

[A semi-analytical decomposition analysis of surface plasmon generation and the optimal nanoledge plasmonic device](#)

By: Zheng Zeng, Madu N. Mendis, David H. Waldeck, and [Jianjun Wei](#)

Z. Zeng, M.N. Mendis, D.H. Waldeck, J. Wei, A semi-analytical decomposition analysis of surface plasmon generation and the optimal nanoledge plasmonic device, *RSC Advances*, **2016**, 6(21), 17196 – 17203. DOI: 10.1039/C6RA01105E

Made available courtesy of the Royal Society of Chemistry:

<https://doi.org/10.1039/C6RA01105E>

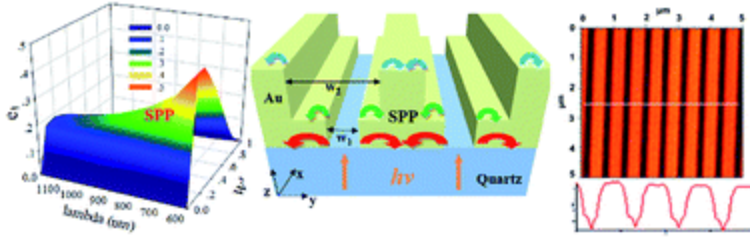
*****© 2016 The Royal Society of Chemistry. Reprinted with permission. No further reproduction is authorized without written permission from The Royal Society of Chemistry. This version of the document is not the version of record. Figures and/or pictures may be missing from this format of the document. *****

Abstract:

Surface plasmon resonance (SPR) of nanostructured thin metal films (so-called nanoplasmonics) has attracted intense attention due to its versatility for optical sensing and chip-based device integration. Understanding the underlying physics and developing applications of nanoplasmonic devices with desirable optical properties, *e.g.* intensity of light scattering and high refractive index (RI) sensitivity at the perforated metal film, is crucial for practical uses in physics, biomedical detection, and environmental monitoring. This work presents a semi-analytical model that enables decomposition and quantitative analysis of surface plasmon generation at a new complex nanoledge aperture structure under plane-wave illumination, thus providing insight on how to optimize plasmonic devices for optimal plasmonic generation efficiencies and RI sensitivity. A factor analysis of parameters (geometric, dielectric-RI, and incident wavelength) relevant to surface plasmon generation is quantitatively investigated to predict the surface plasmon polariton (SPP) generation efficiency. In concert with the analytical treatment, a finite-difference time-domain (FDTD) simulation is used to model the optical transmission spectra and RI sensitivity as a function of the nanoledge device's geometric parameters, and it shows good agreement with the analytical model. Further validation of the analytical approach is provided by fabricating subwavelength nanoledge devices and testing their optical transmission and RI sensitivity.

Keywords: nanoplasmonics | surface plasmon polariton (SPP) | refractive index (RI) | nanoledge devices

Article:



Introduction

Surface plasmon resonance (SPR), an optical phenomenon that is very sensitive to the near surface dielectric constant (refractive index, RI),¹ is well-suited to the detection of surface binding events of chemical and biological agents,^{2,3} with single molecule sensitivity⁴⁻⁶ and compatibility with point-of-care (POC) platforms.⁷⁻⁹ Similarly, metal films that are perforated by subwavelength holes (or slits) display extraordinary optical transmission (EOT) in the nanostructure apertures,¹⁰⁻¹⁵ which arises from strong surface plasmon excitation, and display high refractive index unit (RIU) sensitivity. Consequently, understanding the underlying physics and developing applications of nanoplasmonics with desirable optical properties,¹⁶ *e.g.* intensity of light scattering and high RIU sensitivity at the perforated metal film,¹⁷ are of particular interest for realizing their promise and integrating them into on-chip photonic sensing platforms.¹⁸

Real metals with a finite conductivity are capable of sustaining surface plasmon polariton (SPP) modes, which are bounded at the interface, and mediate the interaction between the nanoapertures at visible or near-infrared frequencies.¹⁹⁻²¹ The SPP generation at the input and output aperture sides of an isolated subwavelength slit, when illuminated by an incident plane-wave or a slit-mode, has been described in a quantitative manner.²²⁻²⁴ The essential results can be generalized and applied to more complicated nano-aperture array structures, allowing for a quantitative analysis of SPP generation and its dependence on different device parameters. This analytical approach can be tested by numerical techniques: finite-element methods (FEM), finite-difference time-domain (FDTD), discrete dipole approximation (DDA), multiple multipole (MMP), and more recent a combination of surface integral equation (SIE) method of moments (MoM) formulation. They have all been applied for modeling the electromagnetic dynamics of nanoplasmonic systems.²⁵⁻²⁷ Among them, the well-established FDTD technique solves Maxwell's equations and provides both qualitative insight and a quantitative link between the optical properties and the underlying SPP properties of the nanoaperture arrays.²⁸

The present work considers a semi-analytical analysis and numerical simulations to investigate a complex nanoaperture–nanoedge device (Fig. 1), which displays SPP phenomena and the extraordinary optical transmission (EOT) of light, with the aim of elucidating the criteria for optimal optical performance and improving its refractive index sensitivity for sensing applications. First, we present an approximate model to examine the generation of surface plasmons on the nanoedge aperture and then combine it with plane wave and slit-mode illumination to quantify the interaction. Through a corresponding factor analysis we identify how the geometric features of the nanoedge structure affect the plasmon generation. This semi-analytical model is applied to predict the SPP generation in nanoedge structures and investigate the origin of their high plasmonic generation efficiencies. In concert, the FDTD method is used

to predict the optical transmission spectra and RI sensitivity as a function of the nanoledge structure's geometric parameters. Lastly, subwavelength nanoledge devices are fabricated and their optical response is measured in order to validate the results obtained from the semi-analytical analysis and FDTD modelling.

