

Article

Epsilon-Near-Zero Absorber by Tamm Plasmon Polariton

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Abstract: Two schemes of excitation of a Tamm plasmon polariton localized at the interface between a photonic crystal and a nanocomposite with the near-zero effective permittivity have been investigated in the framework of the temporal coupled-mode theory. The parameters of the structure have been determined that correspond to the critical coupling of the incident field with a Tamm plasmon polariton and, consequently, ensure the total absorption of the incident radiation by the structure. It has been established that the spectral width of the absorption line depends on the scheme of Tamm plasmon polariton excitation and parameters of a nanocomposite film. The features of field localization at the Tamm plasmon polariton frequency for different excitation schemes have been examined. It has been demonstrated that such media can be used as narrowband absorbers based on Tamm plasmon polaritons localized at the interface between a photonic crystal and a nanocomposite with the near-zero effective permittivity.

Keywords: photonic crystal; nanocomposite; epsilon near zero material; Tamm plasmon polariton

1. Introduction

Tamm plasmon polaritons (TPPs) are a special type of the electromagnetic surface states, in which the field decays exponentially on each side of the surface [1] and the energy transfer along the surface can be stopped. This state can be experimentally observed as a narrow [2,3] or broadband [4] resonance in the optical transmission or reflection spectrum of a sample at wavelengths inside the band gap of a photonic crystal (PhC). Among the proposed and implemented TPP applications are organic solar cells [5], absorbers [6], lasers [7], sensors [8], integrated optical networks [9], heat emitters [10], and spontaneous emission amplifiers [11]. The high degree of field localization at the TPP frequency makes it possible to enhance second-harmonic generation [12] and implement the extremely high light transmittance through a subwavelength hole [13]. The TPPs and devices based on them are designed, as a rule, using a planar metallic film coupled with a PhC. The potentialities of optimizing the optical properties of such structures by means of the variation in the film parameters are exhausted by choosing a material and thickness of this film. New opportunities are offered by metasurfaces and metal-dielectric nanocomposites (NCs), i.e., artificial media structured in a special way, used as film materials [14]. A nanocomposite is a dielectric matrix with metallic particles uniformly distributed over its volume, which is characterized by the resonant effective permittivity. The optical properties of initial materials have no resonant features [15–17]. The position of a resonance in the visible spectral range, as well as the frequency band where the NC behaves like a metal, are determined by the effective permittivity. The latter, in turn, depends on the permittivities of initial materials and concentration, shape, orientation, and size of metallic particles. In particular, the authors of study [18] demonstrated first the possibility of forming a localized mode at the interface between a PhC and a NC representing a transparent matrix with uniformly dispersed silver nanoparticles. The TPP formation at the interface

35 between a PhC bound by a strongly anisotropic NC layer was studied in [19]. The authors of study
 36 [20] proposed a mechanism for forming two TPPs at the interface between a superlattice and an NC
 37 consisting of spherical nanoparticles with a dielectric core and a metal shell. The possibility of forming
 38 a TPP by using porous and gyroidal plasmon materials was discussed in studies [21,22]. It is worth
 39 noting that metal-dielectric NCs can serve as materials with the near-zero effective permittivity. In
 40 recent years, such materials have been in focus of researchers [23] due to the possibility of controlling
 41 the wave front shape [24], amplifying the light transmission through a subwavelength aperture [25],
 42 and enhancing the nonlinear effects [26,27]. As was shown in [28], such materials can be used for
 43 forming the TPPs. The authors established the dissipative principle of the formation of a localized
 44 mode, in which the TPP can form not only at small negative and positive values of the real part of the
 45 permittivity, but also at its zero value. In this work, we show the possibility of designing narrowband
 46 absorbers based on the TPPs localized at the interface between a PhC and an NC with the near-zero
 47 permittivity. Using the temporal coupled-mode theory, we compare two TPP excitation schemes and
 48 predict spectral properties of the localized modes. We demonstrate that the analytical results agree
 49 well with the numerical calculation.

50 2. Model Description

51 We consider a PhC structure, which represents a layered medium bound by a finite NC layer
 52 (Fig. 1). The PhC unit cell is formed from materials a and b with respective layer thicknesses d_a and d_b
 53 and permittivities ε_a and ε_b . The nanocomposite layer with thickness d_{eff} and permittivity ε_{eff} consists
 of metal nanospheres uniformly distributed in a transparent matrix made of optical glass. Then, we

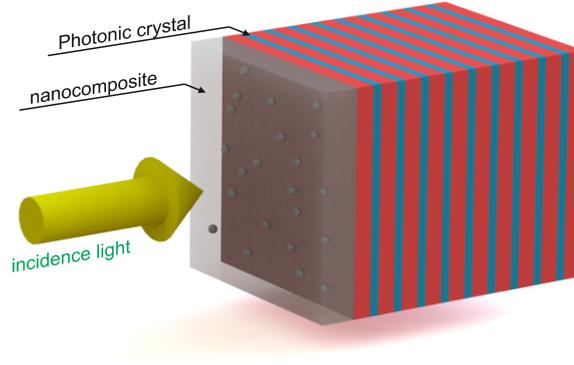


Figure 1. Schematic of a one-dimensional PhC coupled with an NC layer with the near-zero permittivity

54 assume the PhC to be placed in vacuum.

