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Theory of electron energy-loss spectroscopy in atomically thin metallic films

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We study strongly confined plasmons in ultrathin gold and silver films by simulating electron energy-loss spectroscopy. Plasmon dispersion relations are directly retrieved from the energy- and momentum-resolved loss probability under normal incidence conditions, whereas they can also be inferred from aloof parallel beam trajectories from the evolution of the plasmon features in the resulting loss spectra as we vary the impinging electron energy. We find good agreement between nonlocal quantum-mechanical simulations based on the random-phase approximation and a local classical dielectric description for silver films of different thicknesses down to a few atomic layers. We further observe only a minor dependence of quantum simulations for these films on the confining out-of-plane electron potential when comparing density-functional theory within the jellium model with a phenomenological experimentally fitted potential incorporating atomic layer periodicity and in-plane parabolic bands of energy-dependent effective mass. The latter shows also a small dependence on the crystallographic orientation of silver films, while the unphysical assumption of energy-independent electron mass leads to spurious features in the predicted spectra. Interestingly, we find electron band effects to be more relevant in gold films, giving rise to blueshifts when compared to classical or jellium model simulations. In contrast to the strong nonlocal effects found in few-nanometer metal nanoparticles, our study reveals that a local classical description provides excellent quantitative results in both plasmon strength and dispersion when compared to quantum-mechanical simulations down to silver films consisting of only a few atomic layers, thus emphasizing the in-plane nearly free conduction-electron motion associated with plasmons in these structures.

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I. INTRODUCTION

Surface plasmons—the collective electron oscillations at material surfaces and interfaces—provide the means to concentrate and amplify the intensity of externally applied light down to nanoscale regions [1,2], where they interact strongly with molecules and nanostructures, thus becoming a powerful asset in novel applications [3] such as biosensing [2,4,5], photocatalysis [6,7], energy harvesting [8,9], and nonlinear optics [10–13].

Surface plasmons were first identified using electron energy-loss spectroscopy (EELS), starting with the prediction [14] and subsequent measurement of associated loss features in electrons scattered under grazing incidence from Al [15], Na and K [16], and Ag [17,18] surfaces. The main characteristics of surface plasmons in noble and simple metals were successfully explained using time-dependent density-functional theory (TD-DFT) [19] within the jellium model [20,21], while inclusion of electron band effects was required

for other metals [22]. Interestingly, multipole surface plasmons were predicted as additional resonances originating in the smooth electron density profile across metal-dielectric interfaces [22–24] and subsequently found in experiments performed on simple metals such as K and Na [25], but concluded to be too weak to be observed in Al [25] and Ag [26]. These studies focused on the relatively high-energy plasmons supported by planar surfaces in the short-wavelength regime. However, plasmons can hybridize with light forming surface-plasmon polaritons (SPPs) in planar surfaces, which become lightlike modes at low energies, thus losing confinement, as they are characterized by in-plane wavelengths slightly smaller than those of light and long-range penetration into the dielectric material or empty space outside the metal [27–29].

Highly confined plasmons can also be achieved in sharp metallic tips and closely spaced metal surfaces [30], where strong redshifts are produced due to the attractive Coulomb interaction between neighboring noncoplanar interfaces. This effect, which depends dramatically on surface morphology, can also be observed in planar systems such as ultrathin noble metal films [31,32] and narrow metal-dielectric-metal waveguides [12,31,33]. More precisely, hybridization takes place in metal films between the plasmons supported by their two interfaces, giving rise to bonding and antibonding dispersion branches that were first revealed also through EELS in self-standing aluminum foils [34]; in ultrathin films of only a few atomic layers in thickness, the antibonding

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plasmon dispersion is pushed close to the light line, whereas the bonding plasmon becomes strongly confined (reaching the quasistatic limit [31,32]), as experimentally corroborated through angle-resolved low-energy EELS in few-monolayer Ag films [35] and monolayer (ML) DySi₂ [36], as well as in laterally confined wires formed by In [37] and silicide [38], and even in monatomic Au chains grown on Si(557) surfaces [39]. Additionally, graphene has been shown to support long-lived mid-infrared and terahertz plasmons [40] that can be tuned electrically [41,42] and confined vertically down to a few nanometers when placed in close proximity to a planar metal surface [43,44]. While most graphene plasmon studies have been performed using far- and near-field optics setups [45–47], low-energy EELS has also revealed their dispersion relation in extended films [48,49]. Here, we focus instead on visible and near-infrared plasmons supported by atomically thin metal films, which have been recently demonstrated in laterally patterned crystalline Ag layers [32], where they also experience strong spatial confinement.

In this paper, we investigate plasmons in atomically thin noble metal films by theoretically studying EELS for electron beams either traversing them or moving parallel outside their surface, and in this way accessing different regions of the dispersion diagram. Specifically, we explore nonlocal and quantum-confinement effects in the frame of the random-phase approximation (RPA). Our results are in excellent agreement with classical dielectric theory based on the use of frequency-dependent dielectric functions for both Ag and Au films of small thickness down to a few atomic layers. However, this conclusion is in stark contrast to the strong nonlocal effects observed in metal nanoparticles of similar or even larger diameter [50,51], a result that we attribute to the predominance of in-plane electron motion associated with the low-energy plasmons of thin films, unlike the combination of in- and out-of-plane motion in higher-energy SPPs. We also provide a comprehensive description of the influence of various quantum-mechanical effects, such as the variation of the effective mass in the dispersion relation of electronic quantum-well states in the thin films, the detailed dependence on the model used to describe valence electrons, and the inclusion of electron spillout and surface orientation.

II. THEORETICAL FORMALISM

We present the elements needed to calculate EELS probabilities in the nonretarded approximation using the linear response susceptibility to represent the metallic thin film. The latter is obtained in the RPA, starting from the one-electron wave functions of the system, which are organized as vertical quantum-well (QW) states, discretized by confinement along the out-of-plane direction and exhibiting quasifree motion along the plane of the film. We further specify the EELS probability for electron trajectories either parallel or perpendicular with respect to the metal surfaces.

A. Calculation of EELS probabilities from the susceptibility in the nonretarded limit

The loss probability $\Gamma^{\text{EELS}}(\omega)$ measured through EELS in electron microscopes must be normalized in such a way

that $\int_0^\infty d\omega \hbar\omega \Gamma^{\text{EELS}}(\omega)$ gives the average energy loss experienced by the electrons. Taking the latter to follow a straight-line trajectory with constant velocity vector \mathbf{v} parallel to the z axis and impact parameter $\mathbf{R}_0 = (x_0, y_0)$, we can write [52]

$$\Gamma^{\text{EELS}}(\omega) = \frac{e}{\pi \hbar \omega} \int dz \text{Re}\{E_z^{\text{ind}}(\mathbf{R}_0, z, \omega) e^{-i\omega z/v}\} \quad (1)$$

as the integral along the electron trajectory of the frequency-resolved self-induced field $E_z^{\text{ind}}(\mathbf{r}, \omega) = \int dt E_z^{\text{ind}}(\mathbf{r}, t) e^{i\omega t}$, which can be in turn calculated by solving the classical Maxwell equations with the electron point charge acting as an external source in the presence of the sample. This equation is rigorously valid within the approximations of linear response and nonrecoil (i.e., small energy loss $\hbar\omega$ compared with the electron kinetic energy E_0).

In the present study, we consider relatively small electron velocities $v \ll c$ and films of small thickness compared with the involved optical wavelengths. This allows us to work in the quasistatic limit and write the field $E_z^{\text{ind}}(\mathbf{r}, \omega) = -\partial_z \phi^{\text{ind}}(\mathbf{r}, \omega)$ as the gradient of a scalar potential, so Eq. (1) can be integrated by parts to yield

$$\Gamma^{\text{EELS}}(\omega) = \frac{e}{\pi \hbar v} \int dz \text{Im}\{\phi^{\text{ind}}(\mathbf{R}_0, z, \omega) e^{-i\omega z/v}\}. \quad (2)$$

We can now express the induced potential in terms of the induced charge as

$$\phi^{\text{ind}}(\mathbf{r}, \omega) = \int d^3 \mathbf{r}' v(\mathbf{r}, \mathbf{r}') \rho^{\text{ind}}(\mathbf{r}', \omega), \quad (3)$$

where $v(\mathbf{r}, \mathbf{r}')$ is the Coulomb interaction between point charges located at positions \mathbf{r} and \mathbf{r}' . Likewise, we write the induced charge as $\rho^{\text{ind}}(\mathbf{r}, \omega) = \int d^3 \mathbf{r}' \chi(\mathbf{r}, \mathbf{r}', \omega) \phi^{\text{ext}}(\mathbf{r}', \omega)$, where $\chi(\mathbf{r}, \mathbf{r}', \omega)$ is the linear susceptibility, $\phi^{\text{ext}}(\mathbf{r}, \omega) = \int d^3 \mathbf{r}' v(\mathbf{r}, \mathbf{r}') \rho^{\text{ext}}(\mathbf{r}', \omega)$ is the external electric potential generated by the electron charge density $\rho^{\text{ext}}(\mathbf{r}, \omega) = -e \int dt \delta(\mathbf{r} - \mathbf{R}_0 - \mathbf{v}t) e^{i\omega t} = (-e/v) \delta(\mathbf{R} - \mathbf{R}_0) e^{i\omega z/v}$, and we use the notation $\mathbf{r} = (\mathbf{R}, z)$ with $\mathbf{R} = (x, y)$.