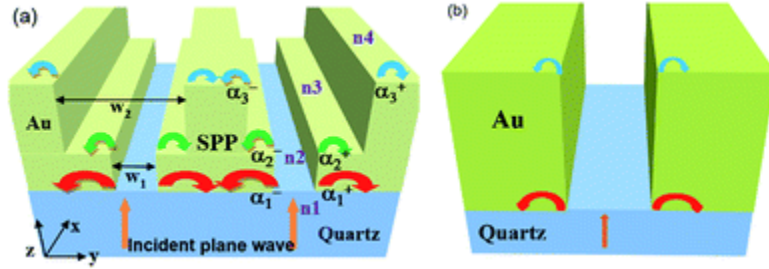


Fig. 1 (a) The schematic illustrates the parameters for the nanoledge structure and SPP generation by a plane wave at normal incidence. The w_1 and w_2 represent the slit widths at the Au–quartz and Au–air interfaces, and the α_1^+ , α_1^- , α_2^+ , α_2^- , α_3^+ , α_3^- represent the SPP generation coefficients at the three interfaces (red, green, and blue arrows, respectively) with inverse propagation directions. The refractive indexes inside the slits are represented by n_2 and n_3 , and those at the outer slits are presented by n_1 and n_4 , when exposed in air, $n_2 = n_3 = n_4 = 1$ (air), and $n_1 = 1.45$ (quartz). (b) The schematic of a straight single nanoslit structure is shown as a comparison.

Results and discussion

Analytical considerations

In order to study nanoledge geometries that are of interest in practice and consider the geometric diffraction with the bounded SPP modes launching on the flat interfaces surrounding the slits, a mechanistic description for SPP generation is needed, especially the SPP scattering coefficients and efficiencies at the slit apertures. Fig. 1 illustrates schematic of a nanoledge structure in subwavelength thick gold film at quartz substrate (Fig. 1a) and a straight nanoslit structure (Fig. 1b) as a comparison. In this study, we focus on the SPP generation at the Au/medium interfaces upon light excitation without considering the height conditions (*i.e.* subwavelength thickness of the metallic film). Note that the thickness (height) predominately affects the SPP fundamental modes in the slit traveling upward and downward, not the SPP generation confined at the flat interfaces;²⁹ hence it is not considered in detail here.

With the semi-analytical model (see Method section), the SPP excitation efficiency e for one side of the aperture is readily calculated with analytical techniques.³⁰ Fig. 2 shows the decomposed SPP excitation efficiency e on one side of the aperture for the interfaces of the nanoledge as a function of the scaled slit width w' and the incident light wavelength, from the visible to near-infrared (600–1200 nm). The SPP excitation is efficient at visible frequencies while e rapidly decreases with the increase of wavelength. For the interfaces surrounding the ledge structure, all of the optimal scaled slit widths are similar with a value of $w' = 0.2$.

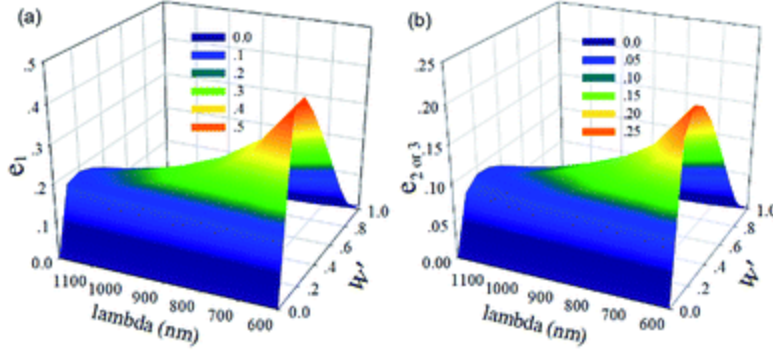


Fig. 2 The SPP generation efficiencies e at the Au–quartz and Au–air interfaces are plotted as a function of λ and w' obtained by the semi-analytical model. (a) Au–quartz interface $e_1 = |\alpha_1^+(w_1/2)|^2 = |\alpha_1^-(w_1/2)|^2$; (b) Au–air interface, $e_2 = |\alpha_2^+(w_1/2)|^2 = |\alpha_2^-(w_1/2)|^2$, the e_3 has the same performance as e_2 (see more in Fig. S1†).

If one selects $n_1 = 1.45$ for quartz and $n_2 = n_3 = n_4 = 1$ for air in the model,³¹ then the optimal nanoedge widths (Fig. 1a) are $w_1 = 0.14\lambda$ for the bottom Au–quartz slit and $w_2 = 0.2\lambda$ for the top Au–air slit. Moreover, at a visible wavelength of 600 nm, the SPP excitation efficiencies are fairly large. The maximum e is calculated as 0.496 for the Au–quartz interface and 0.224 for the Au–air interface of the nanoedge structure in Fig. 1. It is expected that the total SPP excitation efficiency e will result from a “superposition” of the SPP arising from all the interfaces of the nanoedge structures.

A factor analysis³² of the semi-analytical equations was performed in order to assess the correlations of w , λ , n_1 , $n_2/n_3/n_4$ ($n_2 = n_3 = n_4$ in this analysis), and u on the calculation variables of w' , I_0 , I_1 , e , v , and ε in the SPP analysis (see details in ESI, Fig. S2 and ESI†). Fig. 3 shows some radar plots of the influence factor coefficient for different cases. For the Au/quartz interface (Fig. 3a), the factor coefficient of the slit width w on the SPP generation efficiency, e , is 0.12, which is almost the same with that in the Au–air interfaces as shown in Fig. 3b. Moreover, the factor coefficient of the wavelength λ on e becomes larger from the Au–quartz interface to the Au–medium interfaces, in agreement with eqn (6), (11), and (15) (see Method section). Meanwhile, the factor coefficient of the refractive indices, $n_2/n_3/n_4$ on e becomes larger while that of substrate n_1 on e becomes smaller; evident from eqn (4), (9) and (13). Beside this, the factor coefficient of wavelength λ on dielectric constant ε or numerical factor u on v is 1 because of their one–one correspondence. Using Fig. 3a and b, we derived the total factor coefficient of each independent variable (w , λ , n_1 , $n_2/n_3/n_4$, or u) on individual dependent variables (w' , I_0 , I_1 , e , v , or ε) and this is shown in Fig. 3c. In Fig. 3c one can see that the numerical factor u plays the most important part in calculating e , as a factor coefficient of 0.29 according to the yellow area shown. Note that the area is formed by the connection between the six spots (like the six factor coefficients of u on w' , I_0 , I_1 , e , v and ε). In order to more clearly understand the role of the slit width, w , in the analytical part (without u), the influence factor coefficient of w on e was obtained and found to be 0.17; see Fig. 3d. From this analysis and the physical considerations described above, the nanoedge widths, w_1 and w_2 , play an important role in SPP generation.