55 Maxwell's equations are reduced to the Helmholtz equation. For a p -polarized wave propagating
 56 in the xz plane of a layered medium of isotropic materials stacked along the z direction, we have
 57

$$\begin{aligned} \left[\frac{d^2}{dz^2} + \left(\frac{\varepsilon\omega^2}{c^2} - k_x^2 \right) \right] E_x(z) &= 0, \\ H_y(z) &= \frac{-i\omega}{c} \frac{\varepsilon dE_x/dz}{c(\varepsilon\omega^2/c^2 - k_x^2)}, \end{aligned} \quad (1)$$

where $k_x = n(\omega/c)$ is the tangential wavenumber along the x axis, c is the speed of light, and ω is the frequency. The effective permittivity of the NC can be described by the Maxwell-Garnet formula

$$\varepsilon_{eff} = \varepsilon_d \left[1 + \frac{f(\varepsilon_m(\omega) - \varepsilon_d)}{\varepsilon_d + (1-f)(\varepsilon_m(\omega) - \varepsilon_d)1/3} \right], \quad (2)$$

where f – is the filling factor, i.e., the volume fraction of nanoparticles in the matrix; ε_d and $\varepsilon_m(\omega)$ – are the permittivities of the matrix and nanoparticle metal, respectively; and ω – is the radiation frequency. We determine the permittivity of the nanoparticle metal using the Drude approximation

$$\varepsilon_m(\omega) = \varepsilon_0 - \frac{\omega_p^2}{\omega^2 + i\omega\gamma}. \quad (3)$$

58 For certainty, we assume alternating PhC layer materials to be silicon dioxide (SiO₂) with a
 59 permittivity of $\varepsilon_a = 2.10$ and zirconium dioxide (ZrO₂) with a permittivity of $\varepsilon_b = 4.16$. The
 60 respective layer thicknesses are $d_a = 74$ nm and $d_b = 50$ nm. The Drude parameters of silver are
 61 $\varepsilon_0 = 5$, $\omega_p = 9$ eV, and $\gamma = 0.02$ eV [29,30]; for glass, we have $\varepsilon_d = 2.56$.

62 As an example, Fig. 2 shows the $Re \varepsilon_{eff}$ and $Im \varepsilon_{eff}$ dependences calculated using formula (2) with
 regard to Eq. (3) at $f = 0.11$. It can be seen that in the range of $[\lambda_1, \lambda_0]$ we have $Re \varepsilon_{eff} < 0$; i.e., the

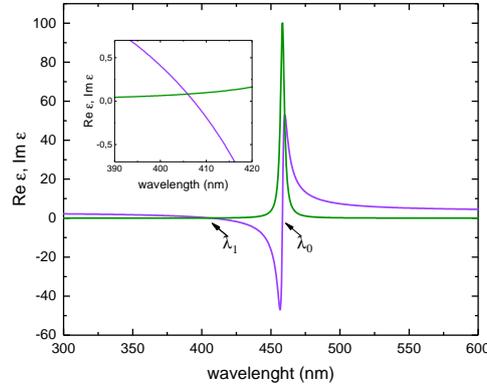


Figure 2. Dependences of the real $Re \varepsilon_{eff}$ (purple line) and imaginary $Im \varepsilon_{eff}$ (green line) parts of the effective permittivity ε_{eff} on the incident light wavelength. The filling factor is $f = 0.11$.

63
 64 NC behaves like a metal. At wavelengths of $\lambda_0 = 458.4$ nm and $\lambda_1 = 407.2$ nm, the real part of the
 65 effective permittivity takes the zero value. In this case, $\lambda_0 = 458.4$ nm is the resonant wavelength, at
 66 which the $Im \varepsilon_{eff}$ value is maximum; therefore, the localized modes cannot form in the vicinity of this
 67 point. In view of the aforesaid, in this study we investigate the TPPs in the vicinity of the point λ_1 .

68 3. Temporal Coupled-Mode Theory

69 The optical properties of the TPP can be described using the temporal coupled-mode theory [31–
 70 34]. This theory is grounded on the fact that any mode (resonance) can be characterized by
 71 eigenfrequency ω_0 and number N of the ports through which the energy passes into this mode
 72 and leaks from it. The energy loss in the channels is described by the relaxation times τ_l , $l = 1 \dots N$ and
 73 the mode is described by the complex amplitude A related to the amplitudes $s_{l\pm}$ of the incoming and
 74 outgoing energy flows. These quantities are described by the ordinary differential equation

$$\frac{dA}{dt} = -i\omega_0 A - \sum_{i=1}^2 \frac{A}{\tau} + \sum_{i=1}^2 \sqrt{\frac{2}{\tau_i}} s_{i+}, \quad (4)$$

The relation between the flow amplitudes is

$$s_{l-} = -s_{l+} + \sqrt{\frac{2}{\tau_l}} A. \quad (5)$$

75 If the resonance is excited through one port with $s_{l+} = -s_{l+}e^{-i\omega t}$, the resonance amplitude can
76 be expressed as

$$A_l(\omega) = \frac{\sqrt{\frac{2}{\tau_l}}}{i(\omega - \omega_0) + \sum_{l=1}^N \frac{1}{\tau_l}} s_{l+}. \quad (6)$$

77 The amplitudes of reflection from port l to port l' form the scattering matrix

$$r_{ll'} = \frac{s_{l'-}}{s_{l+}} = -\hat{\delta}_{ll'} + \frac{\sqrt{\frac{2}{\tau_l}} \sqrt{\frac{2}{\tau_{l'}}}}{i(\omega - \omega_0) + \sum_{l''=1}^N \frac{1}{\tau_{l'}}}, \quad (7)$$

78 where $\hat{\delta}_{ll'}$ is the Kronecker symbol.

