



Cite this: *Nanoscale Horiz.*, 2020, 5, 1088

Received 3rd March 2020,  
Accepted 24th April 2020

DOI: 10.1039/d0nh00139b

[rsc.li/nanoscale-horizons](http://rsc.li/nanoscale-horizons)

# Reconfigurable all-dielectric Fano metasurfaces for strong full-space intensity modulation of visible light†

Sun-Je Kim,<sup>a</sup> Inki Kim,<sup>b</sup> Sungwook Choi,<sup>c</sup> Hyojin Yoon,<sup>d</sup> Changhyun Kim,<sup>a</sup> Yohan Lee,<sup>a</sup> Chulsoo Choi,<sup>a</sup> Junwoo Son,<sup>ib</sup> Yong Wook Lee,<sup>c</sup> Junsuk Rho<sup>ib</sup> and Byoungho Lee<sup>id</sup> <sup>★</sup>

Dynamically reconfigurable nanoscale tuning of visible light properties is one of the ultimate goals both in the academic field of nanophotonics and the optics industry demanding compact and high-resolution display devices. Among various efforts incorporating actively reconfigurable optical materials into metamaterial structures, phase-change materials have been in the spotlight owing to their optical tunability in wide spectral regions including the visible spectrum. However, reconfigurable modulation of visible light intensity has been limited with small modulation depth, reflective schemes, and a lack of profound theoretical background for universal design rules. Here, all-dielectric phase-change Fano metasurface gratings are demonstrated for strong dynamic full-space (reflection and transmission) modulation of visible intensities based on Fano resonances. By judicious periodic couplings between densely arranged meta-atoms containing VO<sub>2</sub>, phase-change induced thermo-optic modulation of full-space intensities is highly enhanced in the visible spectrum. By providing intuitive design rules, we envision that the proposed study would contribute to nanophotonics-enabled optoelectronics technologies for imaging and sensing.

## New concepts

Among various efforts incorporating actively reconfigurable optical materials into metamaterial structures, particularly nanophotonic devices using phase-change materials have been in the spotlight owing to their relatively large optical tunability in wide spectral regions including the visible spectrum. However, in the visible regime, reconfigurable modulation with phase-change materials has been limited with small modulation depth, reflective schemes, and a lack of profound theoretical background for universal design rules. Here, all-dielectric phase-change Fano metasurface gratings are proposed and studied to relieve such conventional drawbacks. We theoretically and experimentally demonstrated strong dynamic full-space (reflection and transmission) modulation of visible intensities based on Fano resonances in all-dielectric metasurface gratings. It is verified that the concept of all-dielectric Fano resonance is highly effective for enhancing phase-change-induced thermo-optic modulation of full-space intensities in the visible spectrum. By providing both intuitive design rules and high performances, we envision that the proposed study would contribute to nanophotonics-enabled optoelectronics technologies for practical imaging and sensing applications.

## 1. Introduction

By the virtue of simple nanofabrication processes and unprecedented optical functionalities within an ultracompact volume, metasurfaces have facilitated the development of a wide range

of flat optical applications including displays and sensors.<sup>1–5</sup> Their geometric structures are designed in terms of individual subwavelength antennas and their periodic coupling to exhibit the ability of resonant trapping and re-scattering of an incident beam with a desired vectorial wavefront. Ultrathin and functional optical-elements have been demonstrated and studied to effectively replace bulky ray optical elements. By modulating the phase, polarization, and amplitude of reflected or transmitted beams, it has been possible to focus,<sup>6</sup> absorb,<sup>7</sup> filter,<sup>8</sup> and steer<sup>9</sup> light with desired properties. These exciting properties of passive functional metasurfaces with fixed optical functions have prompted research on dynamically tunable metasurfaces providing new opportunities, most notably chip-scale beam steering devices and ultrathin modulators. Actively tunable material platforms including graphene,<sup>10–12</sup> semiconductors,<sup>13–15</sup> transparent conducting oxides,<sup>16–18</sup> liquid crystals,<sup>19</sup> phase-change materials (PCM),<sup>20–31</sup> and mechanical actuators<sup>32–34</sup> have been extensively studied in the last decade. However, due to the extremely short

