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Citation: *Journal of Applied Physics* **117**, 133107 (2015); doi: 10.1063/1.4916800

View online: <http://dx.doi.org/10.1063/1.4916800>

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Formation of a hybrid plasmonic waveguide mode probed by dispersion measurement

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(Received 27 January 2015; accepted 23 March 2015; published online 1 April 2015)

Hybrid waveguides, i.e., dielectric waveguides combined with plasmonic waveguides, have great potential for concomitantly exhibiting subwavelength confinement and long range propagation, enabling a highly integrated photonic circuit. We report the characterization of hybrid waveguide modes excited in Si/SiO₂/Al films, by dispersion measurement using angle-resolved electron energy-loss spectroscopy. This experiment directly verifies the formation of the hybrid waveguide mode with a strongly localized electromagnetic field in a 6-nm-thick SiO₂ layer. The results clearly describe the characteristic behavior of the hybrid waveguide mode, which depends on the effective index of the constituent dielectric waveguide and the surface plasmon-polariton modes. © 2015 AIP Publishing LLC. [<http://dx.doi.org/10.1063/1.4916800>]

I. INTRODUCTION

Guiding light with subwavelength confinement is a critical challenge for nanoscale optical applications, such as integrated photonic circuits.¹ A variety of waveguide structures have been proposed for this purpose in previous reports,^{2–10} some of which have been assessed experimentally. Dielectric waveguide structures like dielectric cylinders can allow light propagation over a long distance, although the confinement is limited by the size of the waveguide structure itself. By contrast, waveguide structures using surface plasmon-polaritons (SPPs),^{11,12} such as a metal surface, can achieve deep subwavelength confinement. However, the propagation loss associated with plasmonic waveguides is high because of electron oscillations in the dissipative metallic regions,¹² and the propagation distance is usually limited to several microns.

One strategy that has been predicted to be efficient is coupling a dielectric waveguide with a plasmonic waveguide.^{13–18} A hybrid plasmonic waveguide with a dielectric cylinder above a metal surface was proposed,¹³ and its effectiveness as a nanolaser was demonstrated several years ago.^{19,20} The hybrid waveguide structure can store dense energy in the low-permittivity insulator nano-gap between the dielectric waveguide and the metal surface, leading simultaneously to deep subwavelength confinement and long-range propagation. This waveguide mode has been regarded as a coupling of the dielectric waveguide and the SPPs.^{13,18} Although the properties of hybrid waveguide modes have been analyzed theoretically, experimental verifications have so far been lacking. In particular, although the coupled-mode theory suggests that the character of the hybrid waveguide modes depends mainly on the effective indices of the constituent dielectric waveguide and SPP modes,^{13,18,21} this relationship has not yet been confirmed satisfactorily by experiment. In the present work, we show the results of dispersion

measurements of the hybrid waveguide mode using angle-resolved electron energy-loss spectroscopy (AREELS).^{22–29} In order to investigate the fundamental characteristics of the simplest hybrid waveguide structure, we fabricated three-layered films consisting of a high-permittivity semiconductor layer, a low-permittivity insulator nano-gap (or vacuum) and a metal layer, and a so-called SIM structure. Owing to this simple structure, the dispersion relations observed experimentally can easily be compared with the calculated relationships. Furthermore, the three-layered film with parallel interfaces can be used in real applications, such as integrated photonic circuits,^{14–17} because a structure with a rectangular cross-section is much easier to fabricate using microfabrication technology, e.g., electron beam lithography.

We performed dispersion measurements not only on the SIM films but also on the SI films (without the metal layer) over the energy region encompassing the visible light spectrum to determine the relationship between the effective indices of the constituent modes and the behavior of the hybrid waveguide mode. The low-permittivity insulator gap in the present study was less than 10 nm thick. The formation of a hybrid waveguide mode with extremely dense electromagnetic energy was confirmed through dispersion measurements.

II. METHOD

Figure 1 shows a schematic diagram of AREELS. A narrow slit is placed at the entrance plane of the spectrometer on which the angular dispersive (electron diffraction) pattern is formed. When the slit is set to be perpendicular to the direction of energy dispersion for the spectrometer, the spectral intensity can be obtained as a function of the scattering angle (θ) and the energy loss (E). Since the scattering angle is proportional to the component of the scattering vector perpendicular to the direction of incident electrons (q_{\perp}), the observed two-dimensional (2D) AREELS pattern allows us to directly visualize the energy-loss probability as a function of E and q_{\perp} (E-q map), as shown in Fig. 1. When the

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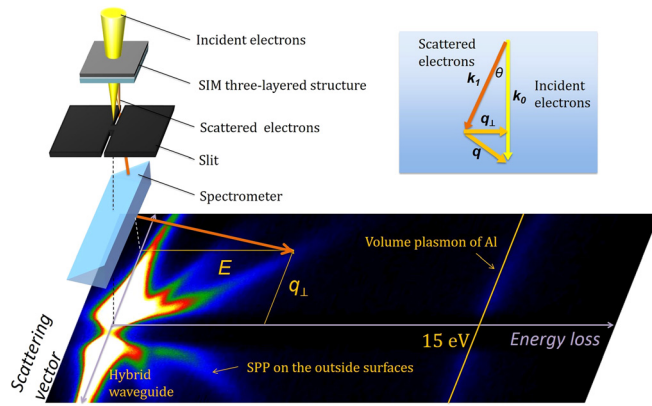


FIG. 1. Schematic diagram of AREELS method. \mathbf{k}_0 and \mathbf{k}_1 represent the wave vectors of incident and scattered electrons, respectively. An angular selection slit is placed at the entrance plane of the spectrometer to limit electron collection to those scattered perpendicular to the direction of energy dispersion. The gap in the center part of the slit is closed to prevent the intense direct beam from saturating the CCD detector. The inserted E-q map is taken from B2.

incident electrons are normal to the multilayered films, q_{\perp} can be assumed to be the wave vector k (propagation constant) of the propagating waves in waveguide structures. Therefore, the E-q map directly yields the dispersion relations of the waveguide modes.