79 These reflectances and transmittances correspond to the spectral peaks in the form of Lorentz
80 contours with the full width at half maximum

$$2\gamma = \sum_{l=1}^N \frac{1}{\tau_l}. \quad (8)$$

81 For one channel, we have $l = l'$. In this case, the reflection amplitude can be written as

$$r_l = -1 + \frac{2\gamma_l}{i(\omega_0 - \omega) + \sum \gamma_{l'}}. \quad (9)$$

The zero reflection is only possible under the critical coupling conditions, i.e., at $\omega = \omega_0$:

$$r_l(\omega = \omega_0) = 0. \quad (10)$$

82 This is sufficient to describe the spectral properties of the TPPs using the temporal coupled-mode
83 theory.

84 4. Study of the Spectral Properties of the TPP Using the Temporal Coupled-Mode Theory

85 The TPP formation is contributed by three energy channels, each characterized by the amplitude
86 relaxation rate γ , which is the ratio between the power of energy relaxation to a channel and the
87 energy accumulated in the TPP. We denote the rates of energy relaxation to the NC transmission and
88 absorption channels and PhC transmission channel as γ_{NC} , γ_A , and γ_{PhC} , respectively. Since the
89 energy accumulated in the TPP is the same for determining the rate of relaxation to each channel, the
90 relaxation rates and corresponding energy coefficients of the structure are expressed by the proportion
91 [34]:

$$\gamma_{NC} : \gamma_A : \gamma_{PhC} = T_{NC} : A_{NC} : T_{PhC}. \quad (11)$$

92 We study the conditions of the critical coupling between the TPP and incident field for two
93 excitation schemes.

If one of the mirrors is opaque, we may ignore one of the energy relaxation channels. Then, critical coupling condition (10) for the scheme of excitation through the NC layer (Fig. 3a) can be written in the form

$$\gamma_{NC} = \gamma_A; \gamma_{PhC} = 0 \Leftrightarrow T_{NC} = A_{NC}; T_{PhC} = 0, \quad (12)$$

94 and, for the scheme of excitation through the PhC (Fig. 3b), in the form

$$\gamma_{PhC} = \gamma_A; \gamma_{NC} = 0 \Leftrightarrow T_{PhC} = A_{NC}; T_{NC} = 0. \quad (13)$$

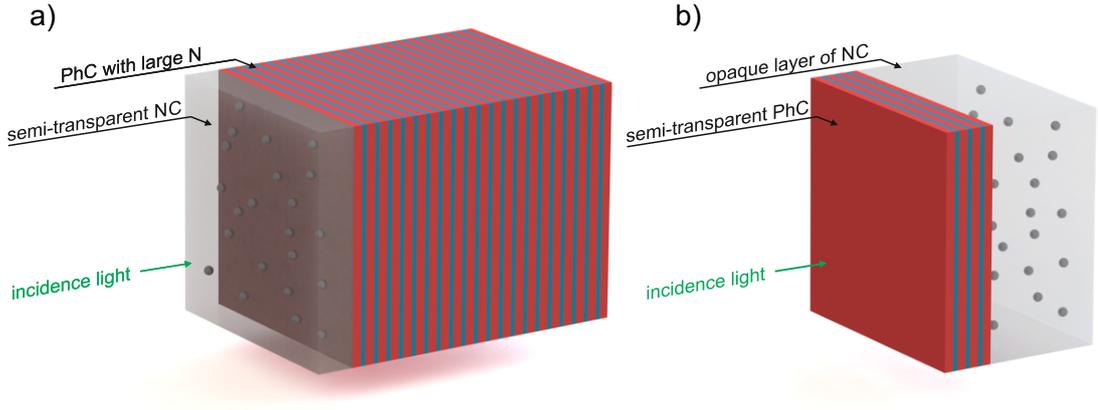


Figure 3. Schematic of the TPP excitation from the side of (a) an NC film and (b) a PhC.

95 Critical coupling equation (12) can be solved graphically. To do that, we should plot together
 96 the dependences of the transmittances and absorptances of the NC film on its thickness and incident
 97 radiation wavelength. The intersection of the surfaces corresponds to the equality of the coefficients
 98 and critical coupling. The energy coefficients for the NC layer can, in turn, be determined using the
 99 Airy formula derived for the metallic layer [35]. For this purpose, we consider a plane-parallel NC
 100 film with the refractive index $n_2 = n'_2 + in''_2$ placed between two dielectric media with the refractive
 101 indices $n_1 = n_{SiO_2}$ and $n_3 = 1$. The transmittances, reflectances, and absorptances of the NC film will
 102 be determined as

$$T_{NC} = \frac{n_3}{n_1} \left| \frac{t_{12} + t_{23}e^{i\beta}}{1 + r_{12}r_{23}e^{2i\beta}} \right|^2, \quad R_{NC} = \left| \frac{r_{12} + r_{23}e^{2i\beta}}{1 + r_{12}r_{23}e^{2i\beta}} \right|^2, \quad (14)$$

$$A_{NC} = 1 - T_{NC} - R_{NC};$$

103 where $\beta = 2\pi n_2 d_{eff} / \lambda_0$ is the phase incoming when the wave passes through the layer, λ_0 is the
 104 wavelength, d_{eff} is the layer thickness, and $t_{12} = 2n_1 / (n_1 + n_2)$, $r_{12} = (n_1 - n_2) / (n_1 + n_2)$, and
 105 $t_{23} = 2n_2 / (n_2 + n_3)$, $r_{23} = (n_2 - n_3) / (n_2 + n_3)$ are the transmission and reflection amplitudes at the
 106 1-2 and 2-3 interfaces, respectively.