<sup>a</sup> Inter-University Semiconductor Research Center and School of Electrical and Computer Engineering, Seoul National University, Gwanakro 1, Gwanak-Gu, Seoul 08826, Republic of Korea. E-mail: [byoungho@snu.ac.kr](mailto:byoungho@snu.ac.kr)

<sup>b</sup> Department of Mechanical Engineering, Pohang University of Science and Technology (POSTECH), Pohang 37673, Republic of Korea

<sup>c</sup> Interdisciplinary Program of Biomedical Mechanical & Electrical Engineering and School of Electrical Engineering, Pukyong National University, Yongso-ro 45, Nam-Gu, Busan 48513, Republic of Korea

<sup>d</sup> Department of Materials Science and Engineering, Pohang University of Science and Technology (POSTECH), Pohang 37673, Republic of Korea

<sup>e</sup> Department of Chemical Engineering, Pohang University of Science and Technology (POSTECH), Pohang 37673, Republic of Korea

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0nh00139b



interaction length for light-matter interactions in resonant metasurfaces, all of these platforms lack good performances of practical value. Thus, it is hard to achieve large tunability, high efficiency, and modulation capability in multiple optical properties using a single resonant metadvice. In particular, modulation of visible light waves in free space has been hardly achieved despite its substantial practical value in miniaturized display and sensing technologies with extremely high resolution. Rather than the other materials and platforms mentioned above, PCMs such as vanadium dioxide ( $\text{VO}_2$ ) and chalcogenide compounds show large thermo-optic tunability of the dielectric functions in the visible and near-infrared regime.<sup>20–31</sup> PCM-based metadvice provide a large design degree of freedom in the optical regime by virtue of their relatively large tunability of the dielectric function compared to other material platforms. Several experimental studies have reported modulation of optical intensities using PCMs in the optical range. Compared to the near and mid infrared range, nanophotonic designs of visible range metadvice based on PCMs are very challenging since they show relatively small tunability and large absorption coefficients with low quality factors at visible wavelengths. The empirical demonstration of large intensity modulation based on a strong physical background and intuition has been elusive. Moreover, despite the clear significance and necessity of extremely miniaturized and integrated flat optics systems, dynamic and effective full-space modulation of both reflection and transmission beams has never been studied thoroughly except for several passive metasurfaces.<sup>35,36</sup>

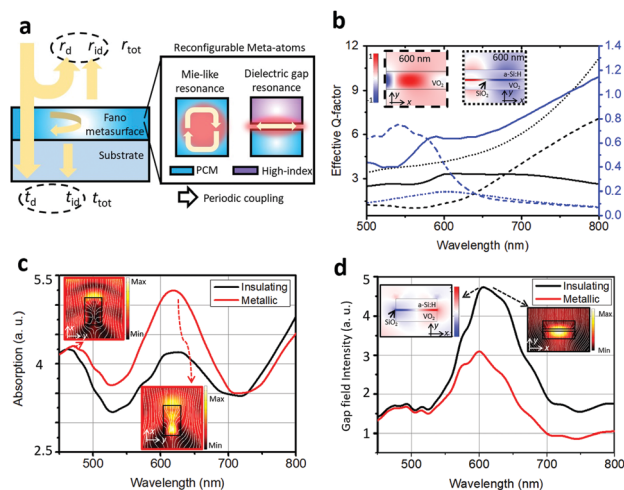
In this paper, we propose a holistic study on simple and effective Fano-resonance methods to tune full-space visible wave intensities using all-dielectric phase-change metasurface gratings based on  $\text{VO}_2$  phase-change. *Via* theoretical and experimental studies, we show that judicious combinations of Fano resonance phenomena in metasurface gratings and the insulator-to-metal transition (IMT) of a  $\text{VO}_2$  film exhibit good modulation performances in simple structures in the visible range. Here, a framework of optical Fano resonance, one of the general phenomena in nanophotonic systems, is introduced for systematic and intuitive understanding.<sup>37–44</sup> For rigorous theoretical design and analysis, zeroth and second-order effective medium theory (EMT), temporal coupled mode theory (TCMT) and Rayleigh-Wood's anomaly (RWA) are introduced in three different types of metasurface devices. The rest of the paper is as follows. Firstly, the thermo-optic properties of a  $\text{VO}_2$  film,  $\text{VO}_2$  nano-waveguide modes, and resonators are investigated as a fundamental step of the study. Then, nanostructured  $\text{VO}_2$  metafilms and diffractive silicon- $\text{VO}_2$  hybrid metagratings are demonstrated with different dynamic modulation characteristics. Last, concluding remarks are made with future perspectives.