AREELS measurements were performed at room temperature using a 200 kV transmission electron microscope (JEM-9980 TKP1) equipped with a cold field emission gun and an omega filter as a spectrometer. The energy resolution determined from the full width at half maximum of the zero-loss peak was 0.4 eV. In order to achieve the high angular dispersion, the specimen was lifted from the normal position in the image mode with spot illumination.²⁶ The scattering

vector resolution determined from the full width at half maximum of a direct beam on the E-q map was $5 \times 10^{-3} \text{ nm}^{-1}$. A specimen area $4 \mu\text{m}$ in diameter was illuminated by an incident electron beam with a convergent semi-angle of 6.4 mrad. The E-q maps were measured by a charge-coupled device (CCD) camera with a pixel size corresponding to 27 meV and $3.8 \times 10^{-4} \text{ nm}^{-1}$ along the loss energy and the scattering vector axes in the E-q maps, respectively. Each E-q map was acquired in 10 s. Ten E-q maps taken under the same conditions were superimposed to improve the signal-to-noise ratio.

The waveguide structures consisting of the multilayered (SI and SIM) films were prepared as shown in Fig. 2(a). Silicon (Si), silicon dioxide (SiO_2), and aluminum (Al) were chosen as the semiconductor layer, the insulator gap, and the metal layer, respectively. First, a 6-nm-thick amorphous SiO_2 film was produced on the sodium chloride crystal (NaCl) substrate by electron beam evaporation. Then, a non-crystalline Si film with a thickness of 52, 105, or 157 nm was deposited on the $\text{SiO}_2(6 \text{ nm})/\text{NaCl}$ substrate by electron beam evaporation. The resulting two-layered films, viz., Si(52 nm)/ $\text{SiO}_2(6 \text{ nm})$, Si(105 nm)/ $\text{SiO}_2(6 \text{ nm})$, and Si(157 nm)/ $\text{SiO}_2(6 \text{ nm})$, are denoted as specimens A1, A2, and A3, respectively. Specimens A1, A2, and A3 were floated on the surface of distilled water by dissolving the NaCl substrates, and transferred onto copper meshes for AREELS measurements. Finally, 35-nm-thick Al films were deposited on the SiO_2 side of specimens A1, A2, and A3 by thermal evaporation; the resulting specimens will hereafter be denoted as specimens B1, B2, and B3, respectively. The thickness of each layer was monitored with a quartz oscillator and confirmed by cross-sectional images taken by a scanning transmission electron microscope, as shown in Fig. 2(b). The total thicknesses of the specimens were small enough to be penetrated by the incident electrons.