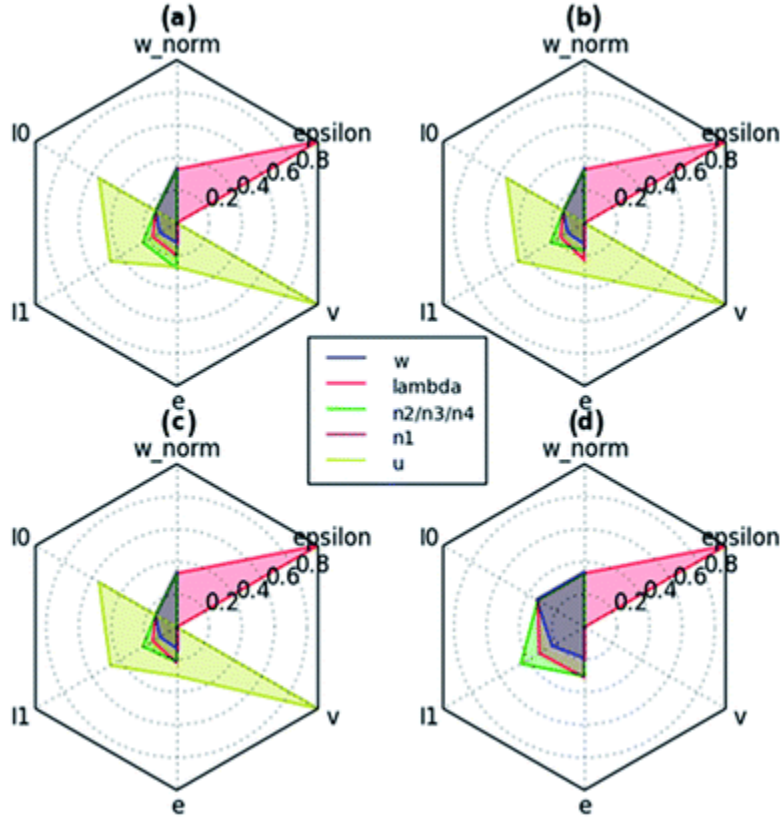


Fig. 3 Radar charts of the factor coefficients of factors (w , λ , $n2/n3/n4$, $n1$, u) on the calculation variables (w' , I_0 , I_1 , e , v , ϵ) in the semi-analytical model are shown for four scenarios. (a) the algebraic operation for e_1 of SPP (red in Fig. 1) at the Au/quartz interface; (b) the algebraic operation for e_2 or e_3 of SPP (green and blue in Fig. 1a) at the Au/Air interfaces; (c) the factor analysis combining (a) with (b); and (d) the analytical part (without u and v) of scenario (c).

Numerical simulation

In order to study the optical transmission properties of the nanoledge structure with different w_1 and w_2 , FDTD calculations were used to simulate the interaction between the metal and the incident light wave.

Fig. 4 summarizes some results of these simulations. Panel (a) in Fig. 4 shows the calculated typical transmission spectra for one selected nanoledge of $w_2 - w_1$ with 280–50 nm (geometries with $w_2 - w_1$ of 280–40 nm, 300–40 nm, and 300–50 nm also performed, see Fig. S8[†]). The four nanoledges were predicted to have high optical transmission (see Fig. 4b and Table S5[†]) because of the transmission resonance corresponding to the Au/quartz mode. The four nanoledge devices have a predicted maximum transmittance of about 27%. The insert in Fig. 4a shows the corresponding TE distribution (more details for the $w_2 - w_1$ of 280–50 nm nanoledge device are shown in Fig. S3a[†]). The SPP generation occurs at three different interfaces which are marked by the white arrows. This was further confirmed by the corresponding TM distributions, given in Fig. S3b.[†] Along with the resulting TM profiles at the reflected surface and at the transmitted surface (see Fig. S4[†]), it suggests that the transmission resonances may arise from localized surface plasmons along the x -direction and SPP propagating along the y -direction.³³

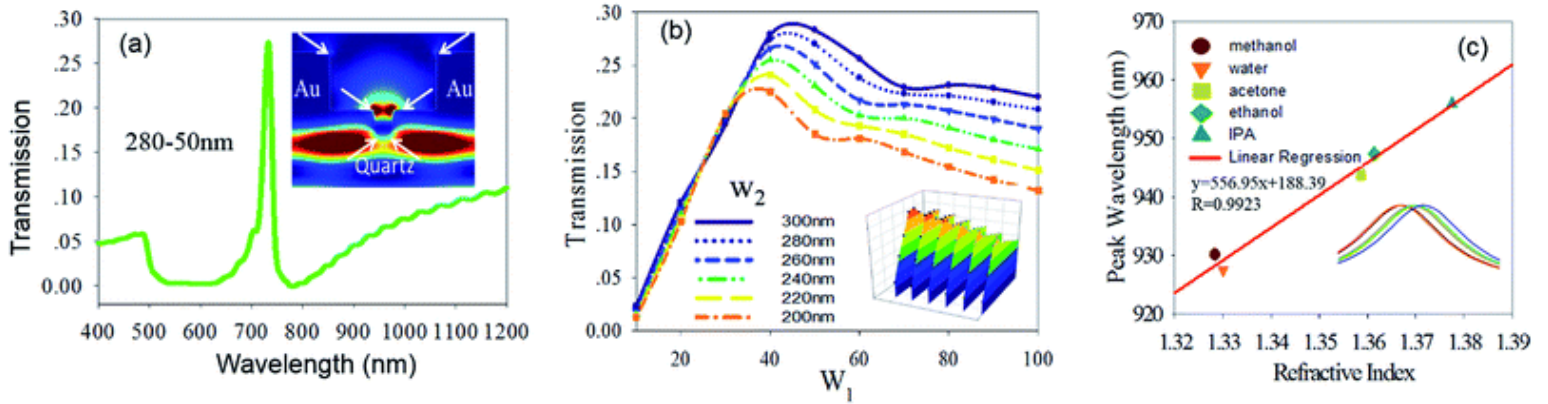


Fig. 4 Panel (a) shows the calculated typical transmission spectra of the $w_2 - w_1$ of 280–50 nm nanoedge system with the inserted corresponding TE field dynamics at 3 seconds calculated for the systems. Panel (b) shows the maximum transmission regarding different w_1 and w_2 in the nanoedge systems with a 3D inset view. Panel (c) shows the peak wavelength of 280–50 nanoslit system vs. refractive indices of bulk solutions, $n_{\text{methanol}} \approx 1.32$, $n_{\text{water}} \approx 1.33$, $n_{\text{acetone}} \approx 1.35$, $n_{\text{ethanol}} \approx 1.36$, and $n_{\text{IPA}} \approx 1.37$ with a peak wavelength shift inset view.