In free space one has $v(\mathbf{r}, \mathbf{r}') = v_0(\mathbf{r} - \mathbf{r}') = 1/|\mathbf{r} - \mathbf{r}'|$, but we are interested in retaining a general spatial dependence of $v(\mathbf{r}, \mathbf{r}')$ in order to describe the polarization background produced in the film by interaction with everything else other than conduction electrons (see below). Combining these elements with Eq. (2), we find the loss probability

$$\Gamma^{\text{EELS}}(\omega) = \frac{e^2}{\pi \hbar v^2} \int d^3 \mathbf{r} \int d^3 \mathbf{r}' w^*(\mathbf{r}) w(\mathbf{r}') \times \text{Im}\{-\chi(\mathbf{r}, \mathbf{r}', \omega)\}, \quad (4)$$

where

$$w(\mathbf{r}) = \int dz' v(\mathbf{r}, \mathbf{R}_0, z') e^{i\omega z'/v} \quad (5)$$

is the *external* potential created by the electron and we have made use of the reciprocity property $\chi(\mathbf{r}, \mathbf{r}', \omega) = \chi(\mathbf{r}', \mathbf{r}, \omega)$ to extract the complex factors w outside the imaginary part. Next, we apply this expression to calculate EELS probabilities from the RPA susceptibility. But first, for completeness, we note that the integral in Eq. (5) can be performed analytically for the bare Coulomb interaction [53] yielding

$$w(\mathbf{r}) = 2K_0(\omega|\mathbf{R} - \mathbf{R}_0|/v) e^{i\omega z/v},$$

where K_0 is a modified Bessel function [53], thus allowing us to write

$$\begin{aligned} \Gamma^{\text{EELS}}(\omega) &= \frac{4e^2}{\pi\hbar v^2} \int d^3\mathbf{r} \int d^3\mathbf{r}' \cos\left[\frac{\omega}{v}(z' - z)\right] \\ &\times K_0\left(\frac{\omega}{v}|\mathbf{R} - \mathbf{R}_0|\right) K_0\left(\frac{\omega}{v}|\mathbf{R}' - \mathbf{R}_0|\right) \\ &\times \text{Im}\{-\chi(\mathbf{r}, \mathbf{r}', \omega)\} \end{aligned}$$

for the loss probability, which we can directly apply to systems in which any background polarization is already contained in χ , or when v is well described by the bare Coulomb interaction (e.g., in simple metals).

B. RPA susceptibility of thin metal films

We follow the same formalism as in Ref. [33], which is extended here to account for an energy dependence of the in-plane electron effective mass. One starts by writing $\chi(\mathbf{r}, \mathbf{r}', \omega)$ in terms of the noninteracting susceptibility $\chi^0(\mathbf{r}, \mathbf{r}', \omega)$ through $\chi = \chi^0 \cdot (\mathcal{I} - v \cdot \chi^0)^{-1}$, where we use matrix notation with spatial coordinates \mathbf{r} and \mathbf{r}' acting as matrix indices, so that matrix multiplication involves integration over \mathbf{r} , and $\mathcal{I}(\mathbf{r}, \mathbf{r}') = \delta(\mathbf{r} - \mathbf{r}')$. We further adopt the RPA by calculating χ^0 as [33,54]

$$\chi^0(\mathbf{r}, \mathbf{r}', \omega) = \frac{2e^2}{\hbar} \sum_{ii'} (f_{i'} - f_i) \frac{\psi_i(\mathbf{r})\psi_i^*(\mathbf{r}')\psi_{i'}^*(\mathbf{r})\psi_{i'}(\mathbf{r}')}{\omega + i\gamma - (\varepsilon_i - \varepsilon_{i'})} \quad (6)$$

from the one-electron wave functions ψ_i of energies $\hbar\varepsilon_i$ and Fermi-Dirac occupation numbers f_i . Here, the factor of 2 accounts for spin degeneracy and γ is a phenomenological damping rate.

We describe metal films assuming translational invariance along the in-plane directions and parabolic electron dispersion with different effective mass m_j^* for each vertical QW band j . This allows us to write the electron wave functions as [55] $\psi_i(\mathbf{r}) = \varphi_j(z)e^{i\mathbf{k}_\parallel \cdot \mathbf{R}}/\sqrt{A}$, where \mathbf{k}_\parallel is the two-dimensional (2D) in-plane wave vector, A is the quantization area, and the state index is multiplexed as $i \rightarrow (j, \mathbf{k}_\parallel)$. Likewise, the electron energy can be separated as $\hbar\varepsilon_{j,\mathbf{k}_\parallel} = \hbar\varepsilon_j^\perp + \hbar^2 k_\parallel^2/2m_j^*$, where $\hbar\varepsilon_j^\perp$ is the out-of-plane energy that signals the QW band bottom (relative to the vacuum level). Inserting these expressions into Eq. (6) and making the customary substitution $\sum_i \rightarrow A \sum_j \int d^2\mathbf{k}_\parallel/(2\pi)^2$ for the state sums, we find [56]

$$\chi(\mathbf{r}, \mathbf{r}', \omega) = \int \frac{d^2\mathbf{Q}}{(2\pi)^2} \chi(Q, z, z', \omega) e^{i\mathbf{Q} \cdot (\mathbf{R} - \mathbf{R}')}, \quad (7)$$

which directly reflects the in-plane homogeneity of the film. We can now work in \mathbf{Q} space, where Eq. (6) reduces, using the above assumptions for the wave functions, to

$$\chi^0(Q, z, z', \omega) = \frac{2e^2}{\hbar} \sum_{jj'} \chi_{jj'}(Q, \omega) \varphi_j(z) \varphi_j^*(z') \varphi_{j'}^*(z) \varphi_{j'}(z'), \quad (8)$$

where

$$\begin{aligned} \chi_{jj'}(Q, \omega) &= \int \frac{d^2\mathbf{k}_\parallel}{(2\pi)^2} (f_{j',|\mathbf{k}_\parallel - \mathbf{Q}|} - f_{j,\mathbf{k}_\parallel}) \\ &\times \frac{1}{\omega + i\gamma - [\varepsilon_j^\perp - \varepsilon_{j'}^\perp + \frac{\hbar}{2}(k_\parallel^2/m_j^* - |\mathbf{k}_\parallel - \mathbf{Q}|^2/m_{j'}^*)]}, \end{aligned} \quad (9)$$

which only depends on the modulus of \mathbf{Q} due to the in-plane band isotropy. We evaluate the integral in Eq. (9) assuming zero temperature [i.e., $f_{j,\mathbf{k}_\parallel} = \theta(E_F - \hbar\varepsilon_{j,\mathbf{k}_\parallel})$, where E_F is the Fermi energy] and taking $\mathbf{Q} = (Q, 0)$ without loss of generality.

Incidentally, simple manipulations of the above expressions reveal a dependence on frequency and damping through $(\omega + i\gamma)^2$ that is maintained in the local limit ($Q \rightarrow 0$), in contrast to $\omega(\omega + i\gamma)$ in the Drude model. The RPA formalism thus produces spectral features with roughly twice the width of the Drude model in the local limit. This problem (along with a more involved issue related to local conservation of electron number for finite attenuation) can be solved through a phenomenological prescription proposed by Mermin [57], which unfortunately becomes rather involved when applied to the present systems. As a practical and reasonably accurate solution, we proceed instead by setting $\gamma = \gamma^{\text{exp}}/2$ in the above expressions (i.e., half the experimental damping rate; see the Appendix).

We obtain the out-of-plane wave functions $\varphi_j(z)$ as the eigenstates of the one-dimensional Hamiltonian $-(\hbar^2/2m_e)\partial_{zz} + V(z)$, using the free-electron mass m_e for the transversal kinetic term and two different models for the confining potential $V(z)$: (i) the self-consistent solution in the jellium (JEL) approximation within density-functional theory (DFT) [20,21] and (ii) a phenomenological atomic-layer potential (ALP) that incorporates out-of-plane bulk atomic-layer corrugation and a surface density profile with parameters fitted to reproduce relevant experimental band-structure features, such as affinity, surface-state energy, and projected bulk band gap, which depend on material and crystal orientation as compiled in Ref. [58].

The JEL model corresponds to the self-consistent DFT solution for a thin slab of background potential and energy-independent effective mass $m_j^* = m_e$ [20,21], computed here through an implementation discussed elsewhere [59].