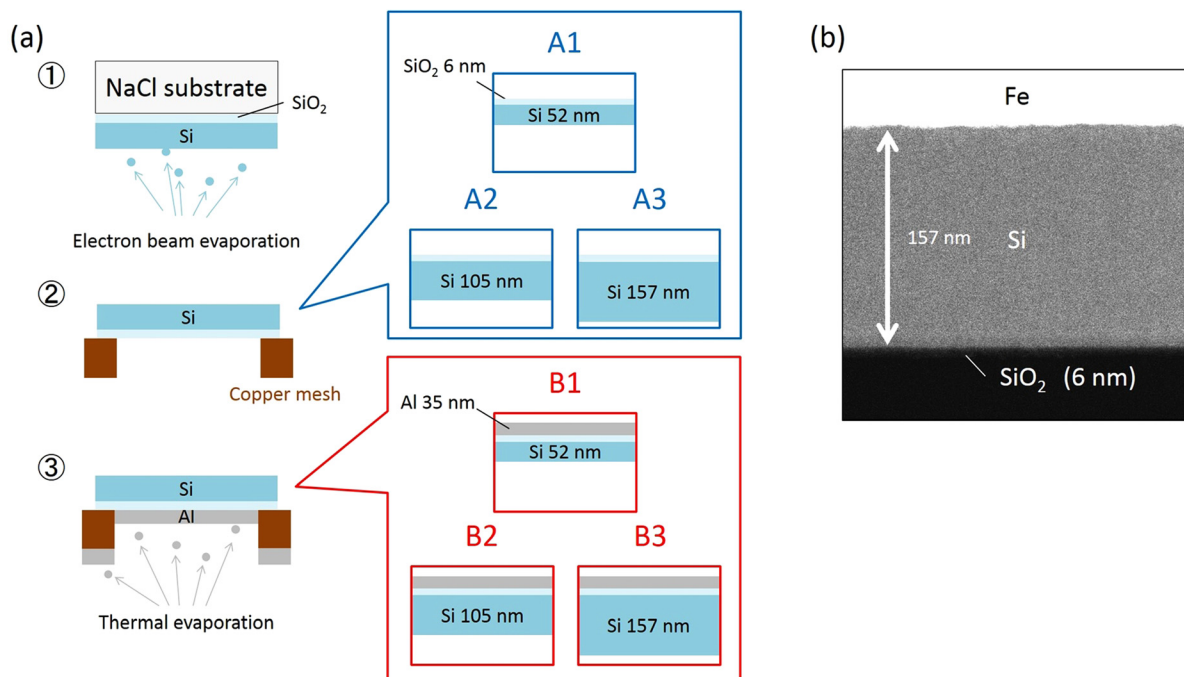


FIG. 2. (a) Schematics of the specimens and the steps of specimen preparation. (b) A cross-sectional STEM image of A3. The brightest region is the iron (Fe) layer deposited to protect the specimen from being damaged by FIB processing.

Since the SPP mode supported by the plane surface is the transverse magnetic (TM) mode, the hybrid waveguide modes supported by the SIM film are also expected to be only TM modes. Moreover, the incident electrons normal to the plane surface can excite only TM modes.³⁰ Therefore, in the present study, we consider the dispersion relations only for TM modes. The experimental E-q maps were compared with the theoretical dispersion curves for TM modes to identify their modes. By solving Maxwell's equations under the appropriate boundary conditions, the dispersion relationship is obtained; it can be expressed in terms of the dielectric function (ϵ) and the thickness (t) of each film via the equation

$$1 + K_{m,i}K_{i,s}e^{2\alpha_i t_i} + K_{i,s}K_{s,0}e^{2\alpha_s t_s} + K_{m,i}K_{s,0}e^{2\alpha_i t_i}e^{2\alpha_s t_s} = 0, \quad (1)$$

where

$$K_{m,i} = \frac{\frac{\alpha_m}{\epsilon_m} + \frac{\alpha_i}{\epsilon_i}}{\frac{\alpha_m}{\epsilon_m} - \frac{\alpha_i}{\epsilon_i}}, \quad (2)$$

$$K_{i,s} = \frac{\frac{\alpha_i}{\epsilon_i} - \frac{\alpha_s}{\epsilon_s}}{\frac{\alpha_i}{\epsilon_i} + \frac{\alpha_s}{\epsilon_s}}, \quad (3)$$

$$K_{s,0} = \frac{\frac{\alpha_s}{\epsilon_s} - \frac{\alpha_0}{\epsilon_0}}{\frac{\alpha_s}{\epsilon_s} + \frac{\alpha_0}{\epsilon_0}}. \quad (4)$$

The indices m , i , s , and 0 denote the metal layer (Al), the insulator gap (SiO₂), the semiconductor layer (Si), and vacuum, respectively. The thickness of the metal layer is assumed to be semi-infinite in this equation. The metal layer changes into vacuum for the dispersion calculation of the SI films. The parameter α , i.e., the transverse wave number (damping factor) of the electromagnetic wave perpendicular to the interface, is given by

$$\alpha_n = \sqrt{k^2 - \epsilon_n \left(\frac{E}{\hbar c}\right)^2} \quad (n = m, i, s, 0), \quad (5)$$

where \hbar and c are the Planck constant divided by 2π and the velocity of light in vacuum, respectively.

The dispersion relations were calculated by solving Eq. (1) numerically. The dielectric function of SiO₂ determined optically was used for the calculation.³¹ The dielectric function of Al was assumed to follow the Drude model without damping

$$\epsilon_m(E) = 1 - \frac{E_p^2}{E^2}, \quad (6)$$

where E_p is the excitation energy for bulk plasmons, which is 15 eV for Al.²³ The reliability of these dielectric functions of SiO₂ and Al was confirmed in our previous work.²⁸ The dielectric function of non-crystalline Si was determined by the present experiment for the SI films (specimens A2 and A3).

III. RESULTS

Figure 3 shows the E-q maps of the SI films (A1, A2, and A3) and SIM films (B1, B2, and B3), in which they are compared with the calculated dispersion relations of light in vacuum, bulk amorphous SiO₂, and Si crystal (white solid lines and curves). Since the real part of the dielectric function of Si is positive and large in the energy range below 4 eV, Čerenkov radiation is generated in Si by 200-keV electrons. This radiation is confined inside the Si slab as discrete waveguide modes²⁷ owing to the boundary effect.³² Therefore, dispersion curves of Si waveguide modes would be expected to be observed below 4 eV in the E-q maps of the SI films. In fact, on the E-q map in Fig. 3(b), the dispersion curve of the lowest TM mode (TM₀) appears in the region between the light lines in vacuum and Si bulk. This is a feature of Si waveguide modes. When the thickness of the Si film is increased, dispersion curves of two TM modes are observed, as shown in Fig. 3(c). The dispersion curve of the TM₀ mode shifts to the lower energies with increasing Si thickness. This behavior is characteristic of dielectric waveguide modes. For the thin Si film (A1), no dispersion curve of the Si waveguide mode was detected below 4 eV, as shown in Fig. 3(a). The suppression of the Si waveguide modes in A1 can be attributed to the decrease in Čerenkov loss whose probability is roughly proportional to the thickness.^{33–35}

On the E-q maps of all SI films shown in Figs. 3(a)–3(c), a broad dispersion curve is observed outside the dispersion relation of light in vacuum above 4 eV. This is the dispersion curve for SPPs excited on the Si/vacuum and Si/SiO₂ interfaces. Since the real part of the dielectric function of Si has a negative value between 4 eV and the volume plasmon energy (16–17 eV), SPPs can be excited on the interface between Si and the insulator (or vacuum) in this energy region. The surface on the vacuum side was presumably oxidized slightly because all specimens were exposed to air for a long time before the AREELS experiments. Therefore, the broad dispersion curve above 4 eV is attributed to SPP on two interfaces.