The electromagnetic field distributions reveal that the plasmonic excitations arise from the Au/quartz interface and the gold/medium interfaces with the strength of Au/quartz > Au/air, which is consistent with the results of the semi-analytical model for SPP efficiencies of $e_1 > e_2 > e_3$ for the four nanoedge devices. According to the analytical considerations, the optimal slit widths are $w_1 = 0.14\lambda$ and $w_2 = 0.2\lambda$ for maximum SPP generation. From the FDTD simulations results for w_1 of 40/50 nm and w_2 of 280/300 nm over a broad spectrum ranging from the visible to near-infrared, the $w_2 - w_1$ of 280 nm–50 nm nanoedge system is optimal; *i.e.*, close to optimal slit widths and with the highest optical transmission (Fig. 4b). If we assume an incident light with wavelength of 600 nm, we obtain the scaled widths $w'_1 = 0.121$, $w'_2 = 0.083$ and $w'_3 = 0.467$, corresponding to the three Au/medium interfaces from bottom to top, and the SPP generation efficiencies of $e_1 = 0.48$, $e_2 = 0.18$, and $e_3 = 0.16$, which is consistent with the order of the TE field intensities of the three Au–medium interfaces; see the white arrows shown in Fig. 4a insert. These results indicate that the semi-analytical approach provides insight into the SPP generation efficiencies by enabling a decomposition analysis of the SP in such a complex perforated metal film nanostructure.

Fig. 4c shows an analysis in which the FDTD simulations are used to calculate the refractive index sensitivity of the optimal $w_2 - w_1$ of 280–50 nm nanoedge device. In these simulations the peak wavelength shift was monitored for different refractive indices of the external medium and chosen to mimic the index of refraction of common solvents; namely, methanol, deionized water, acetone, ethanol, and isopropyl alcohol (IPA).³⁴ The calculated wavelength red shifted as the RI of the solvent increased, and the sensitivity was found to be 556 nm per RIU. Furthermore, the plot shows that the dependence of the peak wavelength on the bulk RI of the medium is linear, which makes the nanoedge system highly suitable for biosensing applications.

A comparison of semi-analytical approach and numerical simulation

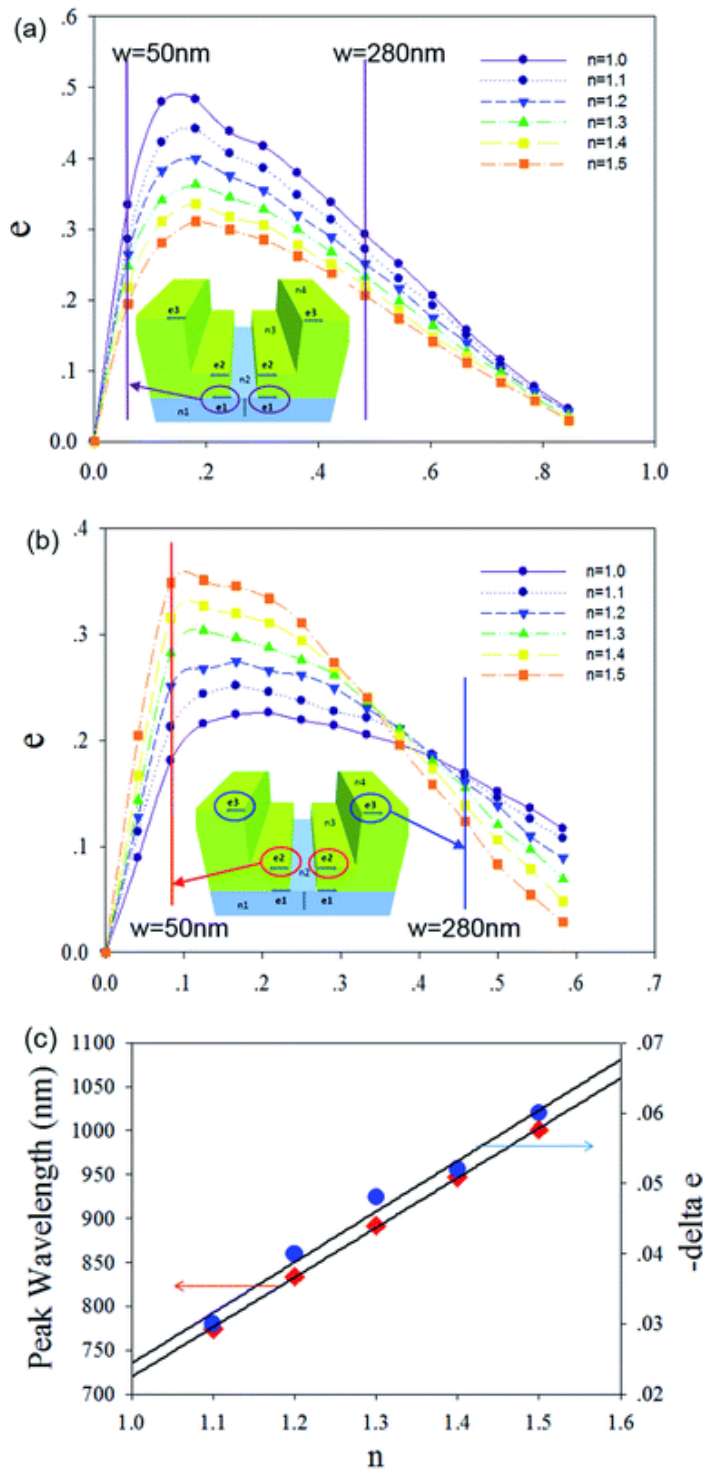


Fig. 5 Decomposition of the SPP generation efficiency e is shown as a function of the nanoledge geometries at different refractive index of $n_2 = n_3 = n_4 (=n)$. Panel (a) shows the SPP generation efficiency e at the Au/quartz interfaces; the two vertical lines indicate the efficiency e at 50 nm and 280 nm slit width. Panel (b) shows the SPP generation efficiency e at the Au/RI-media interfaces, the red and blue lines illustrating the efficiency e of the slit width at 50 nm and 280 nm. Panel (c) compares the EOT peak shift and the total SPP generation efficiency changes as a function of bulk media RIs (see Table S6† and Fig. S5† for individual e values).