107 The NC film spectra calculated using formula (14) are shown in Fig. 4.

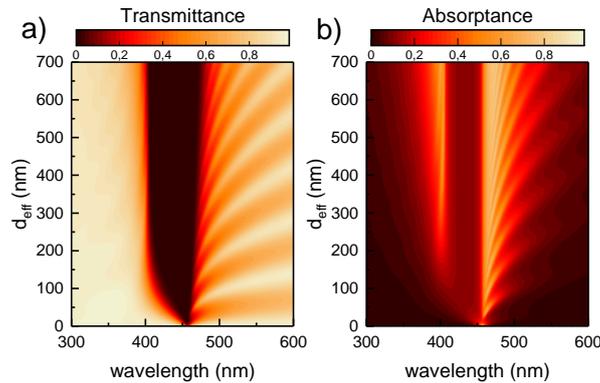


Figure 4. (a) Transmission and (b) absorption spectra of the NC film at different film thicknesses and incident radiation wavelengths. The filling factor is $f = 0.11$.

108 The critical coupling is presented by the line of intersection of these surfaces. According to the
 109 numerical calculation, condition (12) in the wavelength ranges of $(407 < \lambda < 408 \text{ nm})$, where the NC
 110 effective permittivity is near-zero, is satisfied at NC film thicknesses of about 200 nm. Thus, coupling

111 of a PhC with the NC film of such a thickness will lead to the formation of the TPP, at the wavelength
 112 of which the entire radiation incident onto the structure will be absorbed.

113 The graphic solution of the critical coupling conditions for two TPP excitation schemes is
 114 illustrated in Fig. 5.

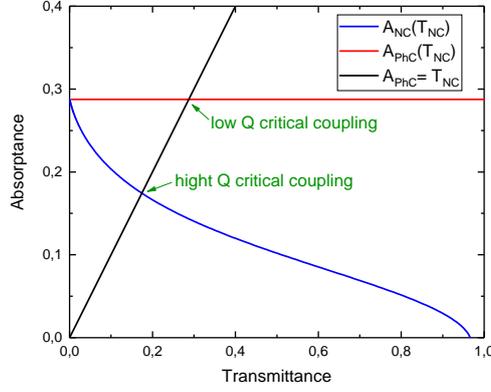


Figure 5. Dependence of absorbance of the NC film on its transmittance at different film thicknesses (blue line) and dependence of the absorbance of an opaque NC film on the PhC transmittance at different numbers of PhC periods (red line). The point of intersection between the black and blue lines corresponds to critical coupling condition (12) and the point of intersection between the black and red lines corresponds to condition (13). The wavelength is $\lambda = 407.1$ nm.

In this case, the dependence of the PhC transmittance on the number of periods N was determined as

$$T_{PhC} = 1 - |r_N|^2, \quad (15)$$

where r_N is the amplitude of reflection from the multilayer structure [36]:

$$r_N = \frac{CU_{N-1}}{AU_{N-1} - U_{N-2}}, \quad (16)$$

$U_N = \sin[(N+1)K\Lambda] / \sin[K\Lambda]$ and $K = 1/\Lambda \arccos[(A+D)/2]$ is the Bloch wavenumber, and N is the number of periods. $A, B, C,$ and D are the elements of the 2×2 transfer matrix, which relates the plane wave amplitudes in layer 1 of the unit cell to the analogous amplitudes for an equivalent layer in the next PhC unit cell.

$$\begin{aligned} A &= e^{ik_{1z}d_a} \left[\cos k_{2z}d_b + \frac{1}{2}i \left(\frac{k_{2z}}{k_{1z}} + \frac{k_{1z}}{k_{2z}} \right) \sin k_{2z}d_b \right], \\ B &= e^{-ik_{1z}d_a} \left[\frac{1}{2}i \left(\frac{k_{2z}}{k_{1z}} - \frac{k_{1z}}{k_{2z}} \right) \sin k_{2z}d_b \right], \\ C &= e^{ik_{1z}d_a} \left[-\frac{1}{2}i \left(\frac{k_{2z}}{k_{1z}} - \frac{k_{1z}}{k_{2z}} \right) \sin k_{2z}d_b \right], \\ D &= e^{-ik_{1z}d_a} \left[\cos k_{2z}d_b - \frac{1}{2}i \left(\frac{k_{2z}}{k_{1z}} + \frac{k_{1z}}{k_{2z}} \right) \sin k_{2z}d_b \right], \end{aligned} \quad (17)$$

115 where $k_{1z} = (\omega/c) \sqrt{\varepsilon_a}$ and $k_{2z} = (\omega/c) \sqrt{\varepsilon_b}$ are the wave vectors of the first and second layers.

116 It can be seen from the figures that in the scheme of TPP excitation through the NC, the critical
 117 coupling conditions are established at lower transmittances and absorbances, i.e., at lower energy
 118 relaxation rates, the sum of which determines the resonance spectral line width. Thus, the scheme
 119 of TPP excitation through the NC is more attractive, since under the critical coupling conditions, the
 120 resonance line and, consequently, absorption band is narrower. This comparison is contrary to the
 121 comparison of IR radiation TPP linewidths experimentally observed in [34]. This can be explained
 122 through the difference between metal dispersion laws for visible and IR radiations.