Figures 3(d)–3(f) show the E-q maps measured from SIM films (B1, B2, and B3). The dispersion curves observed below 4 eV are slightly different from those of the SI films. For B1, in particular, a new dispersion curve clearly appears in the E-q map of Fig. 3(d), whereas no dispersion curve was detected for the corresponding SI film (A1), as seen in Fig. 3(a). This dispersion curve might be related to the SPP excited on the Al/SiO₂ interface. Moreover, when the thickness of the Si film in SIM structures is increased, the dispersion curves shift to lower energies (see Figs. 3(e) and 3(f)), similar to the case of SI films. In order to distinguish the dispersion relations of SIM and SI films, their angle-resolved EELS spectra at $q_{\perp} = 0.02, 0.03, \text{ and } 0.04 \text{ nm}^{-1}$ were compared. Figure 4(a) shows the angle-resolved EELS spectra extracted from the E-q maps of Figs. 3(b) and 3(e). The new peaks observed in the SIM film (B2) are lower than those of the Si waveguide (TM₀) mode in the SI film (A2), which suggests that the new excitation mode arises from the coupling between the SPPs on the Al/SiO₂ interface and the TM₀ modes in the Si film, representing a so-called hybrid

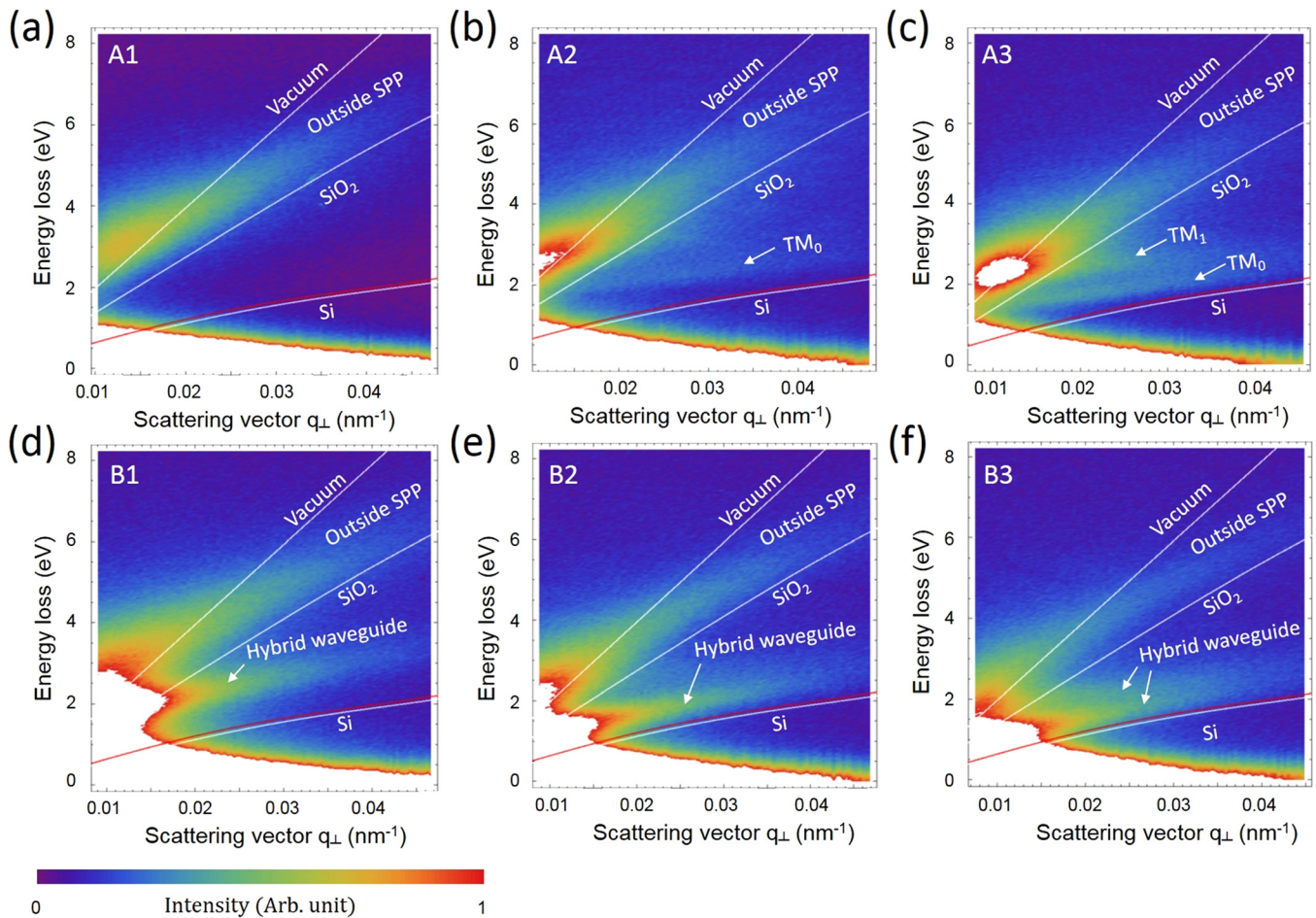


FIG. 3. E-q maps of (a) A1, (b) A2, (c) A3, (d) B1, (e) B2, and (f) B3. The white solid curves and line are the calculated dispersion relations of light in bulk SiO₂ and vacuum. The red curve represents the relationship between the energy and q_{\perp} of Čerenkov radiation excited by 200 keV electrons in bulk Si.

waveguide mode. It should be noted that the energy difference between the hybrid waveguide and the TM₀ modes becomes small at large wave vectors ($q_{\perp} = 0.04 \text{ nm}^{-1}$). Figure 4(b) shows the angle-resolved EELS spectra extracted

from the E-q maps of Figs. 3(c) and 3(f), where the second mode of the hybrid waveguide coupled with the TM₁ mode of the Si waveguide also appears as weak shoulder peaks in the spectra of B3. The first and second modes of the hybrid

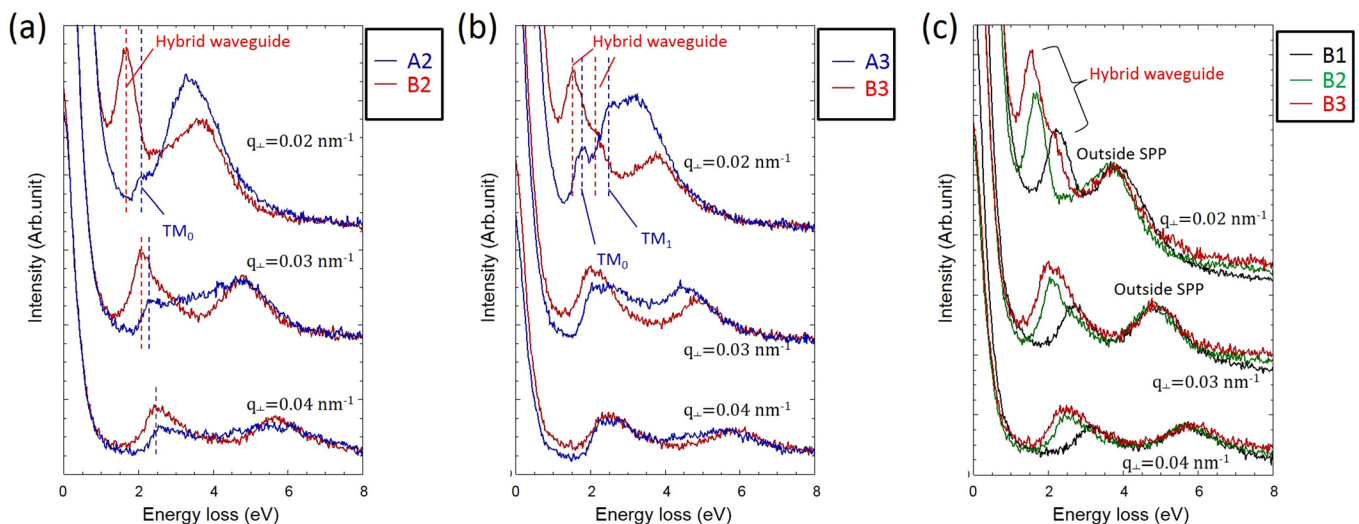


FIG. 4. Angle-resolved EELS obtained from the E-q maps of (a) A2 (blue) and B2 (red), and (b) A3 (blue) and B3 (red) at $q_{\perp} = 0.02 \text{ nm}^{-1}$, 0.03 nm^{-1} , and 0.04 nm^{-1} . (c) Comparison of the angle-resolved EELS of B1 (black), B2 (green), and B3 (red).