As a comparison between the FDTD and the semi-analytical decomposition analysis of SPP generation, Fig. 5 presents the main results of the predicted SPP-generation efficiencies e as a function of the nanoledge widths (w_1, w_2) and RIs ($n_2 = n_3 = n_4 = n$) at the incident wavelength $\lambda = 600$ nm. For the bottom Au/quartz interface, the SPP generation efficiencies e (obtained from eqn (4) with $w'_1 = n_1 w_1 / \lambda$) decrease as the surrounding medium's RI increases (up to 1.5) for the slit width w_1 smaller than λ (Fig. 5a). In contrast, the SPP-generation efficiency e (e_2/e_3) of the Au/medium interfaces increase as the medium's RI increases for slit width below 0.33λ (~ 200 nm at 600 nm incident light), and thereafter transitions to the same dependence as for the Au/quartz interfaces (Fig. 5b). With a specific nanoledge geometry of $w_1 = 50$ nm and $w_2 = 280$ nm, it is interesting to note that the total SPP generation efficiency change, $-\Delta(e_1 + e_2 + e_3)$, has the same dependence on the bulk media RI as the EOT peak shift (Fig. 5c). Given that a weakened SPP generation efficiency correlates with a red shift of the optical transmission peak from a coupling of RI and geometry parameters (w'), the SP decomposition analysis suggests that, for the specific nanoledge $w_2 - w_1$ of 280–50 nm, the SPP generation efficiency (e_2) of Au/medium interface in the center of the nanoledge would result in a blue shift as the RI increases, while a decrease of SPP generation efficiencies (e_1, e_3) at the bottom and top Au/medium interfaces would result in a red shift for the optical transmission. Indeed, the blue shift of optical transmission in a nanoslit cavity has been reported in a previous study.³⁵

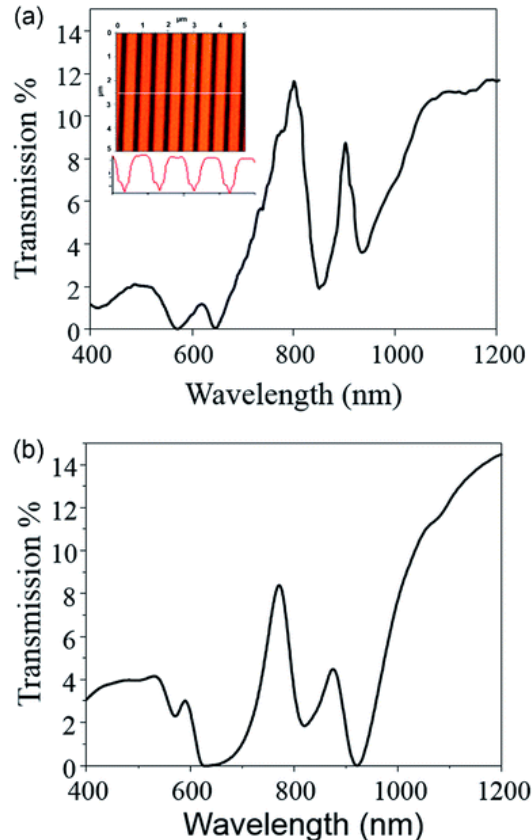


Fig. 6 Experimentally obtained transmission spectrum of a 600 nm periodicity nanoledge structure in air is illustrated in panel (a). The insert in (a) is the AFM image of the nanoledge structure with a cross along the white line is depicted; the scale bar is 1 micron. In panel (b) a FDTD calculated transmission spectrum is shown, in which the slit dimensions/geometry obtained from AFM.

Experimental studies

To further validate our analysis, electron beam lithography (EBL) and focused ion beam (FIB) milling were used to fabricate Au nanoledge structures in arrays ($30 \times 30 \mu\text{m}^2$), and the optical transmission spectra were measured as function of the change of refractive index in the nanoledge area. Fig. 6a inserted and Fig. 7a present AFM and SEM images of geometrically different nanoledge structures that were fabricated.

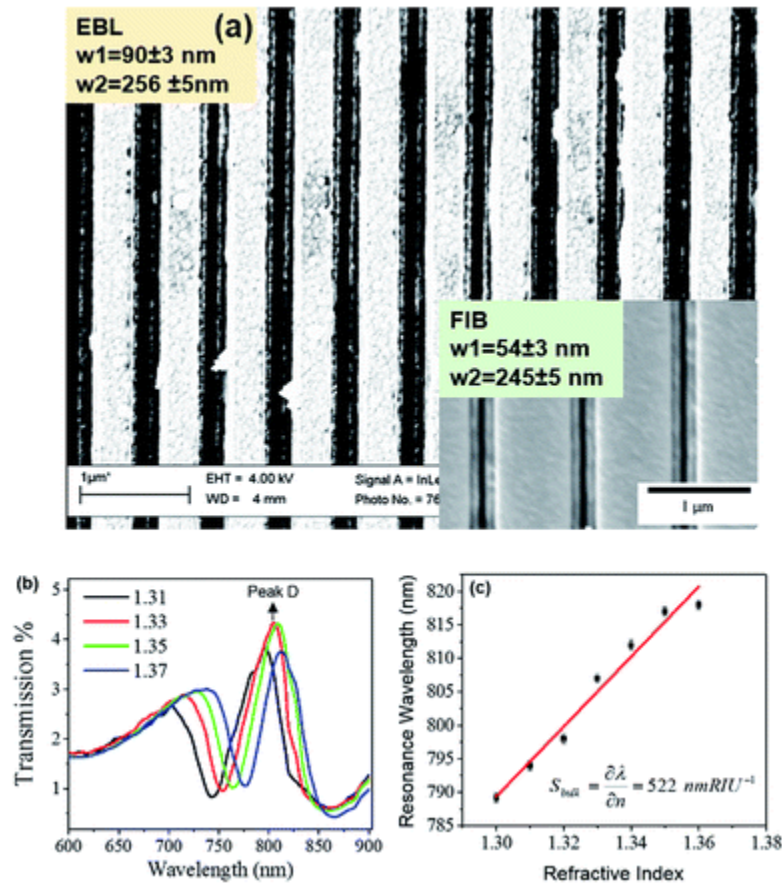


Fig. 7 (a) SEM images of two nanoledge fabricated by EBL and FIB, respectively; (b) the transmission spectra of the FIB fabricated nanoledge arrays ($w_2 - w_1 = 245 - 54 \text{ nm}$) with changes of bulk refractive index; and (c) the primary peak position (peak D) as a function of refractive index obtained from (b).