123 5. Numerical Calculation

124 To numerically compare the two excitation schemes, we calculate the transmittance spectra of the
 125 structures using the transfer matrix method [37]. The results of the calculation are presented in Fig. 6.
 It can be seen in Fig.6a that the critical coupling of the TPP with the incident field upon excitation

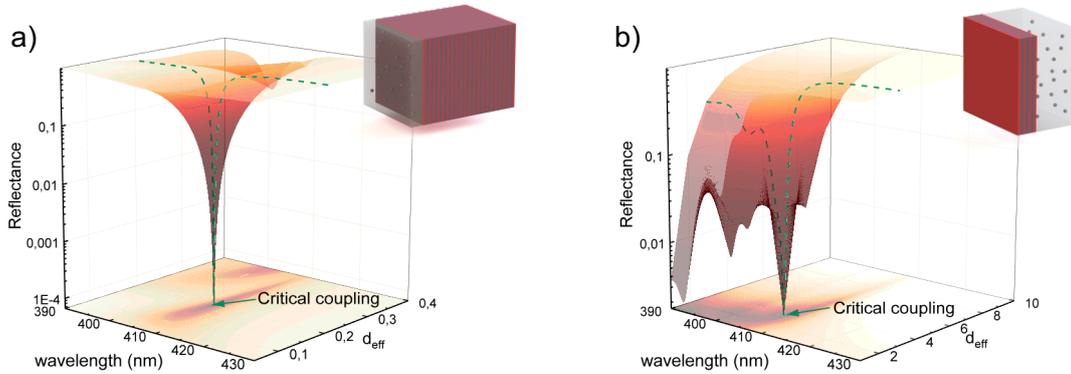


Figure 6. (a) NC-PhC and (b) PhC-NC reflectance spectra of the structure at different NC layer thicknesses d_{eff} and a constant filling factor of $f = 0.11$. The green dashed line shows the reflectance spectra of the structures under critical coupling conditions.

126 through the NC is obtained at an NC layer thickness of $d_{eff} = 201$ nm, which is consistent with the
 127 above-described theory. In addition, the calculation showed that upon TPP excitation from the PhC
 128 side, the critical coupling is obtained at a number of PhC periods of $N = 3$. In this case, the spectral line
 129 for the first excitation scheme appears narrower, which is in good agreement with the data reported
 130 in the previous section. The energy spectra of the structure for the two excitation schemes under the

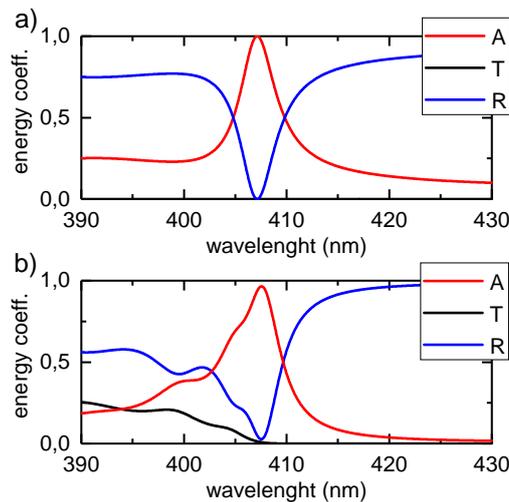


Figure 7. Transmittance (red line), reflectance (blue line), and absorptance (black line) spectra of the structure under the critical coupling conditions upon TPP excitation from (a) the NC and (b) PhC side. The NC layer thickness and number of PhC periods are $d_{eff} = 201$ nm, $N = 25$ and $d_{eff} = 700$ nm, $N = 3$, respectively.

131 critical coupling conditions are shown in Fig. 7.

132 Under the TPP excitation from the NC side, the maximum absorptance is observed at a wavelength
 133 of $\lambda = 407.1$ nm and under the excitation from the PhC side, at a wavelength of $\lambda = 406.7$ nm.
 134

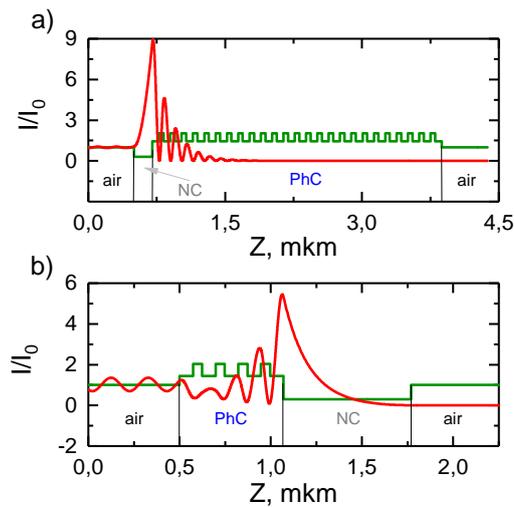


Figure 8. TPP in the region of positive ϵ_{eff} . Spatial distribution of the refractive index of the structure (green line) and local field intensity at the TPP wavelength (red line) for the TPP excitation from (a) the NC and (b) PhC side.

135 The effective permittivity at these wavelengths takes the values $\epsilon_{eff} = 0.0094 + 0.0858i$ and $\epsilon_{eff} =$
 136 $0.0348 + 0.0843i$, respectively.