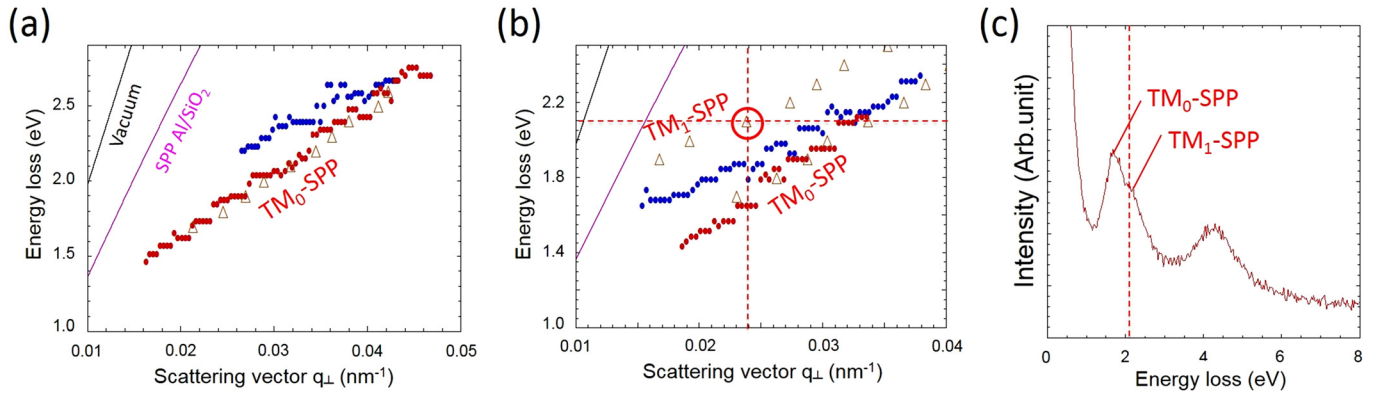


FIG. 5. (a) and (b) Dispersion plots of the hybrid waveguide mode (red filled circles) obtained from (a) B2 and (b) B3, compared with the theoretical dispersion plots (brown open triangles). The dispersion relations of the TM_0 mode extracted from A2 and A3 are shown by blue filled circles. The solid purple curve and black line are the theoretical dispersion relations of SPPs on the Al/SiO₂ interface and of light in vacuum, respectively. (c) Angle-resolved EELS of B3 at $q_{\perp} = 0.024 \text{ nm}^{-1}$.

waveguide shift to lower energies relative to the TM_0 and TM_1 modes of the Si waveguide, respectively. Their differences also decrease with increasing q_{\perp} .

The angle-resolved EELS spectra extracted from the E-q maps of all SIM structures are compared in Fig. 4(c). The intensity of each spectrum is normalized at the peak of the outside SPPs. It should be noted that the ratio of excitation probability of the hybrid waveguide modes to that of the outside SPPs clearly increases with increasing Si layer thickness. This behavior is typical of volume excitations like Čerenkov radiation.^{33–35} Thus, the intensity increases with Si thickness is attributed to the enhancement of the hybrid waveguide modes by coupling with Čerenkov radiation, similar to the abovementioned case for Si waveguide modes.

For accurate assignment of the hybrid waveguide modes, the dispersion relation observed experimentally should be compared to that calculated by solving Eq. (1) numerically. Figures 5(a) and 5(b) compare experimental and calculated dispersion relations. The experimental dispersions (red filled circles) were extracted from the E-q maps in Figs. 3(e) and 3(f) by finding a local maximum in the energy region from 1.45 eV to 2.75 eV and 2.15 eV, respectively. These experimental plots agree well with the calculated dispersion (red open triangles) of the hybrid waveguide mode formed from the coupling of the TM_0 waveguide and SPP mode (TM_0 -SPP mode) with a quasi-even symmetry. For comparison, these figures also plot the dispersion relations of the Si waveguide TM_0 mode extracted from Figs. 3(b) and 3(c) (see blue filled circles). It can be seen that the dispersion for the TM_0 -SPP mode merges with that of the TM_0 mode at large wave vectors. Also seen in Fig. 5(b) on the high-energy side is the theoretical dispersion for the TM_1 -SPP mode with quasi-even symmetry. Although it is difficult to extract the experimental dispersion relation for the TM_1 -SPP mode from the E-q map of Fig. 3(f), the angle-resolved spectrum for $q_{\perp} = 0.024 \text{ nm}^{-1}$ has a shoulder at 2.1 eV as shown in Fig. 5(c). The position of this shoulder agrees with the theoretical prediction shown in Fig. 5(b), so the existence of the TM_1 -SPP mode with quasi-even symmetry is verified experimentally.

IV. DISCUSSION

Hybrid waveguide modes have been demonstrated in SIM waveguide structures in the visible wavelength region by measuring the dispersion relations directly using AREELS. The dispersion curve of the TM_0 -SPP hybrid mode shown in Fig. 5(a) is distinct from those of the TM_0 and SPP modes at low energy, but merged with that of the TM_0 mode at high energy. This suggests that the character of a hybrid waveguide mode depends on the energy. A hybrid mode can be characterized in terms of the effective index, defined as the propagation constant of waveguide mode divided by the wave vector in vacuum. Since the normal component of the scattering vector is equal to the propagation constant, it is easy to convert an E-q map into a dispersion relation of the effective index.

Figure 6(a) shows the effective indices for the TM_0 -SPP hybrid mode, the TM_0 mode supported by a Si film, and the SPP mode supported by an aluminum film as a function of energy, which is converted from the E-q relationship of Fig. 5(a). Since the effective index for the TM_0 mode below 2.2 eV could not be determined experimentally owing to the weak intensity in the E-q map of Fig. 4(b), it was calculated by using the dielectric function evaluated from the results of Fig. 3(c). The effective indices for the TM_0 and SPP modes intersect at about 1.8 eV, which is the critical energy where the effective index of the TM_0 -SPP hybrid mode differs greatly from the average of the effective indices of the TM_0 and SPP modes. The SPPs at the Al/SiO₂ interface and the polarized charges in the Si waveguide oscillate in phase at this critical energy.