In the ALP model we fit m_j^* to experimental data (see Table I) and consider an effective electron density n_{eff} . Upon integration over the density of states of the parabolic QW bands, we can then write the Fermi energy of an N -layer film as

$$E_F = \left(\sum_{j=1}^M m_j^* \right)^{-1} \left(n_{\text{eff}} a_s N \hbar^2 \pi + \sum_{j=1}^M m_j^* \hbar \varepsilon_j^\perp \right), \quad (10)$$

where $j = M$ is the highest partially populated QW band (i.e., $\varepsilon_M^\perp < E_F/\hbar < \varepsilon_{M+1}^\perp$) and a_s is the atomic interlayer spacing [i.e., the film thickness is $d = Na_s$, with $a_s = 0.236$ nm for Ag(111) and Au(111), and $a_s = 0.205$ nm for Ag(100)]. This expression reduces to a similar one in Ref. [33] when m_j^* is independent of j . We adjust n_{eff} for each type of metal surface in such a way that Eq. (10) gives the experimental bulk values

TABLE I. Parameters used to describe the parabolic dispersion of quantum wells in Ag(111), Ag(100), and Au(111) films. We take the effective mass of each QW j to linearly vary as $m_j^*/m_e = a\hbar\varepsilon_j^\perp + b$ with band-bottom energy $\hbar\varepsilon_j^\perp$, where the parameters a and b are taken to match m_0 at the highest occupied QW [below the surface state (SS)] in the semi-infinite surface and $m^* = m_e$ at the bottom of the conduction band. The effective electron density n_{eff} , given here relative to the bulk conduction electron density n_0 , is required to fit the experimentally observed Fermi energy E_F (relative to the vacuum level) and SS energy.

Material	a (eV $^{-1}$)	b	$m^*(\text{SS})/m_e$	m_0/m_e	n_{eff}/n_0	E_F (eV)
Ag(111)	−0.1549	−0.5446	0.40 [60]	0.25 [61]	0.8381	−4.63 [62,64]
Ag(100)	−0.0817	0.2116		0.40 [63]	0.8710	−4.43 [58]
Au(111)	−0.1660	−0.8937	0.26 [60]	0.21 [61]	0.9443	−5.50 [62,64]

of E_F listed in Table I. Incidentally, although the effective mass of surface states also varies with energy [65,66], we take it as constant because of the lack of data for ultrathin Au and Ag films; this should be a reasonable approximation for films consisting of $N \geq 5$ layers, where the surface-state energy is already close to the semi-infinite surface level.

Conduction electrons interact through the bare Coulomb potential in simple metals, which in \mathbf{Q} space reduces to $v(\mathbf{Q}, z, z') = (2\pi/Q)e^{-Q|z-z'|}$. However, polarization of inner electronic bands plays a major role in the dielectric response of Ag and Au. We describe this effect by modifying $v(\mathbf{Q}, z, z')$ in order to account for the interaction between point charges in the presence of a dielectric slab of local background permittivity fitted to experimental data [67] after subtracting a Drude term representing conduction electrons (see the Appendix). We thus adopt the local response approximation for this contribution originating in localized inner electron states, whereas conduction electrons are treated nonlocally through the above RPA formalism. Similar to Eq. (7), translational symmetry in the film allows us to write

$$v(\mathbf{r}, \mathbf{r}') = \int \frac{d^2\mathbf{Q}}{(2\pi)^2} v(\mathbf{Q}, z, z') e^{i\mathbf{Q}\cdot(\mathbf{R}-\mathbf{R}')}, \quad (11)$$

where $v(\mathbf{Q}, z, z')$ is reproduced for convenience from Ref. [33] in the Appendix. We note that Eq. (11) neglects the effect of lateral atomic corrugation in this interaction (i.e., the background permittivity is taken to be homogeneous inside the film).

Finally, we calculate $\chi(\mathbf{Q}, z, z', \omega)$ from the noninteracting susceptibility [Eq. (8)] and the screened interaction by discretizing both of them in real space coordinates (z, z') and numerically performing the linear matrix algebra explained above. We obtain converged results with respect to the number of discretization points and also compared with an expansion in harmonic functions [33].

C. EELS probability under normal incidence

Direct insertion of Eqs. (7) and (11) into Eqs. (4) and (5) leads to the result

$$\Gamma_{\perp}^{\text{EELS}}(\omega) = \int_0^{\infty} dQ \Gamma_{\perp}^{\text{EELS}}(Q, \omega) \quad (12)$$

with

$$\begin{aligned} \Gamma_{\perp}^{\text{EELS}}(Q, \omega) &= \frac{e^2 Q}{2\pi^2 \hbar v^2} \int dz \int dz' I_{\perp}^*(Q, z) I_{\perp}(Q, z') \text{Im}\{-\chi(Q, z, z', \omega)\}, \end{aligned} \quad (13)$$

where

$$I_{\perp}(Q, z) = \int dz' v(\mathbf{Q}, z, z') e^{i\omega z'/v} \quad (14)$$

contains the external electron potential. For completeness, we note that when $v(\mathbf{Q}, z, z')$ is the bare Coulomb interaction $(2\pi/Q)e^{-Q|z-z'|}$, Eq. (14) becomes $I_{\perp}(Q, z) = 4\pi e^{i\omega z/v}/(Q^2 + \omega^2/v^2)$, so Eq. (13) reduces to

$$\begin{aligned} \Gamma_{\perp}^{\text{EELS}}(Q, \omega) &= \frac{8e^2}{\hbar v^2} \frac{Q}{(Q^2 + \omega^2/v^2)^2} \\ &\times \int dz \int dz' \cos[\omega(z-z')/v] \\ &\times \text{Im}\{-\chi(Q, z, z', \omega)\}, \end{aligned} \quad (15)$$

where we have used reciprocity again [i.e., $\chi(Q, z, z', \omega) = \chi(Q, z', z, \omega)$].

In the simulations that we present below, we compare the RPA approach just presented with classical electromagnetic calculations based on the use of a local frequency-dependent dielectric function for the metal. This configuration has been theoretically studied for a long time [68], and in particular, we use the analytical expressions derived in a previous publication for an electron normally incident on a dielectric slab [69] with the bulk contribution integrated up to a cutoff wave vector $Q = 5 \text{ nm}^{-1}$.

D. EELS probability in the aloof configuration

For an electron moving parallel to the film at a distance z_0 from the metal surface, it is convenient to make the substitutions $z \rightarrow x$, $\mathbf{R} \rightarrow (y, z)$, and $\mathbf{R}_0 \rightarrow (0, z_0)$ in Eqs. (4) and (5), so combining them with Eqs. (7) and (11), and retaining $\mathbf{R} = (x, y)$ in the latter, we readily obtain

$$\begin{aligned} \Gamma_{\parallel}^{\text{EELS}}(\omega) &= \frac{e^2 L}{\pi^2 \hbar v^2} \int_0^{\infty} dQ_y \int dz \int dz' v^*(Q, z, z_0) \\ &\times v(Q, z', z_0) \text{Im}\{-\chi(Q, z, z', \omega)\}, \end{aligned} \quad (16)$$

where $Q = \sqrt{\omega^2/v^2 + Q_y^2}$ and L is the electron path length. Again for completeness, when $v(\mathbf{Q}, z, z')$ is the bare Coulomb

interaction, Eq. (16) reduces to

$$\Gamma_{\parallel}^{\text{EELS}}(\omega) = \frac{4e^2L}{\hbar v^2} \int_0^{\infty} \frac{dQ_y}{Q^2} \times \int dz \int dz' e^{-Q(|z-z_0|+|z'-z_0|)} \times \text{Im}\{-\chi(Q, z, z', \omega)\}.$$

The above expressions can be applied to electron impact parameters z_0 both inside and outside the metal, but they can be simplified when the beam is not overlapping the conduction electron charge [see Fig. 3(a)], so that $z_0 > z, z'$ lies in the region inside the above integrals in which $\chi(Q, z, z', \omega)$ is zero, and therefore, changing the variable of integration from Q_y to Q , we can write

$$\Gamma_{\parallel}^{\text{EELS}}(\omega) = \frac{2e^2L}{\pi \hbar v^2} \int_{\omega/v}^{\infty} dQ \frac{e^{-2Qz_0}}{\sqrt{Q^2 - \omega^2/v^2}} \text{Im}\{r_p(Q, \omega)\}, \quad (17)$$

where

$$r_p(Q, \omega) = -\frac{Q}{2\pi} \int dz \int dz' v^*(Q, z, z_0) v(Q, z', z_0) \times e^{2Qz_0} \chi(Q, z, z', \omega) \quad (18)$$

is the Fresnel reflection coefficient of the film for p polarization (i.e., with the magnetic field parallel to the surface) in the quasistatic limit. Incidentally, Eq. (18) is independent of the source location z_0 when it does not overlap the metal because $v(Q, z, z_0)$ then depends on z_0 only through a factor e^{-Qz_0} (see the Appendix). Equation (17), which agrees with previous derivations from classical dielectric theory [70], reveals $\text{Im}\{r_p(Q, \omega)\}$ as a loss function, which is used below to visualize the surface plasmon dispersion. We also provide results from a local dielectric description based on the textbook solution of the Poisson equation for the reflection coefficient [33],

$$r_p^{\text{classical}} = \frac{(\epsilon^2 - 1)(1 - e^{-2Qd})}{(\epsilon + 1)^2 - (\epsilon - 1)^2 e^{-2Qd}}, \quad (19)$$

for a metal film of thickness d and permittivity ϵ . The poles of the reflection coefficient lead to the bonding plasmon dispersion relation $\coth(Qd/2) = \omega_p^2/\omega^2 - \epsilon_b$, which depends on the background permittivity ϵ_b and the plasma frequency $\omega_p = \sqrt{4\pi n_0 e^2/m_e}$. These parameters are in turn dependent on the choice of material, and in particular, the conduction-electron density n_0 indicates the number of electrons that participate in the plasmonic excitations, so they strongly affect the plasmon dispersion relation, the frequency of which is proportional to $\sqrt{n_0}$.