137 The spatial distributions of the local field intensity at the TPP wavelengths are presented in Fig. 8.

138

139 In both cases, the field is localized at the interface between the media in the region comparable
 140 with the wavelength. The localization value for the scheme of excitation through the NC is higher than
 141 that for the scheme of excitation through the PhC by a factor of 1.5.

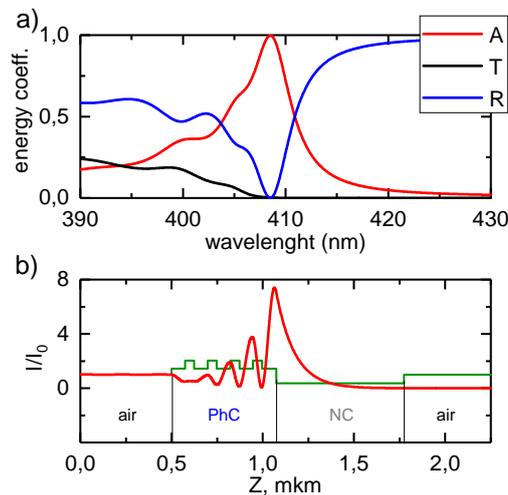


Figure 9. (a) TPP in the metal-like NC region $\epsilon_{eff} < 0$. Transmittance (black line), reflectance (blue line), and absorptance (red line) spectra of the structure under the critical coupling conditions for the scheme of TPP excitation from the PhC side at a thickness of $d_{first} = 78$ nm of the layer adjacent to the PhC and (b) spatial distribution of the local field intensity at the TPP wavelength.

142 It should be noted that under excitation from the PhC side, at the TPP wavelength (Fig. 7b) 2.5%
 143 of the radiation falling onto the structure is not absorbed, but passes through the NC layer. This is

144 related to the TPP formation in the range of small positive epsilon values, where the NC plays the role
145 of a dissipative dielectric [38].

146 The zero transmission can be obtained by changing the TPP wavelength via changing the thickness
147 of the PhC layer adjacent to the NC. As a result, the localized mode wavelength appears in the
148 metal-like NC region and the transmittance decreases (Fig. 9).

149 The local field intensity at the TPP wavelength normalized to the input intensity is shown in in
150 Fig.9b. Here, the field localization is higher than in the case shown in Fig.8a by 20%. This confirms the
151 fact that the critical coupling of the incident field with the TPP is obtained. Comparing with Fig.8b,
152 we can see in Fig.9b that the skin layer of the metal-like NC is thinner and about 10^{-6} of the incident
153 radiation is transmitted at the TPP wavelength.

154 6. Conclusions

155 Thus, we examined the optical properties of TPPs localized at the interface between a PhC and
156 an NC with the near-zero permittivity. We studied two TPP excitation schemes: through a PhC and
157 through an NC layer. Using the temporal coupled-mode theory, we determined the parameters of the
158 structure corresponding to the critical coupling of the incident field with the TPP. It was established
159 that under this condition, the entire radiation falling onto the structure is absorbed by it. It was
160 demonstrated analytically and numerically that the scheme of TPP excitation through the NC is
161 more attractive, since under the critical coupling conditions the resonance line and, consequently, the
162 absorption band is narrower. It was established that to obtain the 100% absorption of the incident
163 radiation upon TPP excitation through the PhC, it is necessary to change the thickness of the first PhC
164 coupled to the NC. This ensures the required phase shift of the wave and TPP localization in the region
165 with the negative epsilon values. The proposed model can be used in designing narrowband absorbers
166 based on Tamm plasmon polaritons localized in resonant PhC structures.

167 **Author Contributions:** R.G.B. performed the calculations, visualized the results and drafted the manuscript. I.V.T.
168 helped with software and methods, S.Y.V. supervised the whole study and finalized the manuscript.

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171 References

- 172 1. Kaliteevski, M.A.; Iorsh, I.; Brand, S.; Abram, R.A.; Chamberlain, J.M.; Kavokin, A.V.; Shelykh, I.A. Tamm
173 plasmon-polaritons: Possible electromagnetic states at the interface of a metal and a dielectric Bragg mirror.
174 *Physical Review B* **2007**, *76*, 165415. doi:10.1103/PhysRevB.76.165415.
- 175 2. Sasin, M.E.; Seisyan, R.P.; Kaliteevski, M.A.; Brand, S.; Abram, R.A.; Chamberlain, J.M.; Egorov, A.Y.;
176 Vasil'ev, A.P.; Mikhlin, V.S.; Kavokin, A.V. Tamm plasmon polaritons: Slow and spatially compact light.
177 *Applied Physics Letters* **2008**, *92*, 251112. doi:10.1063/1.2952486.
- 178 3. Sasin, M.E.; Seisyan, R.P.; Kaliteevski, M.A.; Brand, S.; Abram, R.A.; Chamberlain, J.M.; Iorsh, I.V.; Shelykh,
179 I.A.; Egorov, A.Y.; Vasil'ev, A.P.; Mikhlin, V.S.; Kavokin, A.V. Tamm plasmon-polaritons: First experimental
180 observation. *Superlattices and Microstructures* **2010**, *47*, 44–49. doi:10.1016/j.spmi.2009.09.003.
- 181 4. Svyakhovskiy, S.E.; Bikbaev, R.G.; Myslivets, S.A.; Evlashin, S.A.; Vyunishev, A.M.; Pankin, P.S.;
182 Timofeev, I.V.; Vetrov, S.Y.; Arkhipkin, V.G. Experimental Demonstration of Broadband Optical Tamm
183 States in Photonic Crystal. 2018 International Conference Laser Optics (ICLO), 2018, pp. 309–309.
184 doi:10.1109/LO.2018.8435505.
- 185 5. Zhang, X.L.; Song, J.F.; Li, X.B.; Feng, J.; Sun, H.B. Optical Tamm states enhanced broad-band absorption of
186 organic solar cells. *Applied Physics Letters* **2012**, *101*, 243901. doi:10.1063/1.4770316.
- 187 6. Gong, Y.; Liu, X.; Wang, L.; Lu, H.; Wang, G. Multiple responses of TPP-assisted near-perfect absorption in
188 metal/Fibonacci quasiperiodic photonic crystal. *Optics Express* **2011**, *19*, 9759. doi:10.1364/OE.19.009759.
- 189 7. Symonds, C.; Lheureux, G.; Hugonin, J.P.; Greffet, J.J.; Laverdant, J.; Brucoli, G.; Lemaitre, A.; Senellart, P.;
190 Bellessa, J. Confined Tamm Plasmon Lasers. *Nano Letters* **2013**, *13*, 3179–3184. doi:10.1021/nl401210b.