A simple coupling mode theory suggests that the maximum coupling occurs at this critical energy and the character of the hybrid mode is governed equally by both modes,^{13,21} so that it can be expected that the electromagnetic energy will be confined at the insulator gap region effectively. The electromagnetic energy density per unit area in the propagation plane is given by the following equation:¹³

$$W = \frac{1}{2} \left(\frac{d(\epsilon\omega)}{d\omega} |\mathbf{E}|^2 + \mu_0 |\mathbf{H}|^2 \right), \quad (7)$$

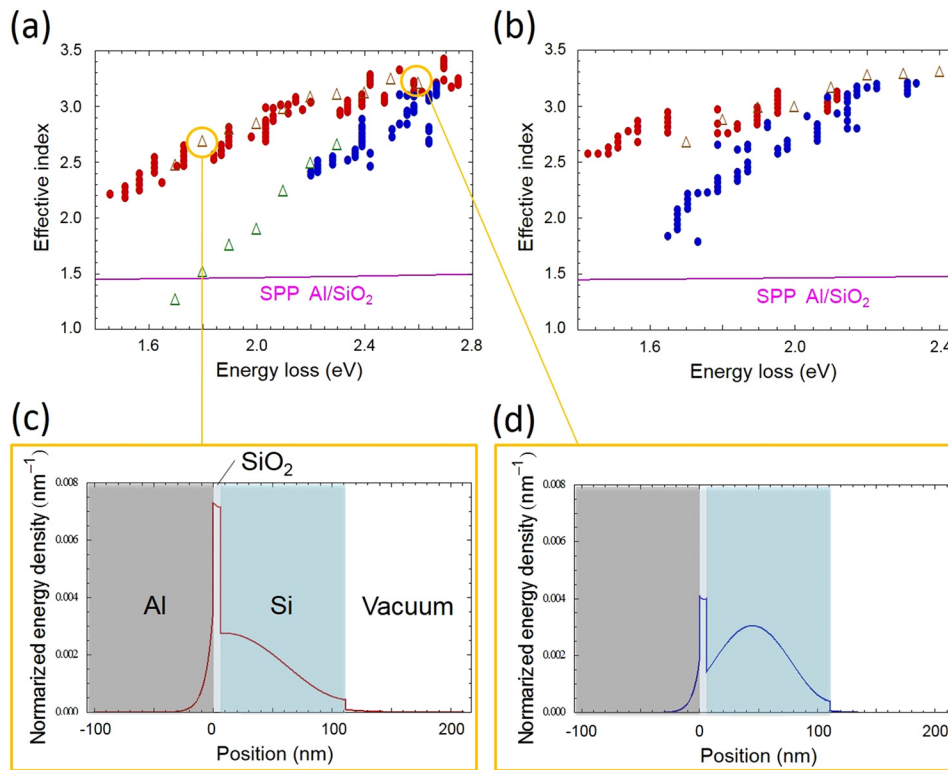


FIG. 6. (a) and (b) Energy dependence of the effective indices for (a) A2 and B2, and (b) A3 and B3 calculated from the dispersion plots (Figs. 5(a) and 5(b)). The red and blue filled circles represent the TM_0 -SPP hybrid mode and the TM_0 mode obtained in the AREELS experiment, respectively. The brown and green open triangles represent the theoretical indices. The solid purple curve is the theoretical indices of SPP on the Al/SiO₂ interface. (c) and (d) Normalized electromagnetic energy density for the TM_0 -SPP mode in B2 calculated from the theoretical dispersion relation (c) near the intersection (1.8 eV) and (d) on the high energy side (2.6 eV).

where ω and μ_0 are the angular frequency and permeability of vacuum, respectively. The expressions for the electric field \mathbf{E} and magnetic field \mathbf{H} are given by solutions of Eq. (1). Figure 6(c) shows the electromagnetic energy density calculated for the SIM structure (specimen B2) at 1.8 eV, which is normalized by the total electromagnetic energy calculated by the integration of W along the normal to the interface. The characteristics of the hybrid waveguide mode with a strong energy confinement at the gap region can be seen in this figure. With increasing energy, the difference between the effective indices for the TM_0 -SPP hybrid mode and the TM_0 mode becomes small, which means that the hybrid mode becomes TM_0 -like. Indeed, the electromagnetic energy density calculated at 2.6 eV (Fig. 6(d)) shows a larger contribution in the Si waveguide region and a smaller energy confinement at the gap region, as compared to Fig. 6(c). At energies below 1.8 eV, the effective index for the TM_0 mode is smaller than that for the SPP mode, and the hybrid mode is expected to become SPP-like. The dispersion for the effective index for the SIM structure of specimen B3 with a thick Si layer is shown in Fig. 6(b). With increasing Si waveguide layer thickness, the critical energy giving maximum coupling seems to be decrease, which suggests that the energy of effective hybrid coupling can be controlled by adjusting the thickness of the dielectric waveguide layer.

V. CONCLUSION

The present dispersion measurements conducted on Si/SiO₂/Al structures by AREELS demonstrated that the SPP mode excited in the interface of Al/SiO₂ can couple with the TM_0 waveguide mode excited in the Si layer, leading to the TM_0 -SPP hybrid waveguide mode. The dispersion of the effective indices for the constituent modes showed that the

maximum coupling occurs at the critical energy. Our experiment with SIM structures having different semiconductor thicknesses indicated that the critical energy for a hybrid waveguide mode decreased with increasing thickness. The electromagnetic energy of a hybrid waveguide mode excited at the critical energy was found to be confined effectively to the insulator gap region. Our results clearly demonstrated that the character and critical energy of a hybrid waveguide mode can be controlled by adjusting the thickness of the semiconductor layer. Finally, we verified the TM_1 -SPP mode with quasi-even symmetry experimentally.