The experimental transmission spectrum for a FIB fabricated nanoledge array collected in air with a periodicity of 600 nm is shown in Fig. 6a. It can be noted that FDTD calculations predict a sharper main transmission feature (Fig. 4a) in the transmission spectrum as compared to what is experimentally achieved. Broadening of the peaks in the experimental transmission spectrum may arise from fabrication imperfections. It should also be noted that the spectrophotometer, which has a wavelength resolution of 0.782 nm, will not be able to capture these sharp features into the experimental transmission spectrum. To account for fabrication defects, FDTD simulations were performed with realistic slit dimensions taken from AFM imaging of the focused ion beam fabricated nanoledge (Fig. 6a insert). The calculated transmission spectra of

the 600 nm periodicity nanoledge structure is illustrated in Fig. 6b. With these dimensions, the main transmission peak appear less sharp and have much reduced intensity compared the idealized structure. This observation could account for differences in coupling of the cavity modes with SPP modes in the two geometries.

Fig. 7b–c shows the representative transmission spectra of the FIB fabricated nanoledge device (gold thin film on quartz) and the primary transmission peak (D) as a function of the medium RI in the nanoledge. As for the geometry $w_2 - w_1$ of 245–54 nm, the RI sensitivity (S_{bulk}) is 522 nm per RIU, while for the geometry 256 nm/90 nm, the RI sensitivity is 311 nm per RIU (see ESI Fig. S9[†]), which demonstrates a good quantitative agreement with the FDTD simulation. The RI sensitivity is somewhat less than the optimal SPP generation efficiency geometry with $w_2 - w_1$ of 280–50 nm obtained from the perfectly-conducting metal approximation.

Methods

Semi-analytical approach

Based on the mode orthogonality condition,^{23,36} the SPP generation efficiency is governed by the following equations:

$$\int_{-\infty}^{\infty} dz H_y \left(\frac{w}{2}, z \right) E_{\text{SP}}(z) = 2 \left(\alpha^+ \left(\frac{w}{2} \right) + \alpha^- \left(\frac{w}{2} \right) \right) \quad (1)$$

and

$$\int_{-\infty}^{\infty} dz E_z \left(\frac{w}{2}, z \right) H_{\text{SP}}(z) = 2 \left(\alpha^+ \left(\frac{w}{2} \right) - \alpha^- \left(\frac{w}{2} \right) \right) \quad (2)$$

where H and E represent the magnetic and electric field amplitudes, respectively, w is the width of the slit, and $\alpha^+(w/2)$ and $\alpha^-(w/2)$ represent SPP excitation coefficients at the exit sides of the slit.²³ The strength of the SPP generation is provided by $|\alpha|^2$. Using this approach one can derive the following equation for the surface plasmon generation by a single straight slit mode.^{36,37}

$$|\alpha|^2 = f \left(\frac{w}{\lambda} \right) \frac{n_1}{n_2} |\epsilon|^{-1/2} \quad (3)$$

with the assumption that the metal is a perfect conductor. This assumption simplifies the dependence of the geometric diffraction on the dielectric properties of the metal, and the dependence of the bounded SPP mode on α on the dielectric properties of the metal–dielectric interface.³⁸ In eqn (3), λ is the wavelength of the incident light, ϵ is the dielectric constant, and n_1 and n_2 represent two refractive indexes of the two media on either side of the interface (see Fig. 1a).

Combining the SPP fundamental mode with transmission mode in the plane wave basis, we can obtain the SPP generation efficiencies on both sides of the aperture, under the assumption of $\alpha^+(-w/2) = \alpha^-(w/2) = 0$, regarding the electromagnetic field below and inside the slit. By

generalizing the procedure used by Lalanne for a single straight slit (Fig. 1b),^{37,39,40} one can obtain the SPP generation efficiencies e at both sides of the apertures on the three Au interfaces; see Fig. 1a. For the Au/quartz interface e (red) is given by:

$$e_1 = \left| \alpha_1^+(w_1/2) \right|^2 = \left| \alpha_1^-(w_1/2) \right|^2 = \frac{4w_1' n_1^3}{\pi n_2^2} \left| \frac{\varepsilon^{1/2}}{\varepsilon + n_1^2} \left\| \frac{I_1}{1 + (n_1/n_2) w_1' I_0} \right\|^2 \right. \quad (4)$$

in which

$$I_0 = \int_{-\infty}^{\infty} du \left[\frac{\sin(\pi w_1' u)}{\pi w_1' u} \right]^2 / v \quad (5)$$

$$I_1 = \int_{-\infty}^{\infty} du \frac{\exp(-i\pi w_1' u) \sin(\pi w_1' u) / (\pi w_1' u)}{v \left\{ v + \left[n_1^2 / (\varepsilon + n_1^2) \right]^{1/2} \right\}} \quad (6)$$

$w_1' = n_1 w_1 / \lambda$ (7) where w' represents the scaled width and u and v are applied for numerical integration with $u^2 + v^2 = 1$.

