eq. (7) in Appendix A 1, we can now derive that the potential in the interior of the metal can be expressed as an integral over the same potential at the surface,

$$\phi_i(\rho) = - \oint_{S_i} d\rho' [p_i^0(\rho, \rho')\phi_i(\rho') + p_i^1(\rho, \rho')\phi_{i,n}(\rho')], \quad (15)$$

with integration kernels

$$p_i^0(\rho, \rho') = \hat{n}_i(\rho') \cdot \nabla_{\rho'} g_i^L(\rho, \rho'), \quad (16a)$$

$$p_i^1(\rho, \rho') = -g_i^L(\rho, \rho'). \quad (16b)$$

Equation (15) is the sought surface integral for the longitudinal fields in the plasmonic nanowire. The surface integrals Eq. (7) for the transverse fields in the metal and Eq. (9) for the fields in the background simply stay the same in the hydrodynamic

Drude model. Only if one would also wish to allow metal constituents also in the background, for example to describe an infinite metal substrate,⁶ and take its nonlocal response into account, would a modification be needed for the background. We briefly discuss such a modification in Appendix A 2.

Additional boundary condition. Besides the three surface integrals (7), (9), and (15), we again need boundary conditions to obtain unique solutions for the electromagnetic fields in all of space. In local-response theory we only needed the usual Maxwell boundary conditions, as we discussed in Sec. IV A, but for nonlocal response, additional boundary conditions (ABCs) are needed. In the present paper, we only consider metal-dielectric, not metal-metal interfaces. We also assume that the static free-electron density is a step function at the interface and constant within the metal, thereby neglecting Friedel oscillations and the electron spill-out associated with quantum tunneling on the subnanometer scale. These assumptions entail that only one ABC is needed for the hydrodynamic Drude model, which is the condition that the normal component of the free-electron current be continuous and hence by charge conservation vanish at the boundary.^{15,51,53} This condition can be combined with the usual Maxwell boundary condition that in the absence of free charges the normal component of the electric displacement field be continuous across the boundary, whereby the ABC can be unambiguously expressed as⁵¹

$$\epsilon_b \hat{n} \cdot \mathbf{E}_b = \epsilon_{bd} \hat{n} \cdot \mathbf{E}_i, \quad (17)$$

with ϵ_{bd} the bound-electron response of the metal as introduced in Eq. (6). This ABC (17) implies that in general the normal components of the electric field makes a jump at the interface. Such a jump is the common situation also in the usual local-response approximation, but here in Eq. (17) the jump is different than for local response, where the jump described by $\epsilon_b \hat{n} \cdot \mathbf{E}_b = \epsilon_m^T \hat{n} \cdot \mathbf{E}_i$ follows instead from the continuity of the normal component of the displacement field alone, and where on the right-hand side the Drude part for the free electrons is included.

For clarity, let us briefly discuss the local-response limit of the HDM. Equation (17) relates electric fields just inside and just outside the metal in the HDM. Here, "just inside" means on a length scale shorter than the thickness of the surface layer in which nonlocal surface charge redistribution occurs. In the limit $\beta \rightarrow 0$, the usual Drude model is recovered from the dynamics (4), and the surface layer in which nonlocal surface charge redistribution occurs becomes infinitely thin. The additional boundary condition (17) does not depend on β and hence does not change when taking the $\beta \rightarrow 0$ limit, but it becomes irrelevant since the field "just inside" that it relates to becomes the field exactly *on* the metal-dielectric interface. Maxwell's equations alone then relate by $\epsilon_b \hat{n} \cdot \mathbf{E}_b = \epsilon_m^T \hat{n} \cdot \mathbf{E}_i$ the fields on both sides of that interface.

In the ABC (17), the electric field \mathbf{E}_i at the interior of the metal interface of the i th nanowire is the sum of the usual divergence-free electric field \mathbf{E}_i^T described by the surface integral (7) and of the (specifically hydrodynamic) rotation-free electric field \mathbf{E}_i^L , described by the surface integral (15) for its corresponding potential. The occurrence of this sum of independent solutions in a bounded region of space

makes it intuitively clear that the ABC is needed for a unique solution in all of space.

Summary of hydrodynamic GSIM. In summary, three independent surface integrals (7), (9), and (15) are needed for the hydrodynamic model, instead of the common first two for local response. These three integrals give rise to a unique and physically meaningful solution of the electromagnetic fields, when used in combination with three boundary conditions, two of which are the usual ones derived from Maxwell's equations. The third one is the additional boundary condition (17), which is derived from local conservation of free charges after neglecting quantum spill-out of free electrons.

V. NUMERICAL IMPLEMENTATION OF NONLOCAL GSIM

For clarity, we first collect the surface integrals and boundary conditions needed for the local and nonlocal Green's function surface-integral methods. Then we address the occurrence of singularities in the integration kernels of the surface integrals. By introducing a new regularization procedure for the scattering part of the Green's tensors, we extend the applicability of GSIM, both the local-response and

the nonlocal-response version, to geometries where arbitrarily shaped nanowires rest on arbitrary multilayer substrates, rather than floating or hanging slightly above them.^{16,43}

Surface integrals and boundary conditions. The first numerical task of the GSIM is to solve the fields along the nanowire boundary from the following surface integrals. For the metal side of the metal-background boundary we have three surface integrals

$$E_{zi}^T(\boldsymbol{\rho}) = - \oint_{S_i} d\boldsymbol{\rho}' [e_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zi}^T(\boldsymbol{\rho}') + e_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zi,n}^T(\boldsymbol{\rho}')], \quad (18a)$$

$$H_{zi}^T(\boldsymbol{\rho}) = - \oint_{S_i} d\boldsymbol{\rho}' [m_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zi}^T(\boldsymbol{\rho}') + m_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zi,n}^T(\boldsymbol{\rho}')], \quad (18b)$$

$$\phi_i(\boldsymbol{\rho}) = - \oint_{S_i} d\boldsymbol{\rho}' [p_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') \phi_i(\boldsymbol{\rho}') + p_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') \phi_{i,n}(\boldsymbol{\rho}')]. \quad (18c)$$

The third surface integral, Eq. (18c), is the additional one in case of nonlocal response, and is left out in the local GSIM. For the background side of the metal-background boundary we only have two surface integrals

$$E_{zb}(\boldsymbol{\rho}) = E_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \oint_S d\boldsymbol{\rho}' [e_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zb}(\boldsymbol{\rho}') + e_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zb,n}(\boldsymbol{\rho}')] + \oint_S d\boldsymbol{\rho}' [f_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zb}(\boldsymbol{\rho}') + f_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zb,n}(\boldsymbol{\rho}')], \quad (19a)$$

$$H_{zb}(\boldsymbol{\rho}) = H_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \oint_S d\boldsymbol{\rho}' [m_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zb}(\boldsymbol{\rho}') + m_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') H_{zb,n}(\boldsymbol{\rho}')] + \oint_S d\boldsymbol{\rho}' [h_b^0(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zb}(\boldsymbol{\rho}') + h_b^1(\boldsymbol{\rho}, \boldsymbol{\rho}') E_{zb,n}(\boldsymbol{\rho}')]. \quad (19b)$$

In combination with the boundary conditions

$$\hat{n}(\boldsymbol{\rho}) \times \mathbf{E}_b(\boldsymbol{\rho}) = \hat{n}(\boldsymbol{\rho}) \times \mathbf{E}_i(\boldsymbol{\rho}), \quad (20a)$$

$$\hat{n}(\boldsymbol{\rho}) \times \mathbf{H}_b(\boldsymbol{\rho}) = \hat{n}(\boldsymbol{\rho}) \times \mathbf{H}_i(\boldsymbol{\rho}), \quad (20b)$$

$$\epsilon_b(\boldsymbol{\rho}) \hat{n}(\boldsymbol{\rho}) \cdot \mathbf{E}_b(\boldsymbol{\rho}) = \hat{n}(\boldsymbol{\rho}) \cdot \epsilon_{\text{bd}} \mathbf{E}_i(\boldsymbol{\rho}), \quad (20c)$$

for $\boldsymbol{\rho}$ on the boundary S , unique solutions of Maxwell's equations in all of space can be found. Equation (20c) is the additional boundary condition for nonlocal response, which is left out in the local GSIM.

In the special case of normally incident light ($k_z = 0$), the above equations decouple into two independent sets. One is for TE-polarized light. In this case, the longitudinal fields cannot be excited. The relevant surface integrals are then Eqs. (18a) and (19a), in combination with only the boundary conditions (20a) and (20b). The other set is for TM-polarized light. In this case, the longitudinal fields can be excited. The required surface integrals are Eqs. (18b), (18c), and (19b), and all three boundary conditions in Eq. (20) play a role.

Singularities in integration kernels. Some of the integration kernels in the surface integrals for nanowires have singularities, which must be treated carefully. First consider the surface integrals for the nanowires in Eqs. (18a)–(18c). There is a singularity that comes from the Green's function of the Hankel-function type, which blows up in the limit $\boldsymbol{\rho}' \rightarrow \boldsymbol{\rho}$. We regularize the singularities following the routine by García de Abajo and Howie in Ref. 40 and by Jung and Søndergaard

in Ref. 43. In particular, we note that our additional surface integral Eq. (18c) for nonlocal response can be regularized in the same way as was known for the two others of the local GSIM,^{40,43} because the same Green's function appears in it, albeit with a different wave vector in the argument [recall Eq. (14)]. Thus the regularized version of Eq. (18c) becomes

$$\frac{1}{2} \phi_i(\boldsymbol{\rho}) = -\mathcal{P} \oint_{S_i} d\boldsymbol{\rho}' [p_i^0(\boldsymbol{\rho}, \boldsymbol{\rho}') \phi_i(\boldsymbol{\rho}') + p_i^1(\boldsymbol{\rho}, \boldsymbol{\rho}') \phi_{i,n}(\boldsymbol{\rho}')], \quad (21)$$

where “ $\mathcal{P} \oint$ ” represents the integration excluding the singular point at $\boldsymbol{\rho}' = \boldsymbol{\rho}$.

Regularization of surface integrals for background. Next we consider the surface integrals for the background in Eqs. (19a) and (19b). These integrals are the same as for local response, at least when neglecting nonlocal response in the background. Nevertheless we dwell upon them here, because even for local response we could not find in the literature the necessary regularization procedure for touching geometries that we here present.

For the background surface integrals, both the homogenous and the scattering Green's functions exhibit singularities. The singularity associated with the homogenous Green's function can be treated as above in Eq. (21). Singularities associated with the scattering part of the Green's function can also arise and must be treated differently. Let us first assume that the

background is a slab in free space. In Eq. (B1), the surface integral kernels are expressed as integrals over the wave vector k_y , with integration limits $\pm\infty$. Singularities in the kernels may arise when the integrands in Eq. (B1) do not fall off rapidly enough as k_y and hence k_{\parallel} tend to infinity. Now in many cases singularities are prevented from occurring because the reflectivities in the integrands vanish in the limit $k_{\parallel} \rightarrow \infty$. For example, $R_{TE} \rightarrow 0$ as $k_{\parallel} \rightarrow \infty$, whether the slab is a dielectric medium or metallic; $R_{TM} \rightarrow 0$ as $k_{\parallel} \rightarrow \infty$ when the slab is composed of a metal with nonlocal response.⁵⁴ By contrast, R_{TM} approaches a nonzero value as $k_{\parallel} \rightarrow \infty$ when the slab is composed of a dielectric medium, and this case includes a local-response metallic medium that is described by the dielectric function of the metal. This indicates that in particular TM-polarized scattering waves induced by dielectric substrates may lead to a singularity in the scattering part of the Green's function.