## 2. Results and discussion

### 2.1 Thermo-optic properties of $\text{VO}_2$ waveguides and resonators

Ahead of reconfigurable metadvice design, we characterized a tunable  $\text{VO}_2$  thin film made by conventional pulsed laser deposition

on a sapphire substrate<sup>45,46</sup> (see part 1 of the ESI†). The dielectric function implies that  $\text{VO}_2$  is a highly absorptive dielectric in the visible range and it shows a decrease and increase of its refractive index in heating and cooling processes. Based on such tunable  $\text{VO}_2$  properties and low loss materials<sup>47</sup> (see part 2 of the ESI† for the optical properties of a-Si:H), the design strategy is focused on incorporating this thin film into all-dielectric Fano metasurfaces for simultaneous effective control of reflection, absorption, and transmission as shown in Fig. 1(a). In general, an optical Fano resonance with an asymmetric spectral lineshape is constructed by coherent interference of bright continuum scattering and resonant dark mode scattering.<sup>37,39</sup> Thus, individual  $\text{VO}_2$ -containing meta-atoms and their periodic coupling effects are considered rigorously to design tunable and strong Fano interference. As depicted in the inset figures of Fig. 1(a), nano-resonators with Mie resonances<sup>48</sup> and gap photonic mode resonances (GPMR),<sup>49</sup> which confine light mostly in the  $\text{VO}_2$  and lossless low-index  $\text{SiO}_2$  regions, respectively, are considered as unit cell meta-atoms. The first one inherits the conventional advantages of



**Fig. 1** (a) Concepts of full-space modulation by a reconfigurable all-dielectric Fano metasurface designed using efficient meta-atoms and their periodic coupling.  $r_d$ ,  $r_{id}$ ,  $t_d$ , and  $t_{id}$  correspond to direct reflection, indirect reflection, direct transmission, and indirect transmission coefficients, respectively. (b) Comparison of the calculated waveguide effect in terms of the effective quality factor in the insulating phase (left y-axis, black lines) and effective mode index change (right y-axis, blue lines). Solid, dashed, and dotted lines denote a homogeneous  $\text{VO}_2$  medium, 100 nm-thick  $\text{VO}_2$  slab waveguide, and hybrid a-Si:H-SiO<sub>2</sub>- $\text{VO}_2$  waveguide, respectively. The inset field profiles show the normalized  $E_y$  field of the guided TM-polarized modes in the  $\text{VO}_2$  slab waveguide and the hybrid waveguide in the insulating phase (wavelength: 600 nm). (c)  $\text{VO}_2$  phase-change-induced absorption modulation of the  $\text{VO}_2$  ridge resonator (width = 100 nm, thickness = 320 nm). The sub-figures denote the normalized magnetic field intensity at the resonance peaks excited by TM-polarized normal incidence from the top. The white stream lines in the sub-figures present the power flow of incident TM illumination around the resonator. (d)  $\text{VO}_2$  phase-change-induced absorption modulation of the hybrid silicon-silicon dioxide- $\text{VO}_2$  resonator. The resonator width, a-Si:H thickness, SiO<sub>2</sub> thickness, and  $\text{VO}_2$  thickness are 200 nm, 50 nm, 10 nm, and 50 nm, respectively. The subplots show the normalized real part of the  $E_y$  field (left) and magnetic field intensity (right) at the insulating phase resonance (wavelength: 600 nm) under TM illumination.