III. RESULTS AND DISCUSSION

We show examples of the two types of confining electron potentials used in our RPA calculations for Ag films in Figs. 1(a) and 1(b), along with the resulting conduction-electron charge densities. The JEL potential is smooth at the surface and describes electron spillover and Friedel oscillations [71]. The phenomenological ALP further incorporates corrugations due to the atomic planes in the bulk, which result in strong oscillations of the density. The computed

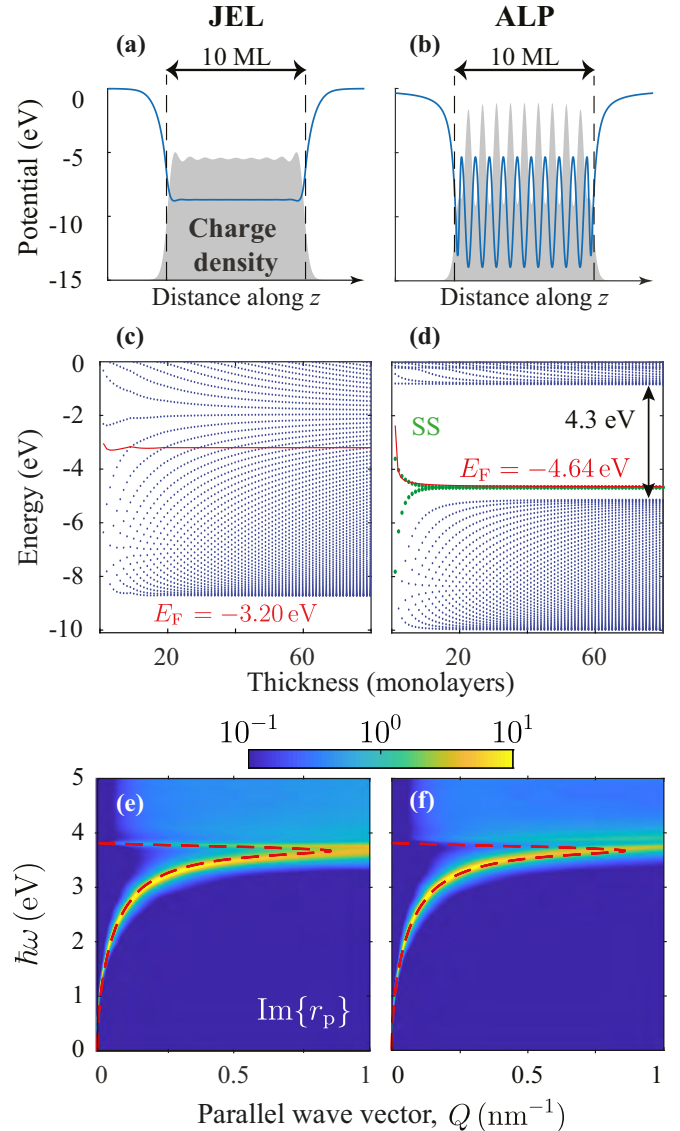


FIG. 1. RPA description of plasmons in atomically thin Ag(111) films. (a, b) Effective confining potential for conduction electrons across a 10-ML film. The conduction charge density is shown as shaded areas. (d, c) Electronic energies as a function of film thickness expressed as the number of (111) atomic layers (blue dots). Red curves and green dots represent the Fermi energy and the surface states (SSs). (e, f) Loss function $\text{Im}\{r_p\}$ calculated in the RPA [color plot, Eq. (18)], compared with the plasmon dispersion relation in the local Drude dielectric model (red curves). Left (right) panels are calculated in the jellium (ALP) model.

electron energies $\hbar\epsilon_j$ (see Sec. II B), which correspond to the bottom points of the QW bands (i.e., for vanishing in-plane momentum), are distributed with N of them below the Fermi level in a Ag(111) film consisting of N monolayers [Figs. 1(c) and 1(d)] [72]. The band structure quickly evolves toward the semi-infinite surface for a few tens of MLs in both models. Additionally, the ALP hosts surface states and a projected bulk gap of energies fitted to experiment [58]. We note that this gap depends on surface orientation: it is present in Ag(111) but absent in Ag(100) at the Fermi

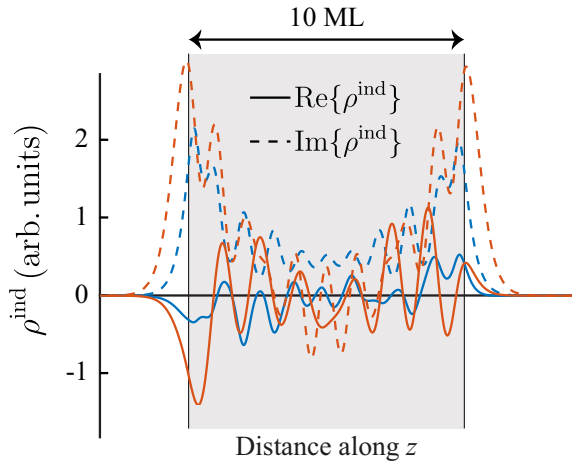


FIG. 2. Plasmon charge density across a thin 10-ML Ag(111) film. We plot the real (solid curves) and imaginary (dashed curves) parts of the induced charge density ρ^{ind} as calculated in the RPA for excitation by a source placed to the left of the film at the plasmon energies $\hbar\omega = 3.54$ eV and $\hbar\omega = 3.47$ eV corresponding to a parallel wave vector $Q = 0.5$ nm $^{-1}$ in the ALP (blue) and JEL (orange) models, respectively.

level, as revealed by photoemission measurements [73] (see also Fig. S1(a) in the Supplemental Material (SM) [74]). Remarkably, despite the important differences in the details of the potentials and electron bands, both models predict a similar plasmon dispersion [Figs. 1(e) and 1(f), density plots, obtained from Eq. (18)], which is in excellent agreement with classical theory [Figs. 1(e) and 1(f), red curves, obtained from the poles of Eq. (19)]. This is mainly attributed to the fact that the dispersion relation is dominated by the background dielectric function and the number of electrons undergoing the plasmonic excitation. Incidentally, we observe the response to also converge toward the semi-infinite surface limit for a few tens of atomic layers (see Fig. S2 in the SM [74]). Similar good agreement is found in the reflection coefficients of Ag films computed for different thickness with either of these potentials, with a square-barrier potential, or with a model potential constructed by gluing on either film side a jellium DFT potential tabulated for semi-infinite surfaces [20] (see Fig. S3 in the SM [74]).

The transversal distribution of charge densities associated with thin-film plasmons shows a clear resemblance when calculated using the ALP or JEL model potentials, although one can still observe substantial discrepancies between the two of them (see for example Fig. 2, where the ALP model charge appears to be smaller in magnitude). However, this different behavior hardly reflects in the dispersion relation and plasmon strength (Fig. 1). Interestingly, the z -integrated charge is nonzero, revealing that plasmons involve net charge oscillations along the in-plane directions for finite wave vector Q .