To clearly illustrate such a scattering singularity, we consider a single nanowire resting on the $x = 0$ top plane of a dielectric slab with permittivity ϵ_d and thickness t . (More general substrates are discussed below.) We take the kernel e_0^{bs} of Eq. (B1a) as an example. In the limit $k_{\parallel} \rightarrow \infty$, the slab reflectivity has the value $R_{TM}(\infty) = (\epsilon_d - 1)/(\epsilon_d + 1)$. This value is independent of the slab thickness, since waves with $k_{\parallel} \rightarrow \infty$ have an infinitely short penetration depth into the slab and hence do not probe its thickness (k_x^2 approaches $-\infty$). We then split e_0^{bs} into two parts, $e_b^{0s} = e_b^{0s1} + e_b^{0s2}$. In e_b^{0s1} we deal with the possible singularity arising due to the large- k_y behavior of the integrand of e_b^{0s} , whereas the integrand of e_b^{0s2} vanishes for large k_y so that e_b^{0s2} does not have a singularity. The possibly singular kernel term is given by

$$\begin{aligned} e_b^{0s1} &= \frac{-ik_z^2 R_{TM}(\infty)}{4\pi k_\rho^2} \int dk_y \frac{1}{k_x} \exp(i\psi) i\mathbf{k}_\rho \cdot \hat{\mathbf{n}}(\boldsymbol{\rho}') \\ &= -\frac{1}{4i} \frac{k_z^2}{k_\rho^2} R_{TM}(\infty) \hat{\mathbf{n}}(\boldsymbol{\rho}') \cdot \nabla_{\boldsymbol{\rho}'} H_0^{(1)}(k_\rho \rho_{os}), \end{aligned} \quad (22)$$

in terms of the angle $\psi = k_y(y - y') - k_x(x + x')$, the wave vectors $k_0 = \omega/c$ and $\mathbf{k}_\rho = (k_x, k_y)$ that satisfy the identity $k_\rho^2 + k_z^2 = k_0^2$, and the length $\rho_{os} = \sqrt{(x + x')^2 + (y - y')^2}$. The identity Eq. (22) follows almost directly from the plane-wave expansion of a cylindrical wave as derived in Eq. (2.2.11) of Ref. 55.

A singularity of the scattering kernel e_b^{0s1} arises when ρ_{os} vanishes for a point on the surface. When does this occur? The kernel $e_b^{0s1}(\boldsymbol{\rho}, \boldsymbol{\rho}')$ appears in the surface integral (19a), and the integration runs on the surface of the nanowire. For a cylindrical nanowire this surface would be parameterized by $(x + r_0)^2 + y^2 = r_0^2$. On the outer surface of the nanowire resting on the $(x = 0)$ plane, x and x' always have the same sign, so that ρ_{os} can only vanish if $x = x' = 0$. For $\boldsymbol{\rho}$ and $\boldsymbol{\rho}'$ on the circle, it follows that the scattering singularity occurs only in $(x, y) = (0, 0)$, where the nanowire and the dielectric substrate touch. It holds more generally for noncylindrical nanowires that scattering singularities occur on the point(s) where nanowires touch dielectric interfaces. Following the same routine as for e_b^{0s} , the singularities in other integration kernels can also be extracted. The singularities all relate to the Hankel function, which can be treated similarly as in

Eq. (21). In doing so, we end up with the following regularized background surface integrals:

$$\begin{aligned} S_e(\boldsymbol{\rho}) E_{zb}(\boldsymbol{\rho}) &= E_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \mathcal{P} \oint_S d\boldsymbol{\rho}' [e_0^b E_{zb}(\boldsymbol{\rho}') + e_1^b \dot{E}_{zb}(\boldsymbol{\rho}')] \\ &\quad + \mathcal{P} \oint_S d\boldsymbol{\rho}' [f_0^b H_{zb}(\boldsymbol{\rho}') + f_1^b \dot{H}_{zb}(\boldsymbol{\rho}')], \end{aligned} \quad (23a)$$

$$\begin{aligned} S_m(\boldsymbol{\rho}) H_{zb}(\boldsymbol{\rho}) &= H_{zb}^{\text{inc}}(\boldsymbol{\rho}) + \mathcal{P} \oint_S d\boldsymbol{\rho}' [m_0^b H_{zb}(\boldsymbol{\rho}') + m_1^b \dot{H}_{zb}(\boldsymbol{\rho}')] \\ &\quad + \mathcal{P} \oint_S d\boldsymbol{\rho}' [h_0^b E_{zb}(\boldsymbol{\rho}') + h_1^b \dot{E}_{zb}(\boldsymbol{\rho}')], \end{aligned} \quad (23b)$$

with

$$S_e(\boldsymbol{\rho}) = \frac{1}{2} \left[1 - \frac{k_z^2}{k_\rho^2} R_{TM}(\infty) \right], \quad (24a)$$

$$S_m(\boldsymbol{\rho}) = \frac{1}{2} \left[1 - \frac{k_0^2}{k_\rho^2} R_{TM}(\infty) \right], \quad (24b)$$

for $\boldsymbol{\rho}$ at the common boundary of the nanowire and the slab, and otherwise $S_e = S_m = 1/2$.

Until now we have assumed that the substrate is a dielectric slab. We already discussed why the regularization procedure does not depend on the thickness of this slab. For the same reasons, we can generalize the substrate to an arbitrary multilayer dielectric. The above regularization in Eqs. (23b) and (24) involves reflectivities in the limit $k_y \rightarrow \infty$. In this limit the reflectivity of a multilayer dielectric will be given by $R_{TM}(\infty) = (\epsilon_d - 1)/(\epsilon_d + 1)$, where ϵ_d is to be understood as the dielectric function of the outer layer of the multilayer substrate on which the nanowires rest. Also, if the substrate is not exactly planar, then locally near the nanowire it can be approximated as planar and again the above regularization can be employed, again involving the limit reflectivity $R_{TM}(\infty)$ of the dielectric material on which the nanowire rests.

By this regularization procedure we have extended the geometries to which the computationally efficient GSIM can be applied to experimentally relevant structures where nanowires of arbitrary shapes rest on arbitrary multilayer substrates. As will be shown in Sec. VII, it is also these touching geometries for which differences between local and nonlocal response of the plasmonic nanowires are largest. Nanowires positioned above the substrate (i.e., nontouching geometries) are slightly simpler to analyze, because a singularity associated with the scattering part of the Green's function does not arise and the above regularization is not needed.

After the above regularizations, the numerical procedure to find solutions with the local or nonlocal GSIM is as follows. By discretizing the regularized surface integrals, and using the boundary conditions, the fields along the metal-background boundaries can be solved. Then, knowing the fields on the boundaries, we can employ the surface integrals once more to obtain the fields in any position of the system, and to extract further physical quantities of our interest.

VI. EFFECTIVE THEORY FOR NONLOCAL BLUESHIFTS

Before applying the hydrodynamic GSIM as developed in the previous sections, we will here give a semiquantitative

analysis of the most conspicuous optical effect of nonlocal response, namely the nonlocal blueshift of plasmonic resonance frequencies. As illustrated below, different plasmonic resonances exhibit different nonlocal blueshifts. Our analysis will explain this, and will guide our numerical investigations in Sec. VII.

For simplicity rather than necessity, we will neglect the dielectric response of the bound electrons in the metal; i.e., we take $\epsilon_{\text{bd}} = 1$. This approximation is better at lower frequencies, in particular below the band gap energy for interband transitions in the metal.

Let us consider a single subwavelength plasmonic resonator with an arbitrary shape in an inhomogeneous dielectric medium. The region of the plasmonic scatterer is denoted by A_m and has a boundary S_m . First we make the usual local-response approximation, and assume that the resonator supports a SP resonance at $\omega_{\text{res}}^{\text{loc}}$. Neglecting loss, the equation of motion for the free electrons is $m_e(\omega_{\text{res}}^{\text{loc}})^2 \mathbf{d} = -e\mathbf{E}$, where \mathbf{d} represents the displacement of the electron. The displacement \mathbf{d} gives rise to a delta-function thin surface charge distribution α_m at the boundary. This α_m then induces the screening charge α_b in the background.

Let us now take instead nonlocal response into account, using the same hydrodynamic model and stepwise equilibrium free-electric density for which we derived the nonlocal GSIM. Quantum spill-out and spatial variations of the equilibrium free-electric density are thus neglected, and consequently the normal component of the linear-response free-electron current vanishes on the metal-dielectric boundary. We then find that the hydrodynamic pressure gradient smears out the linear-response surface charge α_m into a surface charge distribution of finite thickness, decaying away from the surface and *into* the metal approximately exponentially as $\exp(-|k^L|\Delta)$, where k^L is the longitudinal wave number, and Δ the distance to the boundary. Thus, rather than exactly on the surface S_m as for local response, the nonlocal surface charge is effectively accumulated on a smaller boundary S'_m , at a distance of $1/|k^L|$ within S_m . In the region inside S'_m , denoted by A'_m , the electric field \mathbf{E}' is enhanced owing to the inward displacement of the surface charge by the pressure-gradient force. For A'_m to exist, we must of course require that the surface layer thickness $1/|k^L|$ be smaller than the effective radius. We furthermore assume that the free-electron displacement \mathbf{d} within A'_m is unchanged. Then, we approximately have $m_e(\omega_{\text{res}}^{\text{nloc}})^2 \mathbf{d} = -e\mathbf{E}'$, where $\omega_{\text{res}}^{\text{nloc}}$ represents the new resonance frequency modified by the nonlocal response. This explains that the nonlocal response indeed blueshifts the resonance frequency; i.e., $\omega_{\text{res}}^{\text{nloc}} > \omega_{\text{res}}^{\text{loc}}$. Moreover, we can also understand the blueshift quantitatively. By relating $\omega_{\text{res}}^{\text{nloc}}$ to $\omega_{\text{res}}^{\text{loc}}$ by integrating the free-electron equation of motion in the area A'_m , we find the approximate relation

$$\omega_{\text{res}}^{\text{nloc}} \approx \omega_{\text{res}}^{\text{loc}} \left(\frac{\int_{A'_m} d\mathbf{r} |\mathbf{E}'|^2}{\int_{A'_m} d\mathbf{r} |\mathbf{E}|^2} \right)^{1/4}, \quad (25)$$

which we will test in Sec. VII using our nonlocal GSIM.

As a specific example that allows analytical treatment, let us now consider a subwavelength metallic cylinder with radius r_0 in a homogenous dielectric background, and use Eq. (25) to derive the resonance frequency $\omega_{\text{res}}^{\text{nloc}}$.

In local response and in the quasistatic limit, it is well known that the cylinder supports a SP resonance at the frequency $\omega_{\text{res}}^{\text{loc}}$ for which $\epsilon_m^T(\omega_{\text{res}}^{\text{loc}})$ equals $-\epsilon_b$. At the boundary, the surface charge $\alpha = \alpha_m + \alpha_b$ is accumulated such that $\alpha_b/\alpha_m = (\epsilon_m^T \epsilon_b - \epsilon_m)/(\epsilon_b - \epsilon_m^T \epsilon_b)$. The electric field \mathbf{E} in the metal is the sum of two terms, $\mathbf{E} = \mathbf{E}_m + \mathbf{E}_b$, where $\mathbf{E}_{m,b}$ are due to the charge densities $\alpha_{m,b}$, respectively, which control their relative magnitude by $|\mathbf{E}_m|/|\mathbf{E}_b| = \alpha_m/\alpha_b$.