- 191 8. Auguie, B.; Fuertes, M.C.; Angelomé, P.C.; Abdala, N.L.; Soler Illia, G.J.A.A.; Fainstein, A. Tamm Plasmon
192 Resonance in Mesoporous Multilayers: Toward a Sensing Application. *ACS Photonics* **2014**, *1*, 775–780.
193 doi:10.1021/ph5001549.
- 194 9. Chestnov, I.Y.; Sedov, E.S.; Kutrovskaya, S.V.; Kucherik, A.O.; Arakelian, S.M.; Kavokin, A.V.
195 One-dimensional Tamm plasmons: Spatial confinement, propagation, and polarization properties. *Phys.*
196 *Rev. B* **2017**, *96*, 245309. doi:10.1103/PhysRevB.96.245309.
- 197 10. Yang, Z.y.; Ishii, S.; Yokoyama, T.; Dao, T.D.; Sun, M.g.; Nagao, T.; Chen, K.p. Tamm plasmon selective
198 thermal emitters. *Optics Letters* **2016**, *41*, 4453. doi:10.1364/OL.41.004453.
- 199 11. Gubaydullin, A.R.; Symonds, C.; Bellessa, J.; Ivanov, K.A.; Kolykhalova, E.D.; Sasin, M.E.; Lemaitre, A.;
200 Senellart, P.; Pozina, G.; Kaliteevski, M.A. Enhancement of spontaneous emission in Tamm plasmon
201 structures. *Scientific Reports* **2017**, *7*, 9014. doi:10.1038/s41598-017-09245-7.
- 202 12. Afinogenov, B.I.; Bessonov, V.O.; Fedyanin, A.A. Second-harmonic generation enhancement in the presence
203 of Tamm plasmon-polaritons. *Optics Letters* **2014**, *39*, 6895. doi:10.1364/OL.39.006895.
- 204 13. Treshin, I.V.; Klimov, V.V.; Melentiev, P.N.; Balykin, V.I. Optical Tamm state and extraordinary
205 light transmission through a nanoaperture. *Physical Review A* **2013**, *88*, 023832, [arXiv:1305.4340v1].
206 doi:10.1103/PhysRevA.88.023832.
- 207 14. Takayama, O.; Bogdanov, A.A.; Lavrinenko, A.V. Photonic surface waves on metamaterial interfaces.
208 *Journal of Physics: Condensed Matter* **2017**, *29*, 463001. doi:10.1088/1361-648x/aa8bdd.
- 209 15. Oraevsky, A.N.; Protsenko, I.E. Optical properties of heterogeneous media. *Quantum Electronics* **2001**,
210 *31*, 252–256. doi:10.1070/QE2001v031n03ABEH001927.
- 211 16. Moiseev, S.G.; Ostatochnikov, V.A.; Sementsov, D.I. Defect mode suppression in a photonic
212 crystal structure with a resonance nanocomposite layer. *Quantum Electronics* **2012**, *42*, 557–560.
213 doi:10.1070/QE2012v042n06ABEH014822.
- 214 17. Vetrov, S.Y.; Avdeeva, A.Y.; Bikbaev, R.G.; Timofeev, I. Traveling of light through a 1D photonic
215 crystal containing a defect layer with resonant dispersion. *Optics and Spectroscopy* **2012**, *113*, 517–521.
216 doi:10.1134/S0030400X12110070.
- 217 18. Vetrov, S.Y.; Bikbaev, R.G.; Timofeev, I. Optical Tamm states at the interface between a photonic crystal
218 and a nanocomposite with resonance dispersion. *Journal of Experimental and Theoretical Physics* **2013**,
219 *117*, 988–998. doi:10.1134/S1063776113140185.
- 220 19. Vetrov, S.Y.; Bikbaev, R.G.; Timofeev, I. The optical Tamm states at the edges of a photonic crystal bounded
221 by one or two layers of a strongly anisotropic nanocomposite. *Optics Communications* **2017**, *395*, 275–281.
222 doi:10.1016/j.optcom.2016.08.075.
- 223 20. Vetrov, S.Y.; Pankin, P.S.; Timofeev, I. The optical Tamm states at the interface between a photonic
224 crystal and a nanocomposite containing core-shell particles. *Journal of Optics* **2016**, *18*, 065106.
225 doi:10.1088/2040-8978/18/6/065106.
- 226 21. Bikbaev, R.G.; Vetrov, S.Y.; Timofeev, I. The optical Tamm states at the interface between a photonic crystal
227 and nanoporous silver. *Journal of Optics* **2017**, *19*, 015104. doi:10.1088/2040-8986/19/1/015104.
- 228 22. Bikbaev, R.G.; Vetrov, S.Y.; Timofeev, I. Optical Tamm states at the interface between a photonic crystal and
229 a gyroid layer. *Journal of the Optical Society of America B* **2017**, *34*, 2198. doi:10.1364/JOSAB.34.002198.
- 230 23. Javani, M.H.; Stockman, M.I. Real and Imaginary Properties of Epsilon-Near-Zero Materials. *Physical*
231 *Review Letters* **2016**, *117*, 1–6. doi:10.1103/PhysRevLett.117.107404.
- 232 24. Alù, A.; Silveirinha, M.G.; Salandrino, A.; Engheta, N. Epsilon-near-zero metamaterials and
233 electromagnetic sources: Tailoring the radiation phase pattern. *Physical Review B* **2007**, *75*, 155410,
234 [arXiv:cond-mat/0609220]. doi:10.1103/PhysRevB.75.155410.
- 235 25. Inampudi, S.; Adams, D.C.; Ribaudo, T.; Slocum, D.; Vangala, S.; Goodhue, W.D.; Wasserman, D.; Podolskiy,
236 V.A. Epsilon-Near-zero enhanced light transmission through a subwavelength slit. *Physical Review B -*
237 *Condensed Matter and Materials Physics* **2014**, *89*, 1–10. doi:10.1103/PhysRevB.89.125119.
- 238 26. Ciattoni, A.; Rizza, C.; Marini, A.; Di Falco, A.; Faccio, D.; Scalora, M. Enhanced nonlinear effects
239 in pulse propagation through epsilon-near-zero media. *Laser and Photonics Reviews* **2016**, *10*, 517–525.
240 doi:10.1002/lpor.201500326.
- 241 27. Kaipurath, R.P.M.; Pietrzyk, M.; Caspani, L.; Roger, T.; Clerici, M.; Rizza, C.; Ciattoni, A.; Di Falco, A.;
242 Faccio, D. Optically induced metal-to-dielectric transition in Epsilon-Near-Zero metamaterials. *Nature*
243 *Publishing Group* **2016**, pp. 1–7, [1601.07088]. doi:10.1038/srep27700.