We conclude from these results that it is the effective number of valence electrons participating in the plasmons that determines their main characteristics, irrespective of the details of the electron wave functions and induced charge densities. We further note that a classical description of EELS

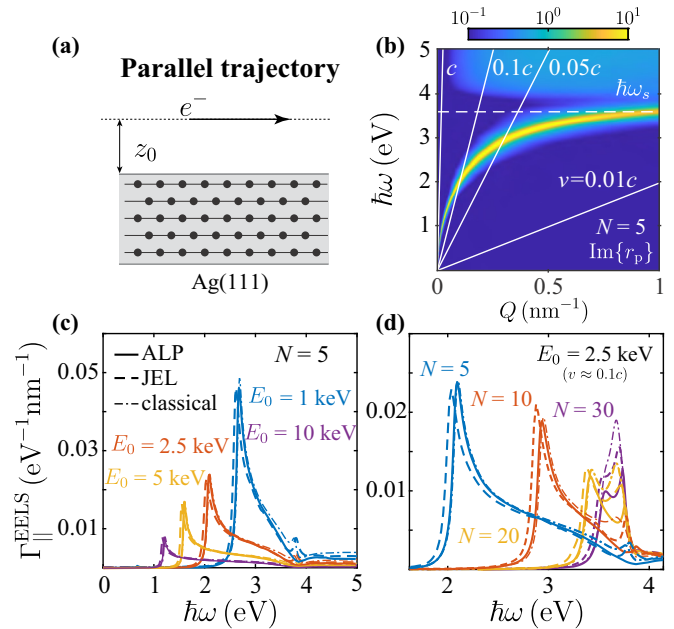


FIG. 3. Alooof EELS in thin Ag(111) films. (a) Scheme showing an electron moving parallel to an $N = 5$ ML Ag(111) metal film at a distance z_0 from its upper surface. (b) Dispersion diagram showing $\text{Im}\{r_p\}$ calculated in the ALP model for the film shown in (a). White solid lines correspond to $\omega = vQ$ for different velocities v , while the dashed horizontal line shows the classical high- Q asymptotic surface-plasmon energy $\hbar\omega_s \simeq 3.7$ eV. (c, d) EELS probability per unit of path length for $z_0 = 0.5$ nm calculated using different models [see legend in (c)] for (c) different electron kinetic energies E_0 with fixed $N = 5$ and (d) different N 's with $E_0 = 2.5$ keV.

is already in good agreement with experimental observation [69], and in particular for silver films [17,18].

The loss function $\text{Im}\{r_p\}$ provides a convenient way to represent the plasmon dispersion relation, as plasmons produce sharp features in the Fresnel reflection coefficient for p polarization. A weighted integral of this quantity over in-plane wave vectors gives the EELS probability under parallel alooof interaction [Fig. 3(a)] according to Eq. (17). However, the integration limit has a threshold at $\omega = Qv$ and the weighting factor multiplying the loss function in the integrand diverges precisely at that point. The cutoff condition $\omega = Qv$ is represented in Fig. 3(b) for different electron velocities (white lines) along with the loss function (density plot). As expected, the points of intersection with the plasmon band produce a dominant contribution that pops up as sharp peaks in the resulting EELS spectra [Figs. 3(c) and 3(d)]. An increase in electron velocity (i.e., in the slope of the threshold line) results in a redshift of the spectral peak [Fig. 3(c)], and likewise, thinner films show plasmons moving farther away from the $\omega = Qc$ light line, thus producing shifts toward lower plasmon energies in the EELS spectra for fixed electron energy. We remark that RPA and classical calculations lead to quantitatively similar results for this configuration, and the former are roughly independent of the choice of confining electron potential.

The ALP model incorporates experimental information on electronic bands, which depend on crystallographic

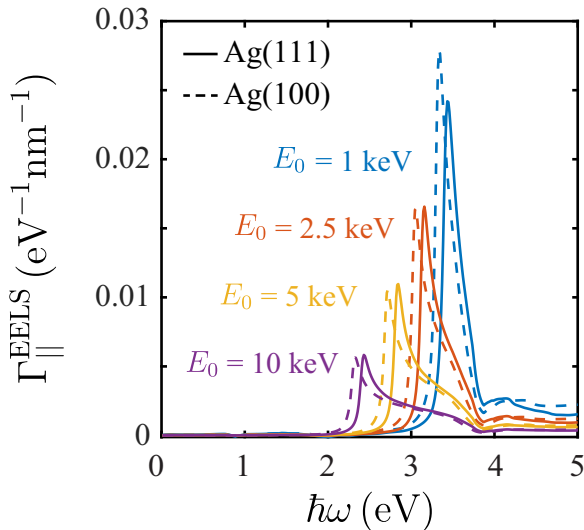


FIG. 4. Plasmon dependence on crystallographic surface orientation: Ag(111) and Ag(100) films. We compare EELS spectra calculated in the ALP model under the same conditions as in Figs. 3(c) and 3(d) for $N = 13$ ML Ag(111) and $N = 15$ ML Ag(100) films (thickness ratio differing by $<0.1\%$).

orientation (see Table I). We explore the effects of this dependence by comparing aloof EELS spectra obtained from Ag(111) and Ag(100) films in Fig. 4. In order to eliminate discrepancies arising from differences in thickness, we consider films consisting of $N = 13$ and $N = 15$ MLs, respectively, so that the thickness ratio is $(2/\sqrt{3}) \times (13/15) \approx 1.001$. We remind that Ag(111) displays a projected bulk gap in the electronic bands, in contrast to Ag(100) (see Fig. S1(a) in the SM [74]); as a consequence the former supports electronic surface states, unlike the latter [58]. Despite these remarkable differences in electronic structure, the resulting spectra look rather similar, except for a small redshift of Ag(100) plasmon peaks relative to Ag(111), comparable in magnitude to those observed in semi-infinite Ag(111) and Ag(110) crystal surfaces through angle-resolved low-energy EELS [75], although the actual magnitude of the shift might be also influenced by electron confinement in our ultrathin films.

The presence of a dielectric substrate of permittivity ϵ_s is known to redshift the plasmon frequency of thin films by a factor of $\sim 1/\sqrt{1 + \epsilon_s}$ due to the attractive image interaction [76]. This effect is observed in our calculated aloof EELS spectra, for which we obtain the combined film-substrate reflection coefficient by using a Fabry-Pérot approach, as discussed elsewhere [33]. We find again excellent agreement between RPA simulations using the ALP and classical calculations (Fig. 5), and in fact, the resemblance between the spectral profiles obtained with both methods increases with ϵ_s .

In Fig. 6 we examine the way lateral dispersion of QW states affects the plasmonic properties of ultrathin Ag films when using the ALP. Comparison of the band structures calculated with [Fig. 6(b)] and without [Fig. 6(a)] inclusion of an energy dependence in the in-plane effective mass anticipates a clear difference between the two of them: the latter shows the same energy jumps between different bands irrespective of the electron parallel wave vector k_{\parallel} ; those energy jumps

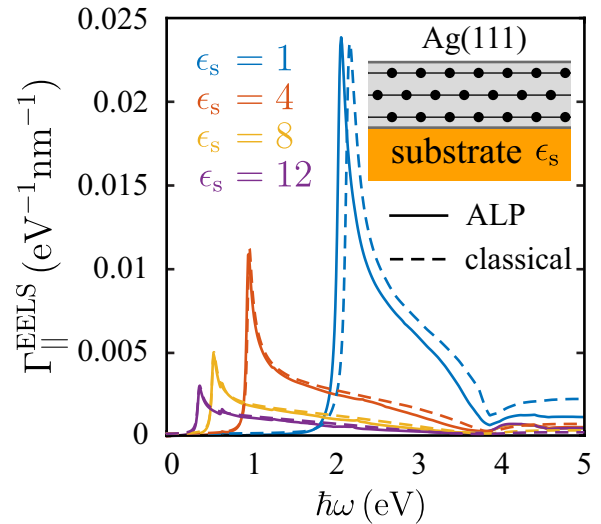


FIG. 5. Substrate-induced plasmon shift. We show EELS spectra for 2.5-keV electrons calculated in either the ALP model or the local classical description under the same conditions as in Fig. 3 for a Ag(111) film consisting of $N = 5$ MLs supported on a planar dielectric substrate of permittivity ϵ_s , as indicated by labels.

will therefore be favored in the optical response, giving rise to spurious spectral features. In contrast, differences in lateral dispersion associated with the energy dependence of the effective mass (described here by fitting existing angle-resolved photoemission data [60,61,63,64,77,78]) should at least partially wash out those spectral features. This is clearly observed in the resulting dispersion diagrams [Figs. 6(c) and 6(d)] and aloof EELS spectra [Figs. 6(e) and 6(f)]. In particular, the dispersion relation for constant m_j^* [Fig. 6(c)] reveals a complex mixture of resonances at energies above 3 eV, which we find to be strongly affected by the HOMO-LUMO gap energy (not shown); these resonances cause fine structure in the EELS spectra that disappears when a realistic energy dependence is introduced in the lateral effective mass [Fig. 6(e)].

We also analyze EELS spectra for normally impinging electron beams (Fig. 7). The momentum- and energy-resolved EELS probability given by Eq. (13) reveals the plasmon dispersion in analogy to the loss function [cf. Figs. 3(b) and 7(b)]. But now, this quantity is directly accessible under normal incidence by recording angle- and energy-dependent electron transmission intensities, as already done in pioneering experiments for thicker Al films showing both bonding and antibonding plasmon dispersions [34]. In contrast to the aloof configuration, the transmission EELS spectra exhibit broader plasmon features [Figs. 7(c) and 7(d)], which in the thin-film limit [70] are the result of weighting the loss function with a profile $Q^2/(Q^2 + \omega^2/v^2)^2$ [see also Eq. (15); the extra factor of Q emerges from χ in the small- Q limit], represented in Fig. 7(b) for 2.5 keV electrons and different energies $\hbar\omega$ (colored curves); these spectra reveal indeed a broad spectral overlap with the plasmon band. Again, we observe very similar results from RPA and classical descriptions, and just a minor dependence on electron potential in the former.