Turning from local to nonlocal response, the charge density α_m is effectively distributed on a smaller surface, as discussed above, characterized by a smaller radius r'_0 equal to $(r_0 - 1/|k^L|)$. We thus have $|k^L|r_0 > 1$ as the consistency requirement for our effective theory. The smaller effective radius results in an enhancement of \mathbf{E}_m approximately by a factor of $(r_0/r'_0)^l$, where l represents the angular momentum of the SP mode. For example, $l = 1$ for the dipole mode, $l = 2$ for the quadrupole mode, etc. Furthermore, the surface charge density α_b is reduced by a factor of $(r'_0/r_0)^l$, and \mathbf{E}_b is correspondingly reduced by the same factor. Taking these effects together, and assuming that nonlocal effects are small, the total electric field in the metal is approximately enhanced by a factor $1 + \epsilon_b l / (|k^L|r_0)$. Inserting these results into Eq. (25), the nonlocal resonance frequency $\omega_{\text{res}}^{\text{nloc}}$ is found to be

$$\omega_{\text{res}}^{\text{nloc}} \approx \omega_{\text{res}}^{\text{loc}} \left(1 + \frac{\epsilon_b l}{2|k^L|r_0} \right) \quad (\text{cylinder}). \quad (26)$$

Thus the magnitude of the nonlocal blueshift essentially depends on four parameters, namely the longitudinal wave vector k^L , the particle size r_0 , the angular-momentum number l of the resonant SP mode under consideration, and finally the background dielectric function ϵ_b . A higher ϵ_b gives rise to a larger blueshift.⁵⁶ Of the two parameters of the nanowire k^L and r_0 , the former is determined by the intrinsic nonlocal β factor and the operating frequency, while the latter can be experimentally varied. Smaller plasmonic resonators give rise to a larger nonlocal blueshift, as is well known. Less well known, although seen but not analyzed in early calculations for a sphere in a homogeneous background,⁵⁷ is our important point that the nonlocal blueshift grows with the angular momentum of the SP mode. In general, mode profiles corresponding to higher values of l show a stronger spatial variation.

By the same approach as leading to Eq. (26), the nonlocal resonance frequency for a plasmonic sphere can be derived as

$$\omega_{\text{res}}^{\text{nloc}} \approx \omega_{\text{res}}^{\text{loc}} \left[1 + \frac{\epsilon_b(l+1)}{2|k^L|r_0} \right] \quad (\text{sphere}). \quad (27)$$

From Eqs. (26) and (27) we find that the relative blueshift $(\omega_{\text{res}}^{\text{nloc}} - \omega_{\text{res}}^{\text{loc}})/\omega_{\text{res}}^{\text{loc}}$ depends on the nature of the plasmonic resonance and grows linearly with the angular momentum number l , both for cylinders and for spheres. We will illustrate this angular momentum dependence by numerically accurate calculations below. Furthermore, this dependence could be tested experimentally. Clearly, from the factors $(l+1)$ in Eq. (27) and l in Eq. (26) it follows that a 3D plasmonic sphere is more sensitive to the nonlocal response than a 2D wire. The relative difference in their blueshifts is a factor of two for the dipole resonance with $l = 1$, and approaches unity for high-order resonances.

Besides the l dependence, our second important point is that Eq. (26) leads to some useful insights about blueshifts of plasmonic nanowires in *inhomogeneous* backgrounds, even though it was derived for a homogeneous background. For example, if we embed a cylindrical plasmonic nanowire on a substrate, then as a result it will typically exhibit a more inhomogeneous and more confined mode profile.^{58,59} Consequently the expansion of the surface charge density into angular-momentum eigenmodes will show a larger contribution from larger angular momenta. Based on the effective theory developed here and in particular in Eq. (26), we expect an accordingly larger nonlocal blueshift. We will quantify and verify this idea in Sec. VII, using our numerically accurate nonlocal GSIM method developed in Sec. IV.

VII. NUMERICAL ANALYSIS: NANOWIRE WITH AND WITHOUT A SUBSTRATE

In this section we employ the hydrodynamic GSIM as developed in Sec. IV to numerically investigate the optical effects of nonlocal response in plasmonic nanowires. The important physical quantity considered here is the extinction cross section σ_{ext} , which in general is defined as the sum of the absorption and scattering cross sections,

$$\sigma_{\text{ext}} = \sigma_{\text{abs}} + \sigma_{\text{sca}}. \quad (28)$$

Cross sections are usually defined as an area, but for the infinitely long nanowires that we consider here, we will instead consider cross sections per length unit of the nanowire, with the dimension of a length. So let us now introduce the cross sections σ_{abs} and σ_{sca} .

We consider TM-polarized incident plane waves with $H_{zb}^{\text{inc}} = \exp(ik_0x)$ and $E_{zb}^{\text{inc}} = 0$. The ratio between the electromagnetic power that is absorbed by the nanowire and the incident electromagnetic power of the plane wave is known as the absorption cross section σ_{abs} , which can be expressed as the surface integral

$$\sigma_{\text{abs}} = \oint_S d\rho \operatorname{Re} \left(\frac{iH_{zb}H_{zb,n}^*}{k_b|H_{zb}^{\text{inc}}|^2} \right), \quad (29)$$

where the superscript “*” represents the complex conjugate operation. Likewise, the scattering cross section σ_{sca} is defined as the ratio between the electromagnetic power that is scattered by the nanowire and the incident power. In a lossless homogeneous background, such as free space, the scattered power can be expressed in terms of only the far-field radiation power. By contrast, in an inhomogeneous and lossy background, the scattered power includes besides the far-field radiation power both the localized waveguide power and the power absorbed by the lossy background. The scattering cross section σ_{sca} can also be written as a surface integral,

$$\sigma_{\text{sca}} = - \oint_S d\rho \operatorname{Re} \left(\frac{iH_{zb}^s H_{zb,n}^{s*}}{k_b|H_{zb}^{\text{inc}}|^2} \right), \quad (30)$$

where H_{zb}^s represents the scattered field defined as $H_{zb}^s = H_{zb} - H_{zb}^{\text{inc}}$, and $H_{zb,n}^s = H_{zb,n} - H_{zb,n}^{\text{inc}}$.

Below we present calculations of extinction cross sections, first for freestanding plasmonic nanowires in Sec. VII A, then

for nanowires above a substrate in Sec. VII B, and finally in Sec. VII C for nanowires resting directly on a substrate.

A. Freestanding nanowire

The extinction cross section of a freestanding nanowire can be computed analytically, not only in local response but also in the hydrodynamic Drude model.^{15,18} It is thus an excellent benchmark problem for numerical methods. In fact, this same benchmark problem was used independently by two groups to show the accuracy of their finite-element method implementations of the hydrodynamic model.^{22,46} Here we put our nonlocal GSIM to the same test.

We consider an Au cylinder with a radius r_0 in a free-space background. An exact solution can be found by a nonlocal extension of Mie scattering theory.^{15,18,23} We use the following parameters for gold: $\hbar\omega_p = 8.812$ eV, $\hbar\gamma_f = 0.0752$ eV, and $v_F = 1.39 \times 10^6$ m/s. As in Sec. VI, for simplicity we neglect the contribution of the bound electrons; i.e., we take $\epsilon_{\text{bd}} = 1$.

In Figs. 2(a)–2(c), the extinction cross section curves are depicted for the local model as well as the nonlocal HDM, by using both the GSIM and Mie scattering theory. Clearly, the results from two different methods agree very well with each other. This verifies the validity of the GSIM as a numerically accurate method, both for local response (as is known in the literature) and for nonlocal response (which is our new result). For nonlocal response the benchmark is more stringent, since not only the blueshifted resonances should come out right with the nonlocal GSIM, but also the series of hydrodynamic resonances above the plasma frequency. And indeed they do.

In Fig. 2(a), two peaks P_1 and P_2 are observed below the plasma frequency. They correspond to the dipole and quadrupole resonances, respectively. P_1 is broader than P_2 because the dipole resonance is more radiative than the quadrupolar one. Comparing the local and nonlocal curves, their resonance frequencies are nearly the same, because nonlocal response has a weak effect on the structure that is much larger than the Thomas-Fermi screening length.^{15,18,22} For Fig. 2(b) we reduce r_0 to 10 nm and observe that the P_2 disappears from the extinction cross sections, exemplifying that the quadrupole resonance becomes harder to excite by a plane wave as the nanowire becomes smaller. Furthermore, P_1 is narrower than in Fig. 2(a) because the smaller scatterer is less radiative. A tiny nonlocal blueshift of P_1 becomes just visible, and it is indeed known that nonlocal blueshifts increase as the size r_0 decreases. Here for $r_0 = 10$ nm, the relative blueshift $\Delta\omega_{\text{res}} \equiv (\omega_{\text{res}}^{\text{nlc}} - \omega_{\text{res}}^{\text{loc}})/\omega_{\text{res}}^{\text{loc}}$ is a mere 0.6%. In Fig. 2(c), we take $r_0 = 2$ nm. The nonlocal response blueshift for P_1 resonance becomes clearly noticeable with $\Delta\omega_{\text{res}} \approx 3\%$. Additionally, a series of peaks corresponding to the optical excitations of resonant longitudinal modes appear above the plasma frequency,^{15,18,22} the so-called unusual resonances.¹⁵ By contrast, these longitudinal modes were not visible in Figs. 2(a) and 2(b), because the frequency spacing between the different longitudinal modes becomes smaller for larger nanowires, and the damping loss smears out these modes.

In the three insets of Fig. 2(c), the electric-field distributions for the P_1 dipole resonances of the local and nonlocal cases, and also for the first-order hydrodynamic resonance L_1 , are shown. The electric field of the P_1 resonance in the local

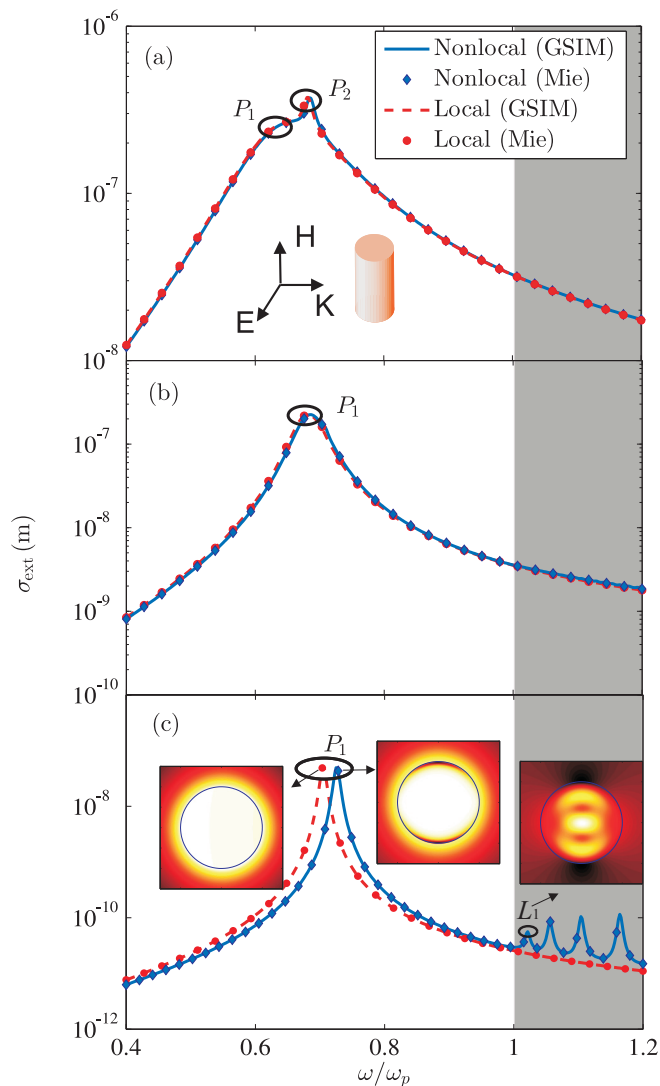


FIG. 2. (Color online) Extinction cross section of an Au cylinder in a free-space background, for a TM-polarized incident plane wave. The cylinder is described both by the usual local-response model and by the hydrodynamic Drude model. The simple wire geometry serves as an excellent benchmark problem: Analytically exact calculations (local and nonlocal Mie theory) are compared with two numerically accurate methods (local and our nonlocal GSIM). The cylinder radius is (a) 20 nm, (b) 10 nm, and (c) 2 nm. The insets in (c) show the electric field distributions $|\mathbf{E}|$ for the P_1 dipole resonance, and for the L_1 hydrodynamic resonance.

case is nearly constant inside the cylinder, as expected for the local-response dipole mode of a strongly subwavelength structure. By contrast, the corresponding nonlocal mode shows a sharp field variation inside the cylinder, a variation that can be ascribed to the evanescent longitudinal waves that are generated at the boundaries [compare Fig. 2(c) of Ref. 48]. For the L_1 hydrodynamic mode, a standing-wave pattern due to the propagating longitudinal wave is observed with our surface-integral method, similar to Fig. 2(b) of Ref. 15. The analytical method and our nonlocal GSIM produce the same fields on the cylinder boundary. By Eqs. (29) and (30) the extinction curves are then also the same, as Fig. 2(c) illustrates. Furthermore, the agreement of the fields on the surface by

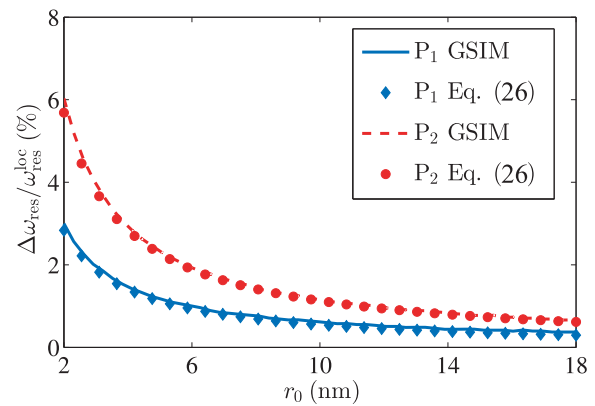


FIG. 3. (Color online) The nonlocal blueshift $\Delta\omega_{\text{res}}$ of the dipole and quadrupole resonances for an Au cylinder in free space, as a function of cylinder radius r_0 .