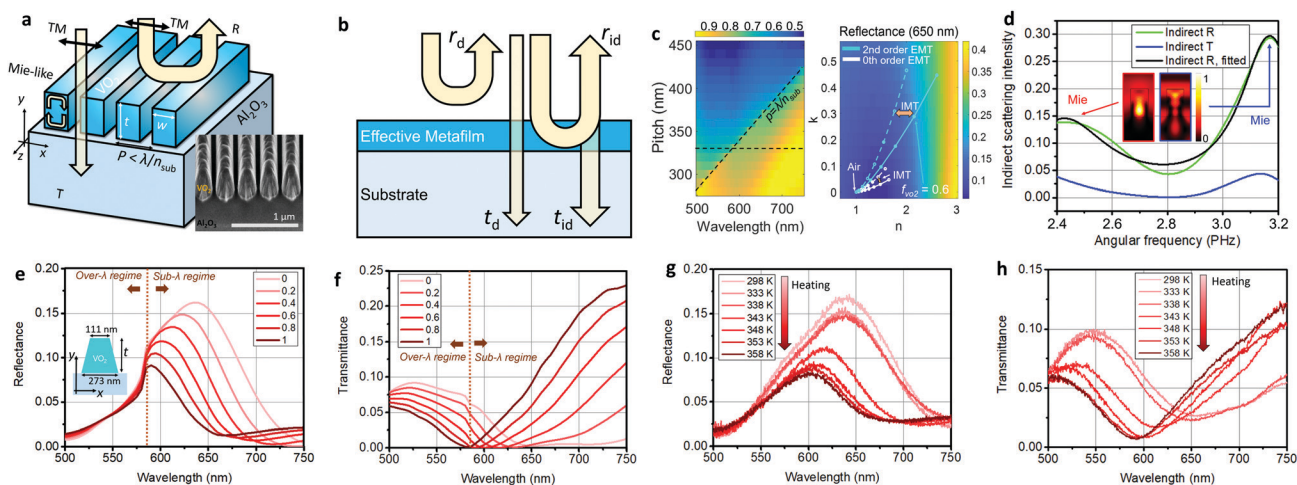


strong magnetic Mie resonances owing to the relatively large refractive index of VO<sub>2</sub> so that light-VO<sub>2</sub> interactions are enhanced easily. On the other hand, a novel GPMR meta-atom shows more advantages with respect to signal efficiency and quality factor as the field penetration into an active VO<sub>2</sub> region is lowered compared to the Mie type meta-atom. The GPMR meta-atom shows interesting field-confining features resembling gap plasmon resonances<sup>50</sup> which exhibit plasmon-assisted ultra-compact capacitive field confinement. However, as a-Si:H and SiO<sub>2</sub> show low absorption loss in the visible regime, GPMR is more advantageous for modulating both reflection and transmission beams rather than gap plasmon resonances. Fig. 1(b) shows an analytical comparison of the effective quality factors and tunability of the refractive index in a homogeneous VO<sub>2</sub> medium, 100 nm-thick VO<sub>2</sub> waveguide, and hybrid a-Si:H (50 nm-thick)-SiO<sub>2</sub> (10 nm-thick)-VO<sub>2</sub> (50 nm-thick) waveguide at room temperature. Numerical full-field simulations are conducted using the RF solver of COMSOL Multiphysics 5.3 and custom-built rigorous coupled wave analysis. The solid, dashed, and short dashed lines refer to the homogeneous VO<sub>2</sub>, VO<sub>2</sub> waveguide, and hybrid a-Si:H-SiO<sub>2</sub>-VO<sub>2</sub> waveguide, respectively. It is clearly shown that the intrinsic material quality factor,  $Q = |\varepsilon_1/\varepsilon_2|$ , of about 3 is too low to make a strong resonance in the bare film. Here,  $\varepsilon_1$  and  $\varepsilon_2$  are the real and imaginary parts of the dielectric function, respectively. On the other hand, the nanostructured waveguides show improved effective quality factor ( $Q_{\text{eff}} = |\varepsilon_{1,\text{eff}}/\varepsilon_{2,\text{eff}}|$ ) according to their