We conclude by showing EELS calculations for Au(111) films in Fig. 8. This noble metal has a similar

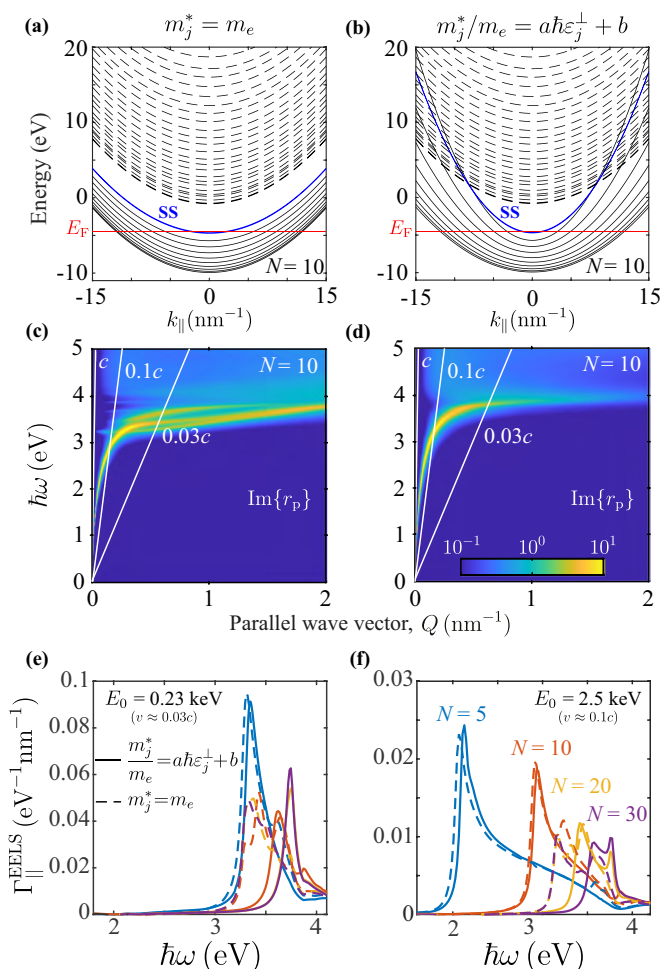


FIG. 6. The role of the electron effective mass. (a, b) In-plane parabolic QW bands of an $N = 10$ ML Ag(111) film in the ALP model with (a) constant and (b) energy-dependent effective mass ($m_j^* = m_e$ and $m_j^* = (a\hbar\varepsilon_j^\perp + b)m_e$, respectively; see Table I). The surface-state band in (b) (blue curve) has a mass $0.4m_e$. Solid (dashed) curves represent bands that are occupied (unoccupied) at $k_{\parallel} = 0$. The Fermi level is shown as a horizontal red line. (c, d) Loss function $\text{Im}\{r_p\}$ under the conditions of (a) and (b), respectively. (e, f) EELS probability under parallel aloof interaction at a distance $z_0 = 0.5$ nm for two different electron energies corresponding to the $\omega = Qv$ lines shown in (c) and (d) and different film thicknesses (see labels) calculated in the ALP model with constant (dashed curves) and energy-dependent (solid curves) electron effective mass.

conduction-electron density as Ag, but the Au d band is closer to the Fermi energy, therefore producing large screening ($\epsilon_b \sim 9$ in the plasmonic region) compared with Ag ($\epsilon_b \sim 4$; see Fig. S4 in the SM [74]). This causes a shift of the high- Q surface plasmon asymptote down to $\hbar\omega_s \approx 2.5$ eV. Additionally, damping is also stronger (more than three times larger than in Ag; see the Appendix), which results in broader spectral features [cf. Fig. 8 for Au and Figs. 3(c) 3(d), 7(c) and 7(d) for Ag]. Interestingly, we observe significant blueshifts in the plasmon spectral features when using the ALP as compared with both jellium DFT and classical models. This effect could originate in a more substantial role played by the electronic band structure in Au(111) because

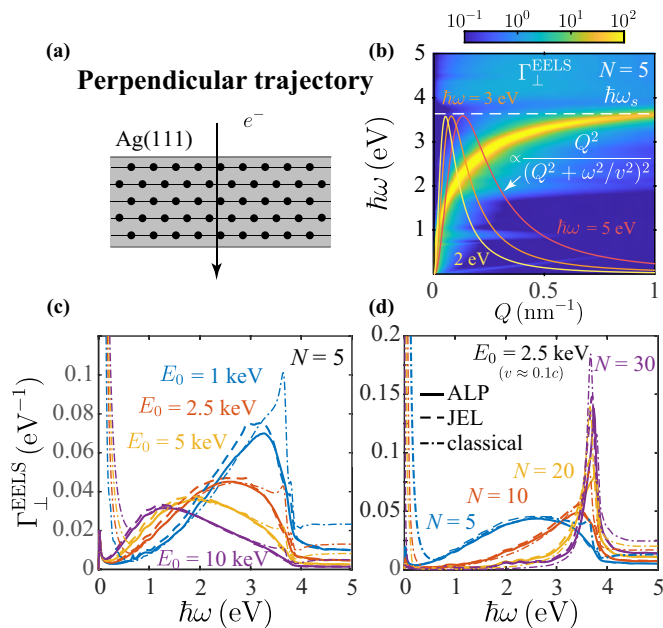


FIG. 7. EELS in thin Ag(111) films under normal incidence. (a) Scheme showing an electron normally traversing a $N = 5$ ML Ag(111) metal film. (b) Momentum- and energy-resolved EELS probability $\Gamma_{\perp}^{\text{EELS}}(Q, \omega)$ [Eq. (13)] calculated for $E_0 = 2.5$ keV electrons ($v/c \approx 0.1$) in the ALP model for the film shown in (a). Colored solid curves show $Q^2/(Q^2 + \omega^2/v^2)^2$ profiles as a function of Q for different energy losses $\hbar\omega = 2, 3$, and 5 eV, while the dashed horizontal line indicates $\hbar\omega_s$. (c, d) EELS probability calculated using different models [see legend in (d)] for (c) different electron kinetic energies E_0 with fixed $N = 5$ and (d) different N 's with $E_0 = 2.5$ keV.

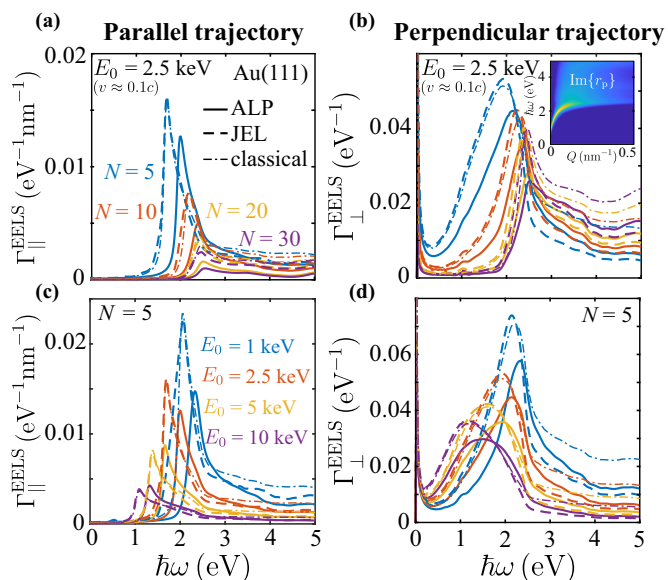


FIG. 8. EELS spectra for gold Au(111) films. We consider (a, c) aloof and (b, d) normal trajectories for either (a, b) fixed electron energy ($E_0 = 2.5$ keV) and varying film thickness ($N = 5$ –30 MLs) or (c, d) fixed $N = 5$ and varying electron energy. Calculations for the same models as in Fig. 3 are presented. The plasmon dispersion is shown for $N = 10$ MLs using the ALP model in the inset of (b).

the projected bulk gap extends further below the Fermi level, and additionally, the surface-state band is also more deeply bound (see Fig. S1(b) in the SM [74]). This is consistent with the general dependence of the optical surface conductivity on Fermi momentum k_F and velocity v_F : in the Drude model for graphene and the two-dimensional electron gas, this quantity is proportional to $k_F v_F$ and the surface plasmon frequency scales as $\propto \sqrt{k_F v_F}$; the situation is more complicated in our thin films because they have multiple 2D bands crossing the Fermi level, but the presence of a deeper gap in Au(111) indicates that the effective band-averaged value of $k_F v_F$ (i.e., with k_F defined by the crossing of each QW at the Fermi level and v_F as the slope of the parabolic dispersion at that energy) is larger than in Ag surfaces, characterized by the presence of shallower bands near E_F . We thus expect an increase in Drude weight, and consequently a plasmon blueshift, in Au(111) relative to Ag; this argument is reinforced by the small effective mass of surface states in Au(111) compared with Ag(111), which also pushes up their associated v_F . In summary, the plasmon blueshifts observed in Au(111) when using the realistic ALP seem to have a physical origin, although more sophisticated first-principles simulations might be needed to conclusively support this finding.