Eqs. (18a) and (19a) also implies that the field distributions are the same in all of space.

To test the effective theory of Sec. VI, in Fig. 3 we show the relative blueshift $\Delta\omega_{\text{res}}$ as a function of r_0 , both for the dipole and the quadrupole resonances of the freestanding nanowire. To excite the quadrupole resonance P_2 , we used a cylindrical wave with angular momentum $l = 2$ as the incident wave. The relative blueshifts $\Delta\omega_{\text{res}}$ are calculated twice, using our nonlocal GSIM and our approximate expression Eq. (26). Clearly, the results from the two methods are in good agreement, which is a first test of the validity of the effective theory in Sec. VI. The information contained in Fig. 3 is twofold: (i) The nonlocal blueshift increases for smaller radius r_0 , as was known before;¹⁵ (ii) the relative nonlocal blueshift for the quadrupole resonance is indeed two times larger than that for the dipole resonance, in agreement with Eq. (26). This significant l dependence is a new result. Our GSIM confirms that higher order SP resonances are significantly more sensitive to nonlocal response.

B. Nanowire above a dielectric substrate

Let us now consider the extinction cross section of a plasmonic nanowire positioned at a finite height above a semi-infinite dielectric substrate, and investigate the interactions between wire and substrate. The cross sections are computed using Eqs. (28)–(30). In comparison to other methods such as the finite-element method, the unique advantage of the GSIM is that even infinitely long and thick substrates can be taken exactly into account, by using the exact background Green's function (given in Appendix B). This advantage of the GSIM was stressed in Ref. 43 for local response, and here we illustrate that the nonlocal GSIM has the same advantage.

Substrate enhances nonlocal blueshift. In Figs. 4(a)–4(c), we present extinction cross sections of the same three nanowires as in Fig. 2, but now positioned a single nanometer above a dielectric substrate with refractive index 1.5. In comparison to Fig. 2, both local and nonlocal resonances are now redshifted. A simple explanation is that the substrate increases the average background permittivity of the wire, and plasmonic nanowires in homogeneous backgrounds with higher permittivities have lower resonance frequencies.⁵⁶

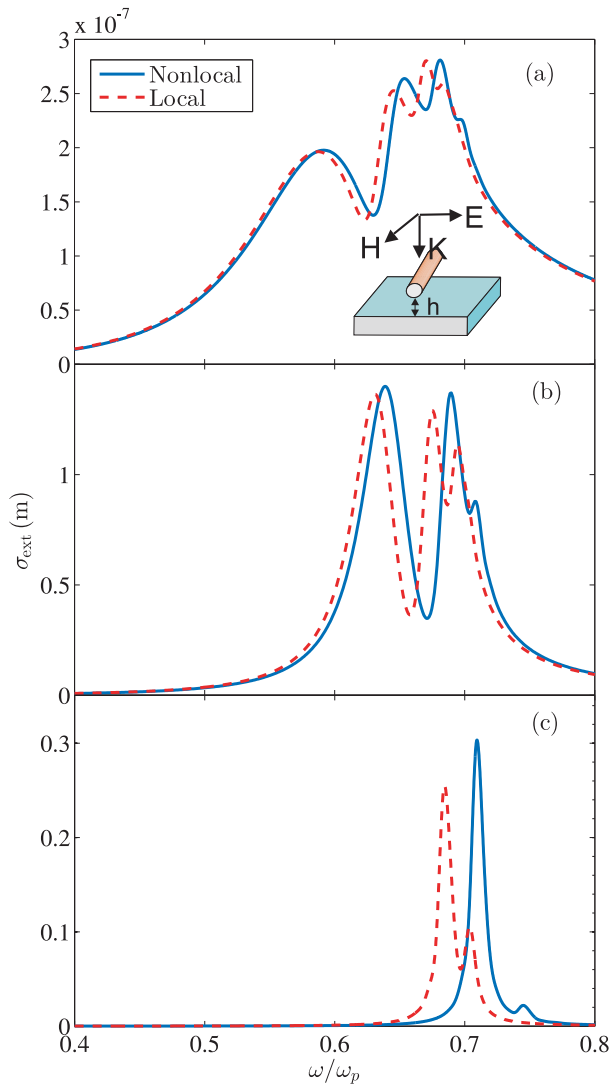


FIG. 4. (Color online) The extinction cross section of an Au cylindrical nanowire positioned above a dielectric substrate of refractive index 1.5, for a TM-polarized plane wave incident from the top. The distance between the Au cylinder and the substrate is $h = 1$ nm. The wire radius r_0 is (a) 20 nm, (b) 10 nm, and (c) 2 nm. Local and nonlocal results are calculated with standard local GSIM and with our generalized nonlocal GSIM, respectively.

Alternatively, the redshift could also be explained from the hybridization theory by considering the interactions between the nanowire and its electromagnetic image induced by the substrate.^{22,60–63} For the hydrodynamic Drude model, in Fig. 4 again blueshifts are observed with respect to the local-response resonances. In particular, for the first-order SP resonance mode, we find the relative blueshifts $\Delta\omega_{\text{res}} \approx 0.7\%$ for $r_0 = 20$ nm, and $\Delta\omega_{\text{res}} \approx 1.25\%$ for $r_0 = 10$ nm, and finally $\Delta\omega_{\text{res}} \approx 3.6\%$ for $r_0 = 2$ nm. These relative blueshifts are larger than those of Sec. VII A without the substrate, so bringing a substrate close to nanowires enhances their nonlocal blueshifts. This is interesting and of practical importance for the interpretation of experiments—for example, for the EELS experiments on few-nanometer-sized plasmonic spheres on substrates of Ref. 56, where larger blueshifts were observed

than calculated hydrodynamic blueshifts for freestanding nanospheres.

Qualitative explanation of larger blueshift. To qualitatively explain why the substrate enhances the nonlocal effects, we boldly apply Eq. (26) that was originally derived for homogeneous backgrounds to inhomogeneous ones. According to Eq. (26), the background may affect the nonlocal blueshift through effectively modifying the background dielectric function ϵ_b and the angular momentum l associated with the resonance. First, in the presence of the substrate one can interpret ϵ_b as an average value, which characterizes the screening charge contribution from the substrate. Independent of how this average is computed, this average value goes up when adding the substrate to the initial free-space environment. Equation (26) then tells us that the nonlocal blueshift of the plasmonic wire's resonance increases. Second, regarding the angular-momentum parameter l , the substrate breaks the symmetry of the background, and makes the mode profile more inhomogeneous, as is known for local response. To illustrate this both for local and nonlocal response, we plot the radial component of the outer electric field along the nanowire boundary for the first- and second-order resonance modes in Figs. 5(a) and 5(b), respectively. Despite the nearby substrate, these modes are still similar to the pure dipole and quadrupole modes in the absence of a substrate. The radial components E_r for local and nonlocal response differ

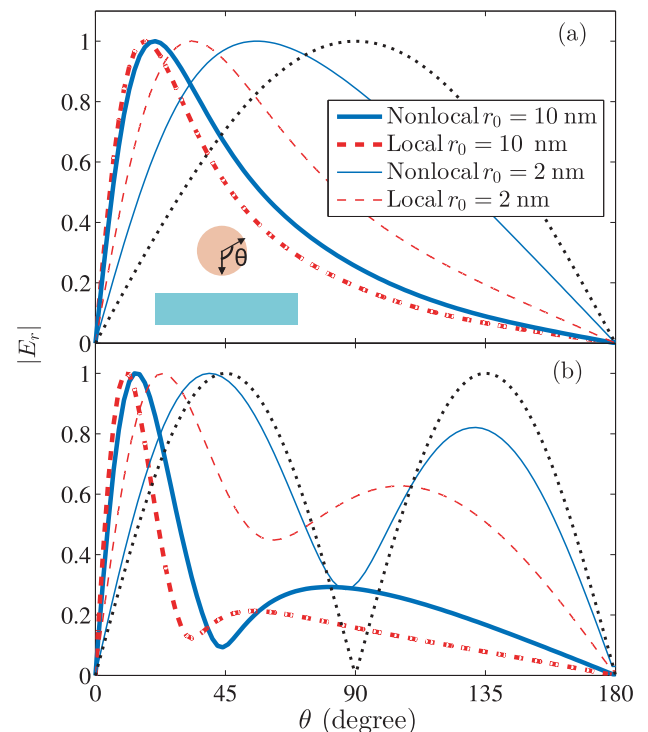


FIG. 5. (Color online) For the wire with $h = 1$ nm above the substrate as shown in the inset of Fig. 4, the radial component of the electric field (scaled with respect to its maximal value) along the nanowire boundary of the background side for (a) the first-order SP resonance mode and (b) the second-order SP resonance mode, calculated with the local GSIM as well as the nonlocal GSIM. The dotted curves in (a) and (b) represent the pure dipole and quadrupole modes, respectively, for the same nanowire but without a substrate.

considerably for all angles, and one reason is the additional boundary condition (17) for nonlocal response. However, the important point is that all field distributions shown in Fig. 5 are more concentrated on the substrate side, and in that sense are more inhomogeneous. This implies that the effective angular momenta of the lowest two resonance modes should be larger than $l = 1$ and 2 , respectively. Based on the hydrodynamic Drude model and in particular on Eq. (26), one therefore expects concomitant larger nonlocal blueshifts due to the presence of the substrate. Thus we can indeed qualitatively explain that the substrate enhances the nonlocal blueshift. It does so by increasing both the average ϵ_b and the effective l .

Quantitative explanation of larger blueshift. To support the above qualitative arguments by numbers, we will quantify how the substrate modifies the parameters ϵ_b and l . We first define the *effective angular momentum* l_{eff} for an arbitrary mode profile in the local-response approximation, by expanding its associated surface charge on the metal surface $\alpha_m(\rho)$ in cylindrical harmonics, with weights α_{ml} . The derivation can be found in Appendix C, and for local response the result is

$$l_{\text{eff}}^{-1} = \frac{\sum_{l \neq 0} |\alpha_{ml}|^2 l^{-1}}{\sum_{l \neq 0} |\alpha_{ml}|^2}. \quad (31)$$

In the case without the substrate, Eq. (31) reproduces the exact angular momenta $l = 1$ and $l = 2$ for the dipole and quadrupole modes, respectively. With the substrate as in Fig. 4,

we numerically calculate with Eq. (31) the effective angular momentum for the first-order resonance mode, as shown in Fig. 6(a1). As expected, we see that $l_{\text{eff}} > 1$ and increases with r_0 . This is consistent with the field distributions in Fig. 5(a), where the field is more enhanced on the substrate side for $r_0 = 10$ nm than for 2 nm.