For the slit mode case, the corresponding SPP efficiencies can be obtained by the following equation:³⁷

$$\left| \alpha(w/2) \right|_{\text{slit-mode}}^2 = \frac{N_p}{N_0} \left| \alpha(w/2) \right|_{\text{plane-wave}}^2 \quad (8)$$

with the assumption that the forward and backward fundamental modes compose the field in the slit and the normalization constants N_0 and N_p are given by $N_0 = w / (2\varepsilon n_2)$ and $N_p = w / (2\varepsilon n_1)$. For the inside Au/medium interface of the ledge (green), e is expressed as:

$$e_2 = \left| \alpha_2^+(w_1/2) \right|^2 = \left| \alpha_2^-(w_1/2) \right|^2 = \frac{4w_2' n_3^2}{\pi n_2} \left| \frac{\varepsilon^{1/2}}{\varepsilon + n_3^2} \left\| \frac{I_1'}{1 + (n_3/n_2) w_2' I_0'} \right\|^2 \right. \quad (9)$$

with

$$I_0' = \int_{-\infty}^{\infty} du \left[\frac{\sin(\pi w_2' u)}{\pi w_2' u} \right]^2 / v \quad (10)$$

$$I_1' = \int_{-\infty}^{\infty} du \frac{\exp(-i\pi w_2' u) \sin(\pi w_2' u) / (\pi w_2' u)}{v \left\{ v + \left[n_4^2 / (\varepsilon + n_4^2) \right]^{1/2} \right\}} \quad (11)$$

$$w_2' = n_2 w_1 / \lambda \quad (12)$$

Similarly, for the Au/medium interface at the top of the nanoledge aperture (blue), e is given by

$$e_3 = \left| \alpha_3^+(w_2/2) \right|^2 = \left| \alpha_3^-(w_2/2) \right|^2 = \frac{4w_3'n_4{}^2}{\pi n_3} \left| \frac{\varepsilon^{1/2}}{\varepsilon + n_4{}^2} \left\| \frac{I_1''}{1 + (n_4/n_3)w_3'I_0''} \right\| \right|^2 \quad (13)$$

with

$$I_0'' = \int_{-\infty}^{\infty} du \left[\frac{\sin(\pi w_3'u)}{\pi w_3'u} \right]^2 / v \quad (14)$$

$$I_1'' = \int_{-\infty}^{\infty} du \frac{\exp(-i\pi w_3'u) \sin(\pi w_3'u) / (\pi w_3'u)}{v \left\{ v + \left[n_4{}^2 / (\varepsilon + n_4{}^2) \right]^{1/2} \right\}} \quad (15)$$

$$w_3' = n_3 w_2 / \lambda \quad (16)$$

Details of the derivation of these results are provided in the ESI.† Because the full integrand, weighted by v , is singular over the interval of -1 and $+1$ and complex for $|u| > 1$, the integrals $I_0(I_0, I_0'')$, and $I_1(I_1, I_1'')$, were calculated numerically.^{37,41} Tables S1–S4† provide numerical results for different values of the normalized slit width w' and wavelength λ with the corresponding dielectric constant values of quartz.

Numerical simulation

The metal's dielectric response was modeled by a Drude–Lorentz model,⁴²

$$\varepsilon_r = \varepsilon_\infty + \sum_{n=1}^N \frac{x_0 G_n \omega_{0n}{}^2}{\omega_{0n}{}^2 + i\Gamma_n \omega - \omega^2} \text{ and } \sum_{n=1}^N G_n = 1 \quad (17)$$

where ε is the permittivity, ω_{0n} is the resonant frequency, Γ_n is the damping coefficient, and x_0 is the permittivity at ω_0 . By using a single resonance and the Fourier transform of the polarization in the algorithm, the FDTD formalism was used to calculate the transverse electric (TE) and transverse magnetic (TM) fields.²⁸

The geometry of the nanoledge structure was modeled in three-dimensional (3D) environments. The dielectric function of Au used in the simulations was from Johnson and Christy⁴³ and the substrate was simulated as an infinite block with a dielectric constant of silicon dioxide taken from Palik.⁴⁴ The simulations were performed using a single aperture as the unit cell with periodic boundary conditions in the y -direction to describe an infinite rectangular array and perfectly matched layers in the boundary along the x and z -direction. The period of the nanoslit array is 600 nm and the heights of the two slits are 50 nm and 150 nm, respectively. Note that the period and height are chosen to correspond to our previous work with nanoplasmonic nanofluidics.⁴⁵ As with the experiments, the calculations consider a linearly polarized broadband plane wave source, which propagates through the quartz substrate and is incident on the back surface of the gold layer at normal incidence.

Device fabrication and testing

Subwavelength nanoledge structures were fabricated using electron beam lithography (EBL) and focused ion beam (FIB) techniques. Sample imaging was done using an SEM in the nanofabrication systems. Spectral characterization of the nanoledge arrays was carried out using a microspectrophotometer (Craic QDI 2010).⁴⁶ The details of the experimental procedures and supporting data are included in the ESI.†

Conclusions

In conclusion, we extended a semi-analytical model to perform a decomposition analysis of the SPP wave generation at metallic interfaces perforated by a subwavelength gold nanoledge structure. The factor analysis of parameters (geometric, dielectric-RI, and incident wavelength) relevant to surface plasmon (SP) generation has been quantitatively investigated for the prediction of surface plasmon polariton (SPP) generation efficiency. The rigorous formalism for the model has been validated by comparisons with the FDTD modelling of the EOT and its sensitivity of RI changes and by experimental testing of fabricated nanoledge devices through measurement of their optical transmission and RI sensitivity. The analysis shows that SPP-generation is very efficient for the gold film nanoledge device. The reported semi-analytical approach provides a new tool for a quantitative decomposition analysis of SPP generation in other related slit structures and should prove useful for plasmonic device development.

Acknowledgements

J. W. and Z. Z. acknowledge the financial support from US NSF (Grant # 1511194). J. W. and Z. Z. also kindly acknowledge the support from JSNN. DHW and MM acknowledge financial support from NIH Grant # 1R21RR026221-01.

Footnote

† Electronic supplementary information (ESI) available: Additional mathematical derivation, programming and data analysis, experimental description, supporting figures and tables. See DOI: [10.1039/c6ra01105e](https://doi.org/10.1039/c6ra01105e)

Notes and references

1. S. Lal , S. Link and N. J. Halas , *Nat. Photonics*, 2007, **1** , 641 —648.
2. J. Li , C. Chen , L. Lagae and P. Van Dorpe , *J. Phys. Chem. C*, 2015, **119** , 29116 —29122.
3. R. Quidant and M. Kreuzer , *Nat. Nanotechnol.*, 2010, **5** , 762 —763.
4. M. I. Stockman *Opt. Express*, 2011, **19** , 22029 —22106.
5. M. A. Beuwer , M. W. J. Prins and P. Zijlstra , *Nano Lett.*, 2015, **15** , 3507 —3511.
6. A. A. Al Balushi and R. Gordon , *ACS Photonics*, 2014, **1** , 389 —393.
7. A. G. Brolo *Nat. Photonics*, 2012, **6** , 709 —713.