IV. CONCLUSION

In summary, we have shown that a local classical dielectric model predicts reasonably well the intensities and dispersion relations of plasmons in ultrathin silver films when compared to quantum-mechanical simulations based on the RPA with different potentials used to simulate the conduction one-electron wave functions. We attribute the small effects of non-locality and quantum confinement in the plasmonic response of these films to the fact that their associated electron motion takes place along in-plane directions, in contrast to metal nanoparticles with a similar size as the film thickness here considered (i.e., electron surface scattering is unavoidable in such particles, thus introducing important nonlocal effects). We confirm this agreement between classical and quantum simulations in Ag and Au films down to a few atomic layers in thickness [33,59], consistent with previous smooth-interface hydrodynamic theory [79].

Additionally, our quantum RPA simulations are relatively insensitive to the details of the confining electron potential, so similar results are obtained when using either a smooth jellium DFT model or a phenomenological potential that incorporates atomic-layer corrugation to fit relevant elements of the electronic band structure. In particular, the latter produces results that are rather independent of the crystallographic orientation of the film. These results support the intuition that the plasmons under consideration are integrated quantities, which essentially depend on the number of electrons in the system, as well as the effect of background screening produced by polarization of inner shell electrons, and not so much on the details of the electronic wave functions. Nonetheless, it is important to introduce the correct energy dependence of the out-of-plane effective mass in the phenomenological

potential model, as otherwise spurious features show up in the calculated plasmon spectra. Although jellium DFT and phenomenological potentials lead to substantially different plasmon charge distributions, spatial integration gives rise to similar plasmon dispersion relations. Interestingly, band effects described in the ALP model are more significant in Au, where they produce plasmon blueshifts relative to the predictions of classical and jellium DFT simulations; we attribute this different behavior in Au(111) relative to Ag(111) and Ag(100) to the fact that the former surface exhibits a projected bulk gap that extends further below the Fermi level, and additionally this gives rise to more bound surface states. We remark that EELS provides the means to access the dispersion relations of strongly confined plasmons in ultrathin metal films, which are too far from the light line to be measured by means of optical techniques.

ACKNOWLEDGMENTS

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APPENDIX: BACKGROUND SCREENED INTERACTION

We introduce the effect of interband polarization in the plasmonic spectral region of noble metals through a dielectric slab of permittivity $\epsilon_b(\omega) = \epsilon(\omega) + \omega_p^2/\omega(\omega + i\gamma^{\text{exp}})$, that is, the local dielectric function of the bulk metal $\epsilon(\omega)$ from which we subtract a classical bulk Drude term representing the contribution of conduction electrons. In practice, we take $\epsilon(\omega)$ from measured optical data [67] and use parameters $\hbar\omega_p = 9.17$ eV and $\hbar\gamma^{\text{exp}} = 21$ meV for Ag, and $\hbar\omega_p = 9.06$ eV and $\hbar\gamma^{\text{exp}} = 71$ meV for Au. The resulting $\epsilon_b(\omega)$ is plotted in Fig. S4 of the SM [74]. Incidentally, as we explain in Sec. II B, we set the damping parameter to $\gamma = \gamma^{\text{exp}}/2$ in the RPA formalism in order to fit the experimental plasmon width. Following previous work [19], we take the background dielectric slab to have a thickness $d = Na_s$, where N is the number of atomic layers and a_s is the interlayer spacing, so that it extends symmetrically a distance $a_s/2$ outside the outer atomic plane on each side of the film.

We reproduce for convenience a previously reported expression [33] for the screened interaction, used here to account for background polarization in a self-standing metal film of thickness d and background permittivity ϵ_b contained in the $0 < z < d$ region:

$$v(Q, z, z') = v^{\text{dir}}(Q, z, z') + v^{\text{ref}}(Q, z, z'),$$

where

$$v^{\text{dir}}(Q, z, z') = \frac{2\pi}{Q} e^{-Q|z-z'|} \times \begin{cases} 1, & z, z' \leq 0 \text{ or } z, z' > d \\ 1/\epsilon_b, & 0 < z, z' \leq d \\ 0, & \text{otherwise,} \end{cases}$$

and

$$v^{\text{ref}}(Q, z, z') = \frac{(2\pi/Q)}{(\epsilon_b + 1)^2 - (\epsilon_b - 1)^2 e^{-2Qd}} \times \begin{cases} (1 - \epsilon_b^2)(e^{2Qd} - 1)e^{-Q(z+z')}, & d < z, z' \\ 2[(\epsilon_b + 1)e^{-Q(z-z')} + (\epsilon_b - 1)e^{-Q(z+z')}], & 0 < z' \leq d < z \\ 4\epsilon_b e^{-Q(z-z')}, & z' \leq 0 \text{ and } d < z \\ 2[(\epsilon_b + 1)e^{Q(z-z')} + (\epsilon_b - 1)e^{-Q(z+z')}], & 0 < z \leq d < z' \\ (1/\epsilon_b)\{(\epsilon_b^2 - 1)[e^{-Q(z+z')} + e^{-Q(2d-z-z')}] \\ \quad + (\epsilon_b - 1)^2[e^{-Q(2d+z-z')} + e^{-Q(2d-z-z')}] \}, & 0 < z, z' \leq d \\ 2[(\epsilon_b + 1)e^{-Q(z-z')} + (\epsilon_b - 1)e^{-Q(2d-z-z')}], & z' \leq 0 < z \leq d \\ 4\epsilon_b e^{Q(z-z')}, & z \leq 0 \text{ and } d < z' \\ 2[(\epsilon_b + 1)e^{Q(z-z')} + (\epsilon_b - 1)e^{-Q(2d-z-z')}], & z \leq 0 < z' \leq d \\ (1 - \epsilon_b^2)(1 - e^{-2Qd})e^{Q(z+z')}, & z, z' \leq 0. \end{cases}$$

For completeness, we illustrate the dramatic effects of interband processes in Fig. S5 of the SM [74] by comparing calculations obtained for Ag films using either screened or bare Coulomb interactions.