Next, let us consider how one could define the *effective background permittivity* ϵ_b^{eff} . As seen in Fig. 5, the field distributions of the first- and second-order modes still resemble the pure dipole and quadrupole resonance modes of freestanding nanowires. We define the effective permittivity ϵ_b^{eff} as the homogeneous background permittivity around the nanowire that would produce the same local resonance frequency as does the nanowire in the inhomogeneous background. The same definition was used in Ref. 5.

In Fig. 6(a1), where the nanowire system is as in Fig. 4, we plot ϵ_b^{eff} for the first-order resonance mode. Clearly, ϵ_b^{eff} is larger than unity, the value in the absence of the substrate. It is important to notice that ϵ_b^{eff} increases approximately by 20% as the nanowire radius r_0 grows from 2 nm to 18 nm. Despite the different geometries and materials considered, this increase is somewhat in conflict with Ref. 5, where in the analysis of EELS experiments on spheres on supporting thin substrates, it was assumed that ϵ_b^{eff} is independent of the sphere radius. Panel 6(a2) shows a weaker dependence of ϵ_b^{eff} on the wire-substrate distance h , at least for $h > 1$ nm. The case of wires touching the substrate ($h = 0$ nm) will be addressed in Sec. VII C.

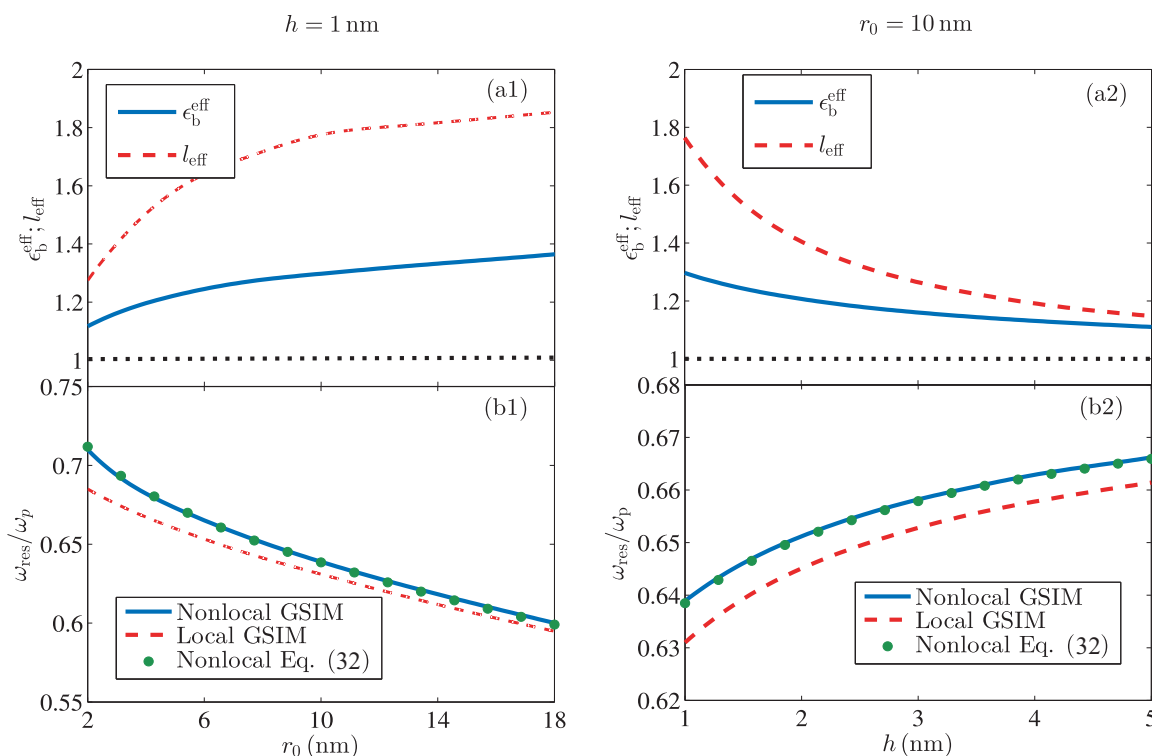


FIG. 6. (Color online) For the wire above the substrate as shown in the inset of Fig. 4, in panels (a1) and (b1) we keep the wire-substrate distance fixed at $h = 1$ nm and vary the wire radius r_0 , while in panels (a2) and (b2) we keep the radius fixed at 10 nm and vary its distance to the substrate. For the first-order resonance, panels (a1) and (a2) show effective angular momenta and average background dielectric functions in nonlocal response, while panels (b1) and (b2) show scaled resonance frequencies both in local and nonlocal response. The dot curves in (a1) and (a2) correspond to the first-order (pure dipole) resonance in nanowires without the substrate.

We defined l_{eff} to quantify effects of the inhomogeneity of the substrate, whereas we defined ϵ_{eff} assuming that the background can be described as an effectively homogeneous one. There is no real contradiction here and the results that we obtain are accurate as long as nonlocal blueshifts are small perturbations, as we shall see. To prove that the definitions of ϵ_b^{eff} and l_{eff} make good physical sense, also in combination, we define the *effective relative nonlocal blueshift* $\omega_{\text{res}}^{\text{nl},\text{eff}}$ based on the expression Eq. (26) for nonlocal blueshift in a homogeneous medium, as

$$\Delta\omega_{\text{res}}^{\text{nl},\text{eff}} = \frac{\epsilon_b^{\text{eff}} l_{\text{eff}}}{2|k^L| r_0}, \quad (32)$$

thus simply replacing l and ϵ_b in Eq. (26) by their effective values as defined above. We calculate this dimensionless blueshift for the first-order SP mode in Figs. 6(b1) and 6(b2), varying r_0 and h , respectively. In the same panels the numerically accurate resonances $\omega_{\text{res}}^{\text{nl},\text{loc}}$ and $\omega_{\text{res}}^{\text{nl}}$ are also shown. It is seen in Figs. 6(b1) and 6(b2) as one of our main results that the effective nonlocal blueshift (32) agrees quite well with the numerically accurate value in the large and physically relevant parameter range where the nonlocal blueshift stays within a few percent. This confirms that our definitions of the effective parameters ϵ_b^{eff} and l_{eff} are useful, and that we can apply our effective theory to explain nonlocal blueshifts of nanoplasmonic wires even in inhomogeneous backgrounds.

C. Nanowire on a dielectric substrate

Having studied nanowires without substrate and above a substrate, in this subsection we consider nanowires directly on a dielectric substrate, i.e., the touching geometry with $h = 0$. This is the first geometry in this paper for which the new regularization of the scattering Green's functions in the GSIM of Sec. V is required. It is required both for the local and for the nonlocal GSIM. In Figs. 7(a)–7(c), extinction curves are depicted for the same three wire radii as before. The substrate is also the same as in Fig. 4. However, this time the local extinction curves and their nonlocal counterparts show completely different features. This is quite unlike the freestanding nanowire in Fig. 2 and the wire above the substrate in Fig. 4, where the nonlocal response as a small perturbation only modifies the local curves slightly. The brief explanation is that the SP mode in the local description diverges in the limit of vanishing gap size between the nanowire and the dielectric substrate, whereas in the nonlocal HDM, no such divergence occurs.

Analogous large differences between local and nonlocal response for touching plasmonic nanoparticles have been predicted for the absorption cross section of two touching plasmonic spheres already by Fuchs and Claro in Ref. 19. Recently Fernández-Domínguez *et al.* elegantly combined transformation optics with the hydrodynamical model to calculate the field enhancement near two touching plasmonic nanowires.²⁰ The general picture is that upon reducing the distance from 1 nm down to zero, the local-response resonances vary wildly even in the final angstrom distance, whereas the nonlocal-response resonances “freeze-out.” Our Fig. 7 illustrates that one does not need plasmonic dimers to see

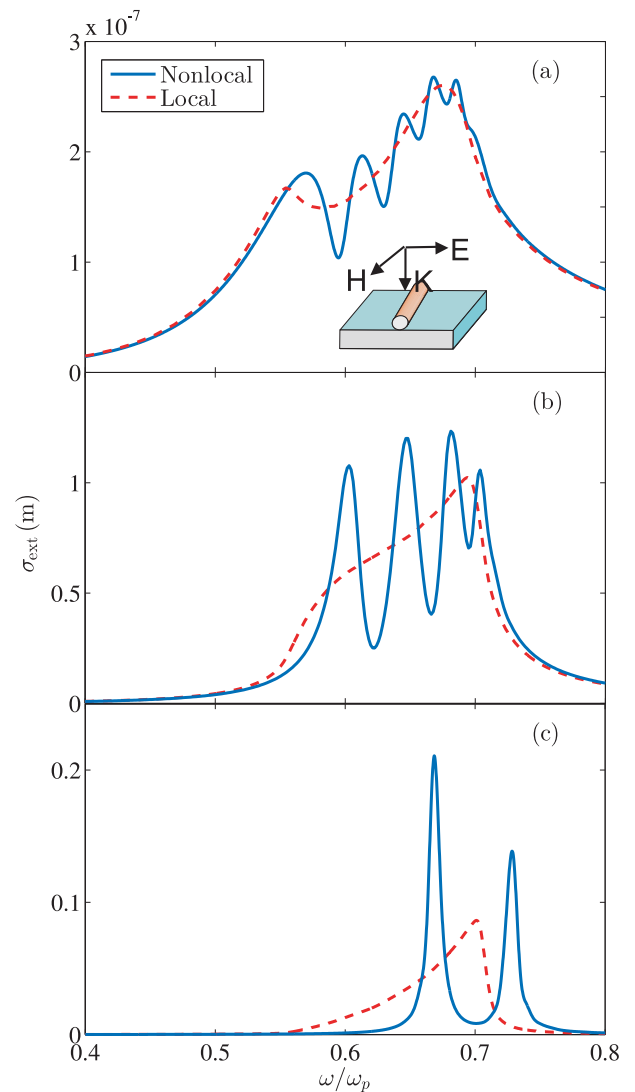


FIG. 7. (Color online) The extinction section for an incident TM-polarized plane wave of a gold cylindrical nanowire resting directly ($h = 0$) on a dielectric substrate of refractive index 1.5. The nanowire radii are (a) 20 nm, (b) 10 nm, and (c) 2 nm.

such large differences between local and nonlocal response in the (almost) touching geometry, since a single plasmonic nanowire above/on a planar dielectric substrate suffices.

To better understand the differences for local and nonlocal response for our geometry, we model the surface plasmons in the (near-)touching region as those of a planar metal-air-dielectric structure. In the quasistatic limit the dispersion relation of the local SP mode supported by the metal-air-dielectric sandwich structure is $\tanh(k_{\text{sp}}h) = -(\epsilon_d + \epsilon_m^T)/(1 + \epsilon_d\epsilon_m^T)$, where h is the thickness of the air gap, and ϵ_d is the permittivity of the semi-infinite dielectric substrate. This dispersion relation entails that k_{sp} diverges as $h \rightarrow 0$. Moreover, the local SP mode exists when $-\epsilon_d < \epsilon_m^T(\omega) < -1$, which agrees well with the frequency range in Fig. 7 where the local extinction cross section is large.