8. M. E. Stewart , C. R. Anderton , L. B. Thompson , J. Maria , S. K. Gray , J. A. Rogers and R. G. Nuzzo , *Chem. Rev.*, 2008, **108** , 494 —521.
9. O. Tokel , F. Inci and U. Demirci , *Chem. Rev.*, 2014, **114** , 5728 —5752.
10. M. Couture , L. S. Live , A. Dhawan and J.-F. Masson , *Analyst*, 2012, **137** , 4162 —4170.
11. E. Hendry , R. V. Mikhaylovskiy , L. D. Barron , M. Kadodwala and T. J. Davis , *Nano Lett.*, 2012, **12** , 3640 —3644.
12. K.-L. Lee , P.-W. Chen , S.-H. Wu , J.-B. Huang , S.-Y. Yang and P.-K. Wei , *ACS Nano*, 2012, **6** , 2931 —2939.
13. T. W. Ebbesen , H. J. Lezec , H. F. Ghaemi , T. Thio and P. A. Wolff , *Nature*, 1998, **391** , 667 —669.
14. A. A. Yanik , M. Huang , O. Kamohara , A. Artar , T. W. Geisbert , J. H. Connor and H. Altug , *Nano Lett.*, 2010, **10** , 4962 —4969.
15. T. G. Mayerhöfer , R. Knipper , U. Hübner , D. Cialla-May , K. Weber , H.-G. Meyer and J. Popp , *ACS Photonics*, 2015, **2** , 1567 —1575.
16. O. Hess , J. B. Pendry , S. A. Maier , R. F. Oulton , J. M. Hamm and K. L. Tsakmakidis , *Nat. Mater.*, 2012, **11** , 573 —584.
17. L. Feuz , P. Jönsson , M. P. Jonsson and F. Höök , *ACS Nano*, 2010, **4** , 2167 —2177.
18. S. S. Aćimović , M. A. Ortega , V. Sanz , J. Berthelot , J. L. Garcia-Cordero , J. Renger , S. J. Maerkl , M. P. Kreuzer and R. Quidant , *Nano Lett.*, 2014, **14** , 2636 —2641.
19. S.-H. Chang , S. Gray and G. Schatz , *Opt. Express*, 2005, **13** , 3150 —3165.
20. P. Lalanne and J. P. Hugonin , *Nat. Phys.*, 2006, **2** , 551 —556.
21. A. E. Schlather , N. Large , A. S. Urban , P. Nordlander and N. J. Halas , *Nano Lett.*, 2013, **13** , 3281 —3286.
22. F. J. Garcia-Vidal , L. Martin-Moreno , T. W. Ebbesen and L. Kuipers , *Rev. Mod. Phys.*, 2010, **82** , 729 —787.
23. P. Lalanne , J. P. Hugonin , H. T. Liu and B. Wang , *Surf. Sci. Rep.*, 2009, **64** , 453 —469.
24. H. Liu and P. Lalanne , *Nature*, 2008, **452** , 728 —731.
25. J. P. Litz , J. P. Camden and D. J. Masiello , *J. Phys. Chem. Lett.*, 2011, **2** , 1695 —1700.
26. T. Tanemura , K. C. Balram , D.-S. Ly-Gagnon , P. Wahl , J. S. White , M. L. Brongersma and D. A. B. Miller , *Nano Lett.*, 2011, **11** , 2693 —2698.
27. D. M. Solís , J. M. Taboada , F. Obelleiro , L. M. Liz-Marzán and F. J. García de Abajo , *ACS Nano*, 2014, **8** , 7559 —7570.
28. Z. Zeng , Y. Liu and J. Wei , *TrAC, Trends Anal. Chem.*, 2016, **75** , 162 —173.

29. L. Aigouy , P. Lalanne , J. P. Hugonin , G. Julié , V. Mathet and M. Mortier , *Phys. Rev. Lett.*, 2007, **98** , 153902.
30. E. Skudrzyk *The Foundations of Acoustics: Basic Mathematics and Basic Acoustics* , Springer-Verlag Wien, 1971.
31. *Handbook of Optical Constants of Solids* , Academic Press, Burlington, 1997.
32. P. Kline *An easy guide to factor analysis* , Taylor and Francis, Hoboken, NJ, 2014.
33. A. E. Cetin , D. Etezadi , B. C. Galarreta , M. P. Busson , Y. Eksioglu and H. Altug , *ACS Photonics*, 2015, **2** , 1167 —1174.
34. M. Sanders , Y. Lin , J. Wei , T. Bono and R. G. Lindquist , *Biosens. Bioelectron.*, 2014, **61** , 95 —101.
35. Y. S. Jung , J. Wuenschell , H. K. Kim , P. Kaur and D. H. Waldeck , *Opt. Express*, 2009, **17** , 16081 —16091.
36. P. Lalanne , J. P. Hugonin and J. C. Rodier , *Phys. Rev. Lett.*, 2005, **95** , 263902.
37. P. Lalanne , J. P. Hugonin and J. C. Rodier , *J. Opt. Soc. Am. A*, 2006, **23** , 1608 —1615.
38. W. Yao , S. Liu , H. Liao , Z. Li , C. Sun , J. Chen and Q. Gong , *Nano Lett.*, 2015, **15** , 3115 —3121.
39. P.-K. Wei , H.-L. Chou , Y.-R. Cheng , C.-H. Wei , W. Fann and J. O. Tegenfeldt , *Opt. Commun.*, 2005, **253** , 198 —204.
40. Y. Xie , A. Zakharian , J. Moloney and M. Mansuripur , *Opt. Express*, 2004, **12** , 6106 —6121.
41. P.Chimento, PHD Doctoral, Leiden University, 2013.
42. A. Vial , A.-S. Grimault , D. Macías , D. Barchiesi and M. L. de la Chapelle , *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **71** , 085416.
43. P. B. Johnson and R. W. Christy , *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1972, **6** , 4370 —4379.
44. H. R. Philipp *Handbook of Optical Constants of Solids* , E. D. Palik Academic Press, Burlington, 1997, pp. 749–763.
45. J. Wei , H. Song , S. Singhal , M. Kofke , M. Mendis and D. H. Waldeck , *ASME Proceedings | Micro/Nanofluidics and Lab-on-a-Chip* , Atlanta, GA, 2012.
46. M. N. Mendis , H. S. Mandal and D. H. Waldeck , *J. Phys. Chem. C*, 2013, **117** , 25693 —25703.