- ¹D. K. Gramotnev and S. I. Bozhevolnyi, *Nat. Photonics* **4**, 83 (2010).
- ²J. Takahara, S. Yamagishi, H. Taki, A. Morimoto, and T. Kobayashi, *Opt. Lett.* **22**, 475 (1997).
- ³P. Berini, *Phys. Rev. B* **63**, 125417 (2001).
- ⁴K. Tanaka and M. Tanaka, *Appl. Phys. Lett.* **82**, 1158 (2003).
- ⁵A. D. Boardman, G. C. Aers, and R. Teshima, *Phys. Rev. B* **24**, 5703 (1981).
- ⁶I. V. Novikov and A. A. Maradudin, *Phys. Rev. B* **66**, 035403 (2002).
- ⁷V. R. Almeida, Q. Xu, C. A. Barrios, and M. Lipson, *Opt. Lett.* **29**, 1209 (2004).
- ⁸R. Zia, J. A. Schuller, and M. L. Brongersma, *Phys. Rev. B* **74**, 165415 (2006).
- ⁹D. F. P. Pile, T. Ogawa, D. K. Gramotnev, T. Okamoto, M. Haraguchi, M. Fukui, and S. Matsuo, *Appl. Phys. Lett.* **87**, 061106 (2005).
- ¹⁰S. I. Bozhevolnyi, V. S. Volkov, E. Devaux, and T. W. Ebbesen, *Phys. Rev. Lett.* **95**, 046802 (2005).
- ¹¹R. H. Ritchie, *Phys. Rev.* **106**, 874 (1957).
- ¹²W. L. Barnes, A. Dereux, and T. W. Ebbesen, *Nature* **424**, 824 (2003).
- ¹³R. F. Oulton, V. J. Sorger, D. A. Genov, D. F. P. Pile, and X. A. Zhang, *Nat. Photonics* **2**, 496 (2008).
- ¹⁴R. Salvador and A. Martínez, *IEEE J. Sel. Top. Quantum Electron.* **14**, 1496 (2008).
- ¹⁵M. Fujii, J. Leuthold, and W. Freude, *IEEE Photonics Technol. Lett.* **21**, 362 (2009).
- ¹⁶D. Dai and S. He, *Opt. Express* **17**, 16646 (2009).
- ¹⁷M. Z. Alam, J. Meier, J. S. Aitchison, and M. Mojahedi, *Opt. Express* **18**, 12971 (2010).

- ¹⁸M. Z. Alam, J. S. Aitchison, and M. Mojahedi, *IEEE J. Sel. Top. Quantum Electron.* **19**, 4602008 (2013).
- ¹⁹R. F. Oulton, V. J. Sorger, T. Zentgraf, R. M. Ma, C. Gladden, L. Dai, G. Bartal, and X. Zhang, *Nature* **461**, 629 (2009).
- ²⁰R. M. Ma, R. F. Oulton, V. J. Sorger, G. Bartal, and X. Zhang, *Nat. Mater.* **10**, 110 (2011).
- ²¹A. W. Snyder and J. D. Love, *Optical Waveguide Theory* (Chapman and Hall, London, New York, 1983).
- ²²H. Watanabe, *J. Phys. Soc. Jpn.* **11**, 112 (1956).
- ²³R. B. Pettit, J. Silcox, and R. Vincent, *Phys. Rev. B* **11**, 3116 (1975).
- ²⁴C. H. Chen, J. Silcox, and R. Vincent, *Phys. Rev. B* **12**, 64 (1975).
- ²⁵P. A. Midgley, *Ultramicroscopy* **76**, 91 (1999).
- ²⁶M. K. Kinyanjui, C. Kramberger, T. Pichler, J. C. Meyer, P. Wachsmuth, G. Benner, and U. Kaiser, *Europhys. Lett.* **97**, 57005 (2012).
- ²⁷H. Saito, C. H. Chen, and H. Kurata, *J. Appl. Phys.* **113**, 113509 (2013).
- ²⁸H. Saito, K. Namura, M. Suzuki, and H. Kurata, *Microscopy* **63**, 85 (2014).
- ²⁹S. C. Liou, C.-S. Shie, C. H. Chen, R. Breitwieser, W. W. Pai, G. Y. Guo, and M.-W. Chu, *Phys. Rev. B* **91**, 045418 (2015).
- ³⁰E. Kröger, *Z. Phys.* **235**, 403 (1970).
- ³¹H. R. Philipp, *J. Phys. Chem. Solids* **32**, 1935 (1971).
- ³²F. J. Garcia de Abajo, A. Rivacoba, N. Zabala, and N. Yamamoto, *Phys. Rev. B* **69**, 155420 (2004).
- ³³M. Stöger-Pollach, H. Franco, P. Schattschneider, S. Lazar, B. Schaffer, W. Grogger, and H. W. Zandbergen, *Micron* **37**, 396 (2006).
- ³⁴L. Gu, V. Srot, W. Sigle, C. Koch, P. van Aken, F. Scholz, S. B. Thapa, C. Kirchner, M. Jetter, and M. Rühle, *Phys. Rev. B* **75**, 195214 (2007).
- ³⁵R. Erni and N. D. Browning, *Ultramicroscopy* **108**, 84 (2008).