In the nonlocal HDM on the other hand, the dispersion relation changes into $\tanh(k_{\text{sp}}h) = -(\epsilon_d + \epsilon_m^T + \epsilon_d\Delta_L)/(1 + \epsilon_d\epsilon_m^T + \Delta_L)$, where $\Delta_L = k_{\text{sp}}(\epsilon_m^T - 1)/\kappa^L$, and

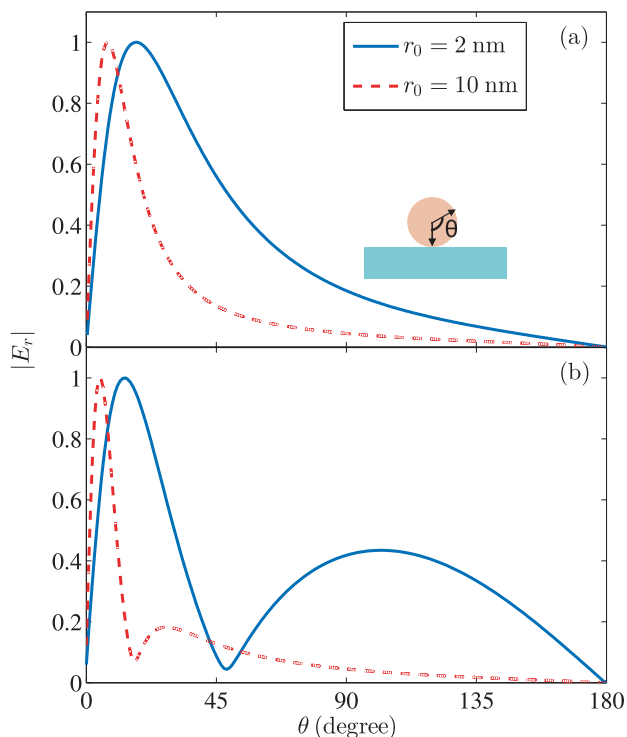


FIG. 8. (Color online) For the same nanowire resting on a substrate as in Fig. 7, the normal component of the electric field scaled with respect to its maximal value along the nanowire boundary, for two nanowire radii. Panel (a): First-order nonlocal SP resonance mode; panel (b): second-order nonlocal SP resonance mode.

$-(\kappa^L)^2 + k_{sp}^2 = (k^L)^2$. The nonlocal correction term Δ_L regularizes the dispersion relation in the limit $h \rightarrow 0$, and thus makes the nonlocal extinction curves for a nanowire resting on a substrate completely different from the local one.

In Figs. 8(a) and 8(b), we plot for nonlocal response the normal component of the electric field along the nanowire boundary for the first- and second-order resonance modes observed in Fig. 7. Compared to Fig. 5 for a nanowire one nanometer away from the substrate, the field distribution in Fig. 8 gained more weight on the substrate side. For example for $r_0 = 10$ nm, the field amplitude peak in Fig. 5(a) occurs near $\theta = 20^\circ$, and decreases to $\theta = 8^\circ$ in Fig. 8(a). The nanowire and the substrate increasingly influence each other as the nanowire approaches the substrate.

For the nanowire touching the substrate, Eq. (32) becomes invalid since its derivation relies on the assumption that differences between local and nonlocal response are small. However, as we will see, the trend described by Eq. (32) that ω_{res}^{nloc} blueshifts as r_0 decreases still holds true, and thinking in terms of the effective parameters l_{eff} and ϵ_b^{eff} is still useful. The argument runs as follows.

Consider two plasmonic nanowires with radii $r_1 > r_2$ and nonlocal SP resonance frequencies $\omega_{res,1}^{nloc}$ and $\omega_{res,2}^{nloc}$. Expanding the coordinate system isotropically, the nanowire with the radius r_2 could equivalently be viewed as the nanowire with the larger radius r_1 in combination with a nonlocal charge layer thickness l_{eff} that is increased by a factor of r_1/r_2 . The increased charge layer results in a stronger field inside the nanowire proportional to l_{eff} as discussed in Sec. VI. There will

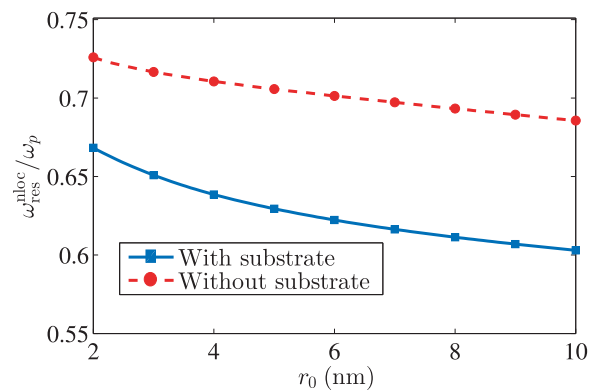


FIG. 9. (Color online) First-order resonance ω_{res}^{nloc} as a function of r_0 for of a gold cylindrical nanowire resting on a semi-infinite dielectric substrate of index 1.5, in a free-space background. The case without a substrate is also shown. The data points are numerically accurate values obtained with our nonlocal GSIM. The curves through the data points are guides to the eye.

also be a weaker screening contribution from the substrate (i.e., the ϵ_b^{eff} is smaller for the smaller wire radius) since the effective distance between the charge and the substrate is increased. These two effects make the resonance frequency $\omega_{res,2}^{nloc}$ for the smaller nanowire radius blueshifted with respect to $\omega_{res,1}^{nloc}$ also in the touching geometry. Incidentally, the latter effect agrees with Fig. 6(a1) where ϵ_b^{eff} also increases as a function of r_0 .

In Fig. 9, we show ω_{res}^{nloc} versus r_0 for the first-order nonlocal resonance. The nanowire-substrate touching geometry is compared to the case without the substrate. It is observed that ω_{res}^{nloc} blueshifts for decreasing wire radius, and more so with the substrate in place. We attribute this to the substrate-increased l_{eff} and ϵ_b^{eff} .

VIII. DISCUSSION AND CONCLUSIONS

In this paper, we generalized the local-response Green's function surface-integral method to a nonlocal version, where the nonlocal response is described by the hydrodynamical Drude model. The method developed here works for arbitrarily shaped nanowires in arbitrary inhomogeneous backgrounds. The key insight that leads to our nonlocal GSIM is that an additional surface integral can be formulated that describes the Maxwell fields associated with the hydrodynamic pressure waves. Spill-out of free electrons is neglected, so their nonlocal response can be described in terms of the fields on the surfaces that confine them. Besides Maxwell's boundary conditions, there is an additional boundary condition that is easily derived once electron spill-out is neglected.

The GSIM has the advantage of being numerically efficient, but until now it was not clear how to apply even the known local-reponse GSIM to nanoparticles resting on substrates, surely a typical situation in experiments. We showed how to apply the GSIM in this case, by regularizing the singularities that only arise for such "touching geometries," by which we mean that the surface that is to be integrated over touches an interface. This regularization procedure works both for the local and for our nonlocal GSIM. This makes the GSIM a more general-purpose numerical method, and we expect that this development will contribute to its popularity.

We expect the nonlocal GSIM also to become a method of choice when studying nonlocal response in complex geometries. Nonlocal response changes the charge distribution especially near the metal-dielectric interfaces, and it is only these interfaces that we need to discretize for the surface method. Thus nonlocal GSIM is computationally efficient and stays close to the action, so to say.

We first compared the nonlocal and local response of nanowires without substrates. We benchmarked the nonlocal GSIM against the analytical solution of the extinction of a cylindrical nanowire and found excellent agreement. We observed the characteristic nonlocal blueshift of extinction resonances. However, our finding that the blueshift is linearly proportional to the angular momentum number of the surface-plasmon resonance is new, as far as we know. We also found an analytical derivation for this phenomenon, based on the fact that nonlocal response effectively pushes the surface-charge density inward into the plasmonic nanowire. It would be interesting for future studies to study the angular-momentum dependence of resonance frequency shifts when also allowing for spill-out of the free electrons.¹⁰

For a nanowire as close as 1 nm to a dielectric substrate, we still can accurately account for the nonlocal blueshift of the resonances, using an effective theory. Besides the nanowire radius and the longitudinal wave vector, this involves an effective angular-momentum number and an effective background dielectric function. The nonlocal blueshift of a cylindrical nanowire is enhanced when close to a substrate. Our explanation can be summarized as follows: For an angular-momentum resonance of the nanowire, the substrate makes the charge distribution on the wire surface more inhomogeneous. The angular-momentum expansion of this charge distribution therefore involves higher angular-momentum numbers, and an effectively higher angular momentum can be defined with concomitant larger nonlocal blueshift. We also find that the substrate increases the effective background dielectric function. Both effects together accurately predict the enhanced nonlocal blueshift. We find that the effective background dielectric function varies by 20% when varying the nanowire radius from 2 nm to 18 nm. By contrast, the effective background dielectric function of nanospheres on a substrate was assumed to be independent of the sphere radius in Ref. 9.

We also calculated extinction spectra of nanowires resting on a dielectric surface. Pronounced differences are found between the local and the nonlocal theory, so that our effective theory for nonlocal blueshifts does not work here. Similar large differences have been predicted before for plasmonic dimers structures (two spheres,¹⁹ or two wires²⁰). Here we show that a single plasmonic nanowire on a dielectric substrate is already enough to observe considerable differences between local and nonlocal response. It may also be the preferred experimental structure to study strong nonlocal effects, since quantum tunneling as for plasmonic dimers is less of a complication.

In this paper we focused on extinction cross sections of nanowires, but also waveguiding, electron energy-loss spectroscopy, and other observables could be calculated using our method. Moreover, the method does not only work for the nanowire structures considered here. We are presently generalizing our nonlocal GSIM to truly three-dimensional geometries, where advantages of surface-integral methods are

even more pronounced. The general idea is the same, namely to add to the known surface integrals a 3D version of the surface integral for the longitudinal field.

ACKNOWLEDGMENTS

This work was financially supported by an H. C. Ørsted Fellowship (W.Y.). W.Y. would like to thank Thomas Søndergaard for introducing him to the Green's function surface-integral method. The Center for Nanostructured Graphene is sponsored by the Danish National Research Foundation, Project DNR58.

APPENDIX A: DERIVATIONS OF SURFACE INTEGRALS

1. Derivation of Eqs. (7) and (15)

Consider the area integral

$$\int_{A_i} dx dy [E_{zi}(\boldsymbol{\rho}) \nabla_{\boldsymbol{\rho}}^2 g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') - g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') \nabla_{\boldsymbol{\rho}}^2 E_{zi}(\boldsymbol{\rho})], \quad (\text{A1})$$

where $x, y \in A_i$. From the definition of the scalar Green's function g_i^T in Sec. IV A, it follows that Eq. (A1) is identical to $E_{zi}(\boldsymbol{\rho}')$. Using the identity $\phi \nabla^2 \psi = \nabla \cdot \phi \nabla \psi - \nabla \phi \cdot \nabla \psi$ and Gauss's integral theorem, Eq. (A1) can be rewritten as

$$\oint_{S_i} d\rho [E_{zi}(\boldsymbol{\rho}) \hat{n}_i(\boldsymbol{\rho}) \cdot \nabla_{\boldsymbol{\rho}} g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') - g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') \hat{n}_i(\boldsymbol{\rho}) \cdot \nabla_{\boldsymbol{\rho}} E_{zi}(\boldsymbol{\rho})]. \quad (\text{A2})$$

By equating Eq. (A2) with E_{zi} , and by using the reciprocity property that $g_i^T(\boldsymbol{\rho}, \boldsymbol{\rho}') = g_i^T(\boldsymbol{\rho}', \boldsymbol{\rho})$, one arrives at the surface integral for the metal domains Eq. (7). The additional surface integral Eq. (15) for nonlocal response in the metal can be derived analogously.