- 244 28. Vetrov, S.Y.; Bikbaev, R.G.; Rudakova, N.V.; Chen, K.p.; Timofeev, I. Optical Tamm states at the interface
245 between a photonic crystal and an epsilon-near-zero nanocomposite. *Journal of Optics* **2017**, *19*, 085103.
246 doi:10.1088/2040-8986/aa75fb.
- 247 29. Yang, H.U.; D'Archangel, J.; Sundheimer, M.L.; Tucker, E.; Boreman, G.D.; Raschke, M.B. Optical dielectric
248 function of silver. *Physical Review B* **2015**, *91*, 235137, [0801.4433]. doi:10.1103/PhysRevB.91.235137.
- 249 30. Johnson, P.; Christy, R.W. Optical constants of the noble metals. *Physical Review B* **1972**, *6*, 4370–4379,
250 [arXiv:1011.1669v3]. doi:10.1103/PhysRevB.6.4370.
- 251 31. Haus, H. *Waves and fields in optoelectronics*; Prentice-Hall: Englewood Cliffs, NJ, 1984.
- 252 32. Joannopoulos, J.D.; Johnson, S.G.; Winn, J.N.; Meade, R.D. *Photonic crystals : molding the flow of light*;
253 Princeton University Press: Princeton, 2008.
- 254 33. Auguié, B.; Bruchhausen, A.; Fainstein, A. Critical coupling to Tamm plasmons. *Journal of Optics* **2015**,
255 *17*, 035003, [arXiv:1411.0608v1]. doi:10.1088/2040-8978/17/3/035003.
- 256 34. Yang, Z.Y.; Ishii, S.; Yokoyama, T.; Dao, T.D.; Sun, M.G.; Pankin, P.S.; Timofeev, I.; Nagao, T.; Chen,
257 K.p. Narrowband Wavelength Selective Thermal Emitters by Confined Tamm Plasmon Polaritons. *ACS*
258 *Photonics* **2017**, *4*, 2212–2219. doi:10.1021/acsp Photonics.7b00408.
- 259 35. Born, M.; Wolf, E. *Principles of optics : electromagnetic theory of propagation, interference and diffraction of light*;
260 Cambridge University Press: Cambridge New York, 1999.
- 261 36. Yariv, A.; Yeh, P. *Optical waves in crystals : propagation and control of laser radiation*; John Wiley and Sons:
262 Hoboken, N.J, 2003.
- 263 37. Yeh, P. Electromagnetic propagation in birefringent layered media. *Journal of the Optical Society of America*
264 **1979**, *69*, 742. doi:10.1364/JOSA.69.000742.
- 265 38. Bikbaev, R.; Vetrov, S.; Timofeev, I. Two Types of Localized States in a Photonic Crystal Bounded by an
266 Epsilon near Zero Nanocomposite. *Photonics* **2018**, *5*, 22. doi:10.3390/photonics5030022.

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