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- [1] D. J. Bergman and M. I. Stockman, *Phys. Rev. Lett.* **90**, 027402 (2003).
- [2] H. Xu, E. J. Bjerneld, M. Käll, and L. Börjesson, *Phys. Rev. Lett.* **83**, 4357 (1999).
- [3] A. Polman, *Science* **322**, 868 (2008).
- [4] J. N. Anker, W. P. Hall, O. Lyandres, N. C. Shah, J. Zhao, and R. P. Van Duyne, *Nat. Mater.* **7**, 442 (2008).
- [5] D. Rodrigo, O. Limaj, D. Janner, D. Etezadi, F. J. García de Abajo, V. Pruneri, and H. Altug, *Science* **349**, 165 (2015).
- [6] Z. W. Seh, S. Liu, M. Low, S.-Y. Zhang, Z. Liu, A. Mlayah, and M.-Y. Han, *Adv. Mater.* **24**, 2310 (2012).
- [7] C. Clavero, *Nat. Photonics* **8**, 95 (2014).
- [8] H. A. Atwater and A. Polman, *Nat. Mater.* **9**, 205 (2010).
- [9] C. F. Guo, T. Sun, F. Cao, Q. Liu, and Z. Ren, *Light Sci. Appl.* **3**, e161 (2014).
- [10] M. Danckwerts and L. Novotny, *Phys. Rev. Lett.* **98**, 026104 (2007).
- [11] R. W. Boyd, *Nonlinear Optics*, 3rd ed. (Academic Press, Amsterdam, 2008).
- [12] A. R. Davoyan, I. V. Shadrivov, and Y. S. Kivshar, *Opt. Express* **16**, 21209 (2008).
- [13] S. Palomba and L. Novotny, *Phys. Rev. Lett.* **101**, 056802 (2008).
- [14] R. H. Ritchie, *Phys. Rev.* **106**, 874 (1957).
- [15] C. J. Powell and J. B. Swan, *Phys. Rev.* **115**, 869 (1959).
- [16] K. D. Tsuei, E. W. Plummer, and P. J. Feibelman, *Phys. Rev. Lett.* **63**, 2256 (1989).
- [17] M. Rocca, L. Yibing, F. Buatier de Mongeot, and U. Valbusa, *Phys. Rev. B* **52**, 14947 (1995).
- [18] J. Daniels, *Z. Phys.* **203**, 235 (1967).
- [19] A. Liebsch, *Phys. Rev. B* **48**, 11317 (1993).
- [20] N. D. Lang and W. Kohn, *Phys. Rev. B* **1**, 4555 (1970).
- [21] N. D. Lang and W. Kohn, *Phys. Rev. B* **7**, 3541 (1973).
- [22] J. M. Pitarke, V. M. Silkin, E. V. Chulkov, and P. M. Echenique, *Rep. Prog. Phys.* **70**, 1 (2007).
- [23] J. F. Dobson and G. H. Harris, *J. Phys. C* **21**, L729 (1988).
- [24] A. Eguiluz, S. C. Ying, and J. J. Quinn, *Phys. Rev. B* **11**, 2118 (1975).
- [25] K. D. Tsuei, E. W. Plummer, A. Liebsch, K. Kempa, and P. Bakshi, *Phys. Rev. Lett.* **64**, 44 (1990).
- [26] A. Liebsch and W. L. Schaich, *Phys. Rev. B* **52**, 14219 (1995).
- [27] H. Raether, *Surface Plasmons on Smooth and Rough Surfaces and on Gratings*, Springer Tracks in Modern Physics Vol. 111 (Springer-Verlag, Berlin, 1988).
- [28] S. A. Maier, *Plasmonics: Fundamentals and Applications* (Springer, New York, 2007).
- [29] W. L. Barnes, A. Dereux, and T. W. Ebbesen, *Nature (London)* **424**, 824 (2003).
- [30] R. A. Álvarez-Puebla, L. M. Liz-Marzán, and F. J. García de Abajo, *J. Phys. Chem. Lett.* **1**, 2428 (2010).
- [31] E. N. Economou, *Phys. Rev.* **182**, 539 (1969).
- [32] Z. M. Abd El-Fattah, V. Mkhitarian, J. Brede, L. Fernández, C. Li, Q. Guo, A. Ghosh, A. Rodríguez Echarrri, D. Naveh, F. Xia *et al.*, *ACS Nano* **13**, 7771 (2019).
- [33] A. Rodríguez Echarrri, J. D. Cox, and F. J. García de Abajo, *Optica* **6**, 630 (2019).
- [34] R. Vincent and J. Silcox, *Phys. Rev. Lett.* **31**, 1487 (1973).
- [35] F. Moresco, M. Rocca, T. Hildebrandt, and M. Henzler, *Phys. Rev. Lett.* **83**, 2238 (1999).
- [36] E. P. Rugeramigabo, T. Nagao, and H. Pfnür, *Phys. Rev. B* **78**, 155402 (2008).
- [37] H. V. Chung, C. J. Kubber, G. Han, S. Rigamonti, D. Sánchez-Portal, D. Enders, A. Pucci, and T. Nagao, *Appl. Phys. Lett.* **96**, 243101 (2010).
- [38] E. P. Rugeramigabo, C. Tegenkamp, H. Pfnür, T. Inaoka, and T. Nagao, *Phys. Rev. B* **81**, 165407 (2010).
- [39] T. Nagao, S. Yaginuma, T. Inaoka, and T. Sakurai, *Phys. Rev. Lett.* **97**, 116802 (2006).
- [40] G. X. Ni, A. S. McLeod, Z. Sun, L. Wang, L. Xiong, K. W. Post, S. S. Sunku, B.-Y. Jiang, J. Hone, C. R. Dean *et al.*, *Nature (London)* **557**, 530 (2018).
- [41] Z. Fei, A. S. Rodin, G. O. Andreev, W. Bao, A. S. McLeod, M. Wagner, L. M. Zhang, Z. Zhao, M. Thiemens, G. Dominguez *et al.*, *Nature (London)* **487**, 82 (2012).
- [42] J. Chen, M. Badioli, P. Alonso-González, S. Thongrattanasiri, F. Huth, J. Osmond, M. Spasenić, A. Centeno, A. Pesquera, P. Godignon *et al.*, *Nature (London)* **487**, 77 (2012).

- [43] M. B. Lundberg, Y. Gao, R. Asgari, C. Tan, B. V. Duppen, M. Autore, P. Alonso-González, A. Woessner, K. Watanabe, T. Taniguchi *et al.*, *Science* **357**, 187 (2017).
- [44] D. Alcaraz Iranzo, S. Nanot, E. J. C. Dias, I. Epstein, C. Peng, D. K. Efetov, M. B. Lundberg, R. Parret, J. Osmond, J.-Y. Hong *et al.*, *Science* **360**, 291 (2018).
- [45] L. Ju, B. Geng, J. Horng, C. Girit, M. Martin, Z. Hao, H. A. Bechtel, X. Liang, A. Zettl, Y. R. Shen *et al.*, *Nat. Nanotechnol.* **6**, 630 (2011).
- [46] A. N. Grigorenko, M. Polini, and K. S. Novoselov, *Nat. Photonics* **6**, 749 (2012).
- [47] D. N. Basov, M. M. Fogler, and F. J. García de Abajo, *Science* **354**, aag1992 (2016).
- [48] Y. Liu, R. F. Willis, K. V. Emtsev, and T. Seyller, *Phys. Rev. B* **78**, 201403(R) (2008).
- [49] Y. Liu and R. F. Willis, *Phys. Rev. B* **81**, 081406(R) (2010).
- [50] U. Kreibig and M. Vollmer, *Optical Properties of Metal Clusters* (Springer-Verlag, Berlin, 1995).
- [51] J. A. Scholl, A. L. Koh, and J. A. Dionne, *Nature (London)* **483**, 421 (2012).
- [52] F. J. García de Abajo, *Rev. Mod. Phys.* **82**, 209 (2010).
- [53] I. S. Gradshteyn and I. M. Ryzhik, *Table of Integrals, Series, and Products* (Academic Press, London, 2007).
- [54] L. Hedin and S. Lundqvist, in *Solid State Physics*, edited by D. T. F. Seitz and H. Ehrenreich (Academic Press, New York, 1970), Vol. 23, pp. 1–181.
- [55] O. Keller, *Phys. Rev. B* **33**, 990 (1986).
- [56] L. Marušić and M. Šunjić, *Phys. Scr.* **63**, 336 (2001).
- [57] N. D. Mermin, *Phys. Rev. B* **1**, 2362 (1970).
- [58] E. Chulkov, V. Silkin, and P. Echenique, *Surf. Sci.* **437**, 330 (1999).
- [59] E. J. H. Skjølstrup, T. Søndergaard, and T. G. Pedersen, *Phys. Rev. B* **99**, 155427 (2019).
- [60] F. Reinert, G. Nicolay, S. Schmidt, D. Ehm, and S. Hüfner, *Phys. Rev. B* **63**, 115415 (2001).
- [61] V. M. Silkin, J. M. Pitarke, E. V. Chulkov, and P. M. Echenique, *Phys. Rev. B* **72**, 115435 (2005).
- [62] R. Paniago, R. Matzdorf, G. Meister, and A. Goldmann, *Surf. Sci.* **336**, 113 (1995).
- [63] A. Garcia-Lekue, J. M. Pitarke, E. V. Chulkov, A. Liebsch, and P. M. Echenique, *Phys. Rev. B* **68**, 045103 (2003).
- [64] S. D. Kevan and R. H. Gaylord, *Phys. Rev. B* **36**, 5809 (1987).
- [65] D. Popović, F. Reinert, S. Hüfner, V. G. Grigoryan, M. Springborg, H. Cercellier, Y. Fagot-Revurat, B. Kierren, and D. Malterre, *Phys. Rev. B* **72**, 045419 (2005).
- [66] M. Kerker, *The Scattering of Light and Other Electromagnetic Radiation: Physical Chemistry: A Series of Monographs* (Academic Press, New York, 2013), Vol. 16.
- [67] P. B. Johnson and R. W. Christy, *Phys. Rev. B* **6**, 4370 (1972).
- [68] E. Kröger, *Z. Phys.* **216**, 115 (1968).
- [69] F. J. García de Abajo, A. Rivacoba, N. Zabala, and N. Yamamoto, *Phys. Rev. B* **69**, 155420 (2004).
- [70] F. J. García de Abajo, *ACS Nano* **7**, 11409 (2013).
- [71] N. W. Ashcroft and N. D. Mermin, *Solid State Physics* (Harcourt College, Philadelphia, 1976).
- [72] T.-C. Chiang, *Surf. Sci. Rep.* **39**, 181 (2000).
- [73] A. Goldmann, V. Dose, and G. Borstel, *Phys. Rev. B* **32**, 1971 (1985).
- [74] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevResearch.2.023096> for results similar to Fig. 1(d) for Ag(100) and Au(111) films; a comparison of the reflection coefficients obtained in the RPA with different confining potentials; an illustration of the convergence of the dispersion relation toward the semi-infinite surface limit with increasing number of atomic layers; the background dielectric function used to account for interband transitions in Ag and Au; and a study of the effect of background screening.
- [75] S. Suto, K.-D. Tsuei, E. W. Plummer, and E. Burstein, *Phys. Rev. Lett.* **63**, 2590 (1989).
- [76] F. J. García de Abajo, *ACS Photonics* **1**, 135 (2014).
- [77] M. A. Mueller, T. Miller, and T.-C. Chiang, *Phys. Rev. B* **41**, 5214 (1990).
- [78] I. Matsuda, T. Tanikawa, S. Hasegawa, H. W. Yeom, K. Tono, and T. Ohta, *e-J. Surf. Sci. Nanotechnol.* **2**, 169 (2004).
- [79] C. David and F. J. García de Abajo, *ACS Nano* **8**, 9558 (2014).