2. Derivation of Eq. (9)

When the background is spatially inhomogeneous, it is difficult to follow the same routine as used in Sec. A 1 above to derive the surface integrals for the dielectric side of the metal-dielectric boundaries. In Ref. 43, the surface integrals for the specific inhomogeneous background with the planar interface are derived by matching the boundary conditions of the Green's function at the interface. Here, we employ an alternative approach based on the surface equivalence theorem to derive the surface integrals for arbitrary backgrounds.⁴⁹ Denoting the actual field distribution as $\{\mathbf{E}_b(\boldsymbol{\rho}), \mathbf{H}_b(\boldsymbol{\rho})\}$ for $\boldsymbol{\rho} \in B$, and $\{\mathbf{E}_i(\boldsymbol{\rho}), \mathbf{H}_i(\boldsymbol{\rho})\}$ for $\boldsymbol{\rho} \in A$. Then, consider a virtual field distribution with $\{\mathbf{E}_i(\boldsymbol{\rho}), \mathbf{H}_i(\boldsymbol{\rho})\}$ replaced by $\{\mathbf{0}, \mathbf{0}\}$. The existence of such a virtual field distribution requires a set of surface currents⁴⁹

$$\tilde{\mathbf{J}}_e(\boldsymbol{\rho}) = \hat{n}(\boldsymbol{\rho}) \times \mathbf{H}_b(\boldsymbol{\rho}), \quad (\text{A3a})$$

$$\tilde{\mathbf{M}}_e(\boldsymbol{\rho}) = -\hat{n}(\boldsymbol{\rho}) \times \mathbf{E}_b(\boldsymbol{\rho}), \quad (\text{A3b})$$

existing only on the metal-dielectric boundary S , and where $\tilde{\mathbf{J}}_e$ and $\tilde{\mathbf{M}}_e$ represent the surface electric and magnetic currents, respectively. The surface currents fix the unphysical field discontinuities across the boundary. The virtual fields are equivalently a result of the fields radiated by $\tilde{\mathbf{J}}_e$, $\tilde{\mathbf{M}}_e$, and also

$\tilde{\mathbf{J}}_b$; i.e.,

$$\begin{aligned} \mathbf{E}_b(\boldsymbol{\rho}) &= \mathbf{E}_b^{\text{inc}}(\boldsymbol{\rho}) + \int d\rho' \mathbf{G}_e(\boldsymbol{\rho}, \boldsymbol{\rho}') \cdot i\omega\mu_0 \tilde{\mathbf{J}}_e(\boldsymbol{\rho}') \\ &+ \int d\rho' \mathbf{G}_e(\boldsymbol{\rho}, \boldsymbol{\rho}') \cdot [(-ik_z \hat{z} \times -\nabla_{\rho'}) \times \tilde{\mathbf{M}}_e(\boldsymbol{\rho}')], \end{aligned} \quad (\text{A4a})$$

$$\begin{aligned} \mathbf{H}_b(\boldsymbol{\rho}) &= \mathbf{H}_b^{\text{inc}}(\boldsymbol{\rho}) + \int d\rho' \mathbf{G}_m(\boldsymbol{\rho}, \boldsymbol{\rho}') \cdot i\omega\epsilon_0 \tilde{\mathbf{M}}_e(\boldsymbol{\rho}') \\ &+ \int d\rho' \mathbf{G}_m(\boldsymbol{\rho}, \boldsymbol{\rho}') \cdot [(ik_z \hat{z} \times +\nabla_{\rho'}) \times \tilde{\mathbf{J}}_e(\boldsymbol{\rho}')]. \end{aligned} \quad (\text{A4b})$$

$\mathbf{E}_b^{\text{inc}}$ and $\mathbf{H}_b^{\text{inc}}$ represent the incident fields from $\tilde{\mathbf{J}}_b$. Furthermore, \mathbf{G}_e and \mathbf{G}_m represent the background electric and magnetic dyadic Green's functions, defined by

$$[\nabla_{k_z} \times \nabla_{k_z} \times -k_0^2 \epsilon_b(\boldsymbol{\rho})] \mathbf{G}_e(\boldsymbol{\rho}, \boldsymbol{\rho}') = \mathbf{I} \delta(\boldsymbol{\rho} - \boldsymbol{\rho}'), \quad (\text{A5a})$$

$$\left[\nabla_{k_z} \times \frac{1}{\epsilon_b(\boldsymbol{\rho})} \nabla_{k_z} \times -k_0^2 \right] \mathbf{G}_m(\boldsymbol{\rho}, \boldsymbol{\rho}') = \mathbf{I} \delta(\boldsymbol{\rho} - \boldsymbol{\rho}'), \quad (\text{A5b})$$

where $\nabla_{k_z} = (\nabla_{\boldsymbol{\rho}} + ik_z \hat{z})$ and \mathbf{I} is the 3×3 unit matrix. Extracting the z component of \mathbf{E}_b and \mathbf{H}_b and taking the expressions of the surface currents into Eq. (A4), we then derive the surface integrals of Eq. (9) for the fields on the dielectric side of the metal-dielectric boundaries, valid for arbitrary spatial inhomogeneity $\epsilon_b(\boldsymbol{\rho})$ of the background.

Until now we have assumed that the metal nanowires are surrounded by a dielectric background, but let us discuss briefly how to describe the situation that there is also metal in the background, for example a metal substrate for plasmonic nanoparticles as in the recent experiments by Oulton *et al.*⁶⁴ and by Ciraci *et al.*⁶ If we neglect possible nonlocal response of the metal in the background, then the optical response of the metal can also be described by the spatially inhomogeneous but local dielectric function $\epsilon_b(\boldsymbol{\rho})$, so the above formalism can be applied.

Alternatively, if one would like to describe the metal in the background also by the hydrodynamical Drude model, then we can do this by taking ϵ_b in the dynamic Green's functions \mathbf{G}_e and \mathbf{G}_m of Eq. (A5) to be a nonlocal operator defined by Eqs. (4) and (5). In other words, we absorb possible nonlocal

effects of the background into the dynamic Green's functions \mathbf{G}_e and \mathbf{G}_m . Then, the nonlocal response is contained in the surface-integral coefficients of Eq. (10). This differs from our treatment of the plasmonic nanowires, where we decomposed the fields into the longitudinal and transverse parts. Absorbing any plasmonic nonlocal response of the background into the background Green's tensor is not just a formal trick. For example, for inhomogeneous backgrounds of a plasmonic slab substrate in free space, the corresponding surface-integral kernels for the nonlocal GSIM can be found in Appendix B.

APPENDIX B: INTEGRATION KERNELS IN EQUATION (10) FOR A LAYERED SUBSTRATE

Here we consider inhomogeneous backgrounds that can be described as substrates that are arbitrary planar multilayer systems in free space. We choose a convenient coordinate system such that the substrate of thickness t is located at $0 < x < t$, with the nanowires in the region $x < 0$. A semi-infinite substrate would correspond to $t = \infty$. The substrate consists of dielectric or metal slabs or a combination thereof. Any metallic layers can either be described with local or with nonlocal response. For all those cases, we present the integration kernels for the surface integral for the fields in the region $x < 0$ outside of the nanowires. This can be done because in the region $x < 0$ the background Green's tensors $\mathbf{G}_{e,m}(\boldsymbol{\rho}, \boldsymbol{\rho}')$ and hence the kernels in Eq. (10) can be expressed in terms of the substrate reflection coefficients at $x = 0$; only the values of these reflection coefficients are different for different metal-dielectric multilayer systems, and also different if the metals are described with local or nonlocal response. For the actual calculation of these reflection coefficients, we refer to textbooks, for example Ref. 55; for the Green's function in layered geometries, to Ref. 52; and for wave propagation in multilayer systems with nonlocal response, to Refs. 25 and 51.

We split the dyadic Green's function $\mathbf{G}_{e,m}$ into a homogeneous and a scattering part; i.e., we write $\mathbf{G}_{e,m} = \mathbf{G}_{e,m}^0 + \mathbf{G}_{e,m}^s$.⁵² Accordingly, the integration kernels in Eq. (10) are split into homogeneous and scattering parts, for example $e_b^0 = e_b^{00} + e_b^{0s}$. The homogeneous parts of the integration kernels are discussed in Sec. IV A, while the scattering parts can be derived as

$$e_b^{0s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{1}{k_x} \exp(i\psi) i\mathbf{k}_{\rho} \cdot \hat{\mathbf{n}}(\boldsymbol{\rho}') \left[-R_{\text{TE}}(k_{\parallel}) \frac{k_y^2 k_0^2}{k_{\rho}^2 k_{\parallel}^2} + R_{\text{TM}}(k_{\parallel}) \frac{k_x^2 k_z^2}{k_{\rho}^2 k_{\parallel}^2} \right], \quad (\text{B1a})$$

$$e_b^{1s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{1}{k_x} \exp(i\psi) \left[-R_{\text{TE}}(k_{\parallel}) \frac{k_y^2 k_0^2}{k_{\rho}^2 k_{\parallel}^2} + R_{\text{TM}}(k_{\parallel}) \frac{k_x^2 k_z^2}{k_{\rho}^2 k_{\parallel}^2} \right], \quad (\text{B1b})$$

$$f_b^{0s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{\omega\mu_0}{k_x} \exp(i\psi) i\mathbf{k}_{\rho} \cdot \hat{\mathbf{n}}(\boldsymbol{\rho}') \frac{k_x k_y k_z}{k_{\rho}^2 k_{\parallel}^2} [R_{\text{TE}}(k_{\parallel}) + R_{\text{TM}}(k_{\parallel})], \quad (\text{B1c})$$

$$f_b^{1s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{\omega\mu_0}{k_x} \exp(i\psi) \frac{k_x k_y k_z}{k_{\rho}^2 k_{\parallel}^2} [R_{\text{TE}}(k_{\parallel}) + R_{\text{TM}}(k_{\parallel})], \quad (\text{B1d})$$

$$h_b^{0s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{-i}{4\pi} \int dk_y \frac{\omega\epsilon_0}{k_x} \exp(i\psi) i\mathbf{k}_{\rho} \cdot \hat{\mathbf{n}}(\boldsymbol{\rho}') \frac{k_x k_y k_z}{k_{\rho}^2 k_{\parallel}^2} [R_{\text{TE}}(k_{\parallel}) + R_{\text{TM}}(k_{\parallel})], \quad (\text{B1e})$$

$$h_b^{1s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{-i}{4\pi} \int dk_y \frac{\omega \epsilon_0}{k_x} \exp(i\psi) \frac{k_x k_y k_z}{k_\rho^2 k_\parallel^2} [R_{\text{TE}}(k_\parallel) + R_{\text{TM}}(k_\parallel)], \quad (\text{B1f})$$

$$m_b^{0s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{1}{k_x} \exp(i\psi) i \mathbf{k}_\rho \cdot \hat{\mathbf{n}}(\boldsymbol{\rho}') \left[R_{\text{TE}}(k_\parallel) \frac{k_x^2 k_z^2}{k_\rho^2 k_\parallel^2} - R_{\text{TM}}(k_\parallel) \frac{k_y^2 k_0^2}{k_\rho^2 k_\parallel^2} \right], \quad (\text{B1g})$$

$$m_b^{1s}(\boldsymbol{\rho}, \boldsymbol{\rho}') = \frac{i}{4\pi} \int dk_y \frac{1}{k_x} \exp(i\psi) \left[R_{\text{TE}}(k_\parallel) \frac{k_x^2 k_z^2}{k_\rho^2 k_\parallel^2} - R_{\text{TM}}(k_\parallel) \frac{k_y^2 k_0^2}{k_\rho^2 k_\parallel^2} \right], \quad (\text{B1h})$$

where $\boldsymbol{\rho} = (x, y)$ is in the region $x < 0$, $\psi = k_y(y - y') - k_x(x + x')$, $k_0 = \omega/c$, $k_x^2 + k_y^2 + k_z^2 = k_0^2$, $k_\rho^2 = k_0^2 - k_z^2$, $k_y^2 + k_z^2 = k_\parallel^2$; R_{TE} and R_{TM} represent the reflection coefficients at $x = 0$ of the multilayer substrate for TE- and TM-polarized plane waves, respectively.

APPENDIX C: DERIVATION OF EQUATION (31)

Here we derive the effective angular momentum number l_{eff} for the SP mode supported by a cylindrically shaped plasmonic nanowire in an inhomogeneous background. This is a key parameter in our explanation of nonlocal blueshifts of nanowires in arbitrary dielectric backgrounds, especially in Eq. (32). The arguments used here are similar to those developed in Sec. VI.

Let us first consider a cylindrical nanowire with local response, in an inhomogeneous background. Define α_m as the surface charge at the nanowire boundary $r = r_0$ of the SP mode of the wire. The surface charge can be decomposed into

$$\alpha_m = \sum_{l \neq 0} \alpha_{ml} \exp(il\phi) \delta(r - r_0), \quad (\text{C1})$$

where l is the angular momentum number of the cylindrical harmonics that ranges from $-\infty$ to ∞ ; the term $l = 0$ is excluded from the summation, as it does not contribute to the surface charge. The surface charge α_m is a source that generates electric fields \mathbf{E} inside the nanowire given by

$$\mathbf{E} = \frac{1}{2\epsilon_0} \sum_{l \neq 0} \alpha_{ml} \left(\frac{r}{r_0} \right)^{l-1} \exp(il\phi) (\hat{r} + i\hat{\phi}). \quad (\text{C2})$$

Second, we consider the same structure, but now we describe the nanowire with nonlocal response. The corresponding surface charge α'_m will now effectively be moved inwards into the nanowire, to $r'_0 = r_0 - 1/k^L$, and is

expressed as

$$\alpha'_m = \sum_{l \neq 0} \alpha_{ml} \frac{r_0}{r'_0} \exp(il\phi) \delta(r - r'_0). \quad (\text{C3})$$

In the region inside r'_0 , denoted by A'_m , the charge density α'_m generates the electric field

$$\mathbf{E}' = \frac{1}{2\epsilon_0} \sum_{l \neq 0} \alpha_{ml} \frac{r_0}{r'_0} \left(\frac{r}{r'_0} \right)^{l-1} \exp(il\phi) (\hat{r} + i\hat{\phi}). \quad (\text{C4})$$

Consequently, the electric field in the area A'_m on average is enhanced by a factor

$$F = \sqrt{\frac{\int_{A'_m} d\mathbf{r} |\mathbf{E}'|^2}{\int_{A'_m} d\mathbf{r} |\mathbf{E}|^2}} \approx \left(\frac{r_0}{r'_0} \right)^{\sum_{l \neq 0} |\alpha_{ml}|^2 l^{-1} / \sum_{l \neq 0} |\alpha_{ml}|^2 l^{-1}}. \quad (\text{C5})$$

Now in a homogenous background, where the angular-momentum number corresponding to the order of the cylindrical harmonics is well defined, the factor F is equal to $(r_0/r'_0)^l$. In our effective description we now identify Eq. (C5) with $F = (r_0/r'_0)^{l_{\text{eff}}}$, which allows us to extract the effective angular momentum number l_{eff} as

$$l_{\text{eff}}^{-1} = \frac{\sum_{l \neq 0} |\alpha_{ml}|^2 l^{-1}}{\sum_{l \neq 0} |\alpha_{ml}|^2}, \quad (\text{C6})$$

which is Eq. (31) of the main text.

The effective angular momentum l_{eff} so defined is only determined by the surface distribution of the free charges and the associated electric field in the plasmonic nanowire. In other words, for l_{eff} we do not consider the electric field that can be associated with screened charges in the inhomogeneous background. Those screened charges influence the other effective parameter, namely the effective background permittivity ϵ_b^{eff} in Eq. (32).

*mwubs@fotonik.dtu.dk

¹S. A. Maier, *Plasmonics: Fundamentals and Applications* (Springer, New York, 2007).

²D. K. Gramotnev and S. I. Bozhevolnyi, *Nat. Photonics* **4**, 83 (2010).

³J. Nelayah, M. Kociak, O. Stephan, F. J. García de Abajo, M. Tence, L. Henrard, D. Taverna, I. Pastoriza-Santos, L. M. Liz-Marzan, and C. Colliex, *Nat. Phys.* **3**, 348 (2007).

⁴O. Nicoletti, M. Wubs, N. A. Mortensen, W. Sigle, P. A. van Aken, and P. A. Midgley, *Opt. Express* **19**, 15371 (2011).

⁵J. A. Scholl, A. L. Koh, and J. A. Dionne, *Nature (London)* **483**, 421 (2012).

⁶C. Ciraci, R. T. Hill, J. J. Mock, Y. Urzhumov, A. I. Fernández-Domínguez, S. A. Maier, J. B. Pendry, A. Chilkoti, and D. R. Smith, *Science* **337**, 1072 (2012).

⁷J. Kern, S. Grossmann, N. V. Tarakina, T. Häckel, M. Emmerling, M. Kamp, J.-S. Huang, P. Biagioni, J. C. Prangsma, and B. Hecht, *Nano Lett.* **12**, 5504 (2012).

⁸K. J. Savage, M. M. Hawkeye, R. Esteban, A. G. Borisov, J. Aizpurua, and J. J. Baumberg, *Nature (London)* **491**, 574 (2012).

⁹J. Scholl, A. Garcia-Etxarri, A. L. Koh, and J. A. Dionne, *Nano Lett.* **13**, 564 (2013).

- ¹⁰T. V. Teperik, P. Nordlander, J. Aizpurua, and A. G. Borisov, *Phys. Rev. Lett.* **110**, 263901 (2013).
- ¹¹F. Bloch, *Z. Phys.* **81**, 363 (1933).
- ¹²S. C. Ying, *Nuovo Cimento B* **23**, 270 (1974).
- ¹³A. Eguiluz and J. J. Quinn, *Phys. Rev. B* **14**, 1347 (1976).
- ¹⁴A. D. Boardman, *Electromagnetic Surface Modes* (John Wiley and Sons, Chichester, 1982).
- ¹⁵S. Raza, G. Toscano, A.-P. Jauho, M. Wubs, and N. A. Mortensen, *Phys. Rev. B* **84**, 121412(R) (2011).
- ¹⁶R. Ruppin, *Phys. Rev. B* **45**, 11209 (1992).
- ¹⁷R. Ruppin, *Phys. Rev. Lett.* **31**, 1434 (1973).
- ¹⁸R. Ruppin, *Opt. Commun.* **190**, 205 (2001).
- ¹⁹R. Fuchs and F. Claro, *Phys. Rev. B* **35**, 3722 (1987).
- ²⁰A. I. Fernández-Domínguez, A. Wiener, F. J. García-Vidal, S. A. Maier, and J. B. Pendry, *Phys. Rev. Lett.* **108**, 106802 (2012).
- ²¹A. Wiener, A. I. Fernández-Domínguez, A. P. Horsfield, J. B. Pendry, and S. A. Maier, *Nano Lett.* **12**, 3308 (2012).
- ²²G. Toscano, S. Raza, A.-P. Jauho, M. Wubs, and N. A. Mortensen, *Opt. Express* **13**, 4176 (2012).
- ²³S. Raza, G. Toscano, A.-P. Jauho, N. A. Mortensen, and M. Wubs, *Plasmonics* **8**, 193 (2012).
- ²⁴G. Toscano, S. Raza, S. Xiao, M. Wubs, A.-P. Jauho, S. I. Bozhevolnyi, and N. A. Mortensen, *Opt. Lett.* **37**, 2538 (2012).
- ²⁵W. L. Mochán, M. del Castillo-Mussot, and R. G. Barrera, *Phys. Rev. B* **35**, 1088 (1987).
- ²⁶F. J. García de Abajo, *J. Phys. Chem. C* **112**, 17983 (2008).
- ²⁷C. David and F. J. García de Abajo, *J. Phys. Chem. C* **115**, 19470 (2011).
- ²⁸J. E. Sipe, V. C. Y. So, M. Fukui, and G. I. Stegeman, *Phys. Rev. B* **21**, 4389 (1980).
- ²⁹C. Ciracì, E. Poutrina, M. Scalora, and D. R. Smith, *Phys. Rev. B* **85**, 201403(R) (2012).
- ³⁰A. Liebsch, *Phys. Rev. B* **48**, 11317 (1993).
- ³¹R. G. Parr and W. Yang, *Density-Functional Theory of Atoms and Molecules* (Oxford University Press, New York, 1994).
- ³²E. Townsend and G. W. Bryant, *Nano Lett.* **12**, 429 (2012).
- ³³D. Marinica, A. Kazansky, P. Nordlander, J. Aizpurua, and A. G. Borisov, *Nano Lett.* **12**, 1333 (2012).
- ³⁴R. Carmina Monreal, T. J. Antosiewicz, and S. P. Apell, *New J. Phys.* **15**, 083044 (2013).
- ³⁵J. J. Zuloaga, E. Prodan, and P. Nordlander, *Nano Lett.* **9**, 887 (2009).
- ³⁶K. S. Yee, *IEEE Trans. Antennas Propag.* **AP-14**, 302 (1966).
- ³⁷A. Taflov and S. C. Hagness, *Computational Electrodynamics: The Finite-Difference Time-Domain Method* (Artech House, Boston, 2005).
- ³⁸J. M. Jin, *The Finite Element Method in Electromagnetics* (Wiley, New York, 2002).
- ³⁹F. J. García de Abajo and A. Howie, *Phys. Rev. Lett.* **80**, 5180 (1998).
- ⁴⁰F. J. García de Abajo and A. Howie, *Phys. Rev. B* **65**, 115418 (2002).
- ⁴¹I. Romero, J. Aizpurua, G. W. Bryant, and F. J. García de Abajo, *Opt. Express* **14**, 9988 (2006).
- ⁴²T. Søndergaard and S. Bozhevolnyi, *Phys. Rev. B* **75**, 073402 (2007).
- ⁴³J. Jung and T. Søndergaard, *Phys. Rev. B* **77**, 245310 (2008).
- ⁴⁴A. M. Kern and O. J. F. Martin, *J. Opt. Soc. Am. A* **26**, 732 (2009).
- ⁴⁵B. Gallinet, A. M. Kern, and O. J. F. Martin, *J. Opt. Soc. Am. A* **27**, 2261 (2010).
- ⁴⁶K. R. Hiremath, L. Zschiedrich, and F. Schmidt, *J. Comput. Phys.* **231**, 5890 (2012).
- ⁴⁷Q. Huang, F. Bao, and S. He, *Opt. Express* **21**, 1430 (2013).
- ⁴⁸G. Toscano, S. Raza, W. Yan, C. Jeppesen, S. Xiao, M. Wubs, A.-P. Jauho, S. I. Bozhevolnyi, and N. A. Mortensen, *Nanophotonics* **2**, 161 (2013).
- ⁴⁹J. A. Kong, *Electromagnetic Wave Theory* (EMW Publishing, Cambridge, MA, 2008).
- ⁵⁰G. Barton, *Rep. Prog. Phys.* **42**, 963 (1979).
- ⁵¹W. Yan, M. Wubs, and N. A. Mortensen, *Phys. Rev. B* **86**, 205429 (2012).
- ⁵²M. S. Tomáš, *Phys. Rev. A* **51**, 2545 (1995).
- ⁵³P. Jewsbury, *J. Phys. F: Met. Phys.* **11**, 195 (1981).
- ⁵⁴G. Ford and W. Weber, *Phys. Rep.* **113**, 195 (1984).
- ⁵⁵W. C. Chew, *Waves and Fields in Inhomogeneous Media* (Van Nostrand Reinhold, New York, 1990).
- ⁵⁶S. Raza, N. Stenger, S. Kadkhodazadeh, S. V. Fischer, N. Kostesha, A.-P. Jauho, A. Burrows, M. Wubs, and N. A. Mortensen, *Nanophotonics* **2**, 131 (2013).
- ⁵⁷A. D. Boardman and B. V. Paranjape, *J. Phys. F: Met. Phys.* **7**, 1935 (1977).
- ⁵⁸J. Jung and T. G. Pedersen, *Opt. Express* **19**, 22775 (2011).
- ⁵⁹S. Zhang and H. Xu, *ACS Nano* **6**, 8128 (2012).
- ⁶⁰J. P. Kottmann and O. J. F. Martin, *Opt. Express* **8**, 655 (2001).
- ⁶¹E. Prodan, C. Radloff, N. J. Halas, and P. Nordlander, *Science* **302**, 419 (2003).
- ⁶²D. W. Brandl, C. Oubre, and P. Nordlander, *J. Chem. Phys.* **123**, 024701 (2005).
- ⁶³T. J. Davis, D. E. Gómez, and K. C. Vernon, *Nano. Lett.* **10**, 2618 (2010).
- ⁶⁴R. F. Oulton, V. J. Sorger, T. Zentgraf, R.-M. Ma, C. Gladden, L. Dai, G. Bartal, and X. Zhang, *Nature (London)* **461**, 629 (2009).