geometry and wavelength, as described in Fig. 1(b). The effective guided mode index,  $n_{\text{eff}}$ , is the square root of  $\varepsilon_{1,\text{eff}} + i\varepsilon_{2,\text{eff}}$ . By sacrificing a bit of effective mode index tunability, it is expected that a more efficient and narrower resonance design could be achieved based on the waveguide study. In Fig. 1(c) and (d), Mie-like resonances and GPMR are numerically investigated with good tunability and enhanced resonance strength compared to the bare film. The two sub-figures in Fig. 1(c) show two strong absorption resonances in a 2-dimensional VO<sub>2</sub> nanobeam resonator. The sub-figures in Fig. 1(d) describe the electromagnetic field properties at the GPMR with enhanced  $Q$ -factor compared to the Mie-like one (Fig. 1(c)). Based on the two distinct resonance properties, tunable Fano metasurface gratings are demonstrated with judicious periodic arrangements of the reconfigurable meta-atoms for modulation of reflection and transmission. A 1-dimensional grating structure is advantageous to study fundamental physical principles of reconfigurable Fano resonances and to apply electrodes for electrical heating without harm to designed optical functions,<sup>14,16–18</sup> rather than 2-dimensionally patterned metasurfaces.

## 2.2 Fano-resonant modulation of a VO<sub>2</sub> metafilm

We first propose a nanostructured metafilm modulator made of periodic subwavelength VO<sub>2</sub> nanobeams. The metafilm strategy has been widely studied for various nanoscale optoelectronics and photonics applications<sup>51–54</sup> in terms of Mie resonances and



**Fig. 2** (a) Schematic diagram of the nanostructured VO<sub>2</sub> metafilm ( $t = 320$  nm and  $w = 180$  nm). The subplot shows a scanning electron micrograph of the fabricated device (b) Schematic diagram of the two-port Fano-resonance mechanism in the metafilm assisted by effective medium theory.  $r_d$ ,  $r_{id}$ ,  $t_d$ , and  $t_{id}$  correspond to the direct reflection, indirect reflection, direct transmission, and indirect transmission coefficients, respectively. (c) Design of the dynamically tunable metafilm. The left 2-dimensional plot shows the absorptance spectra according to varying pitch ( $p$ ) of the unit cell. The oblique and horizontal black dashed lines refer to the diffraction phase-matching condition and  $p = 330$  nm condition, respectively. The right figure shows the reflectance of the approximated metafilm (thickness = 320 nm) effective medium depicted in (b) according to varying refractive index and extinction coefficient at a wavelength of 650 nm. The white and sky blue coloured lines denote the 1st and 2nd order effective medium theory for TM polarization as the filling factor of VO<sub>2</sub> ( $f_{\text{VO}_2}$ ) varies from 0 to 0.7. The solid and dashed lines exhibit the location of the effective index coordinate in the insulating and metallic phases of VO<sub>2</sub>, respectively. (d) Indirect scattering intensity of transmission and reflection. The bright green, blue, and black legends refer to indirect reflectance, indirect transmittance, and doubly resonant Lorentzian fitting results of indirect reflectance. The sub-figures show the two indirect resonances via normalized magnetic field intensity profiles. The full-field simulation results of gradually tunable (e) reflectance and (f) transmittance spectra according to the filling factor (legends) of the metallic phase VO<sub>2</sub>. The simulations are conducted for the tapered metafilm (inset figure of (e)). Measurement results of gradually tunable (g) reflectance and (h) transmittance spectra in the heating process. The legends in (e) and (f) denote the filling factors of metallic VO<sub>2</sub>, and thus the 0 and 1 values correspond to the insulating and metallic phases, respectively. The legends in (g) and (h) denote device temperatures in the heating process.



their periodic coupling effect owing to its simple but extraordinary properties of light trapping and modulation with large design extensibility. Since VO<sub>2</sub> shows significantly tunable dielectric properties in the visible range, densely arranged subwavelength VO<sub>2</sub> nanobeams can be of great advantage to generate strongly reconfigurable Fano resonances with large absorption. As depicted in Fig. 2(a), we propose a Fano-resonant phase-change VO<sub>2</sub> metafilm excited by normal illumination with transverse magnetic (TM) polarization for a dielectric magnetic-like Mie resonance. We fix the thickness ( $t_{\text{VO}_2}$ ) of the VO<sub>2</sub> at 320 nm and the width of each VO<sub>2</sub> nanobeam ( $w$ ) at 180 nm considering fabrication stability of pulsed laser deposition (VO<sub>2</sub> deposition) and focused ion beam milling (nano-patterning). The inset figure in Fig. 2(a) shows a well-defined fabricated metafilm cross-section image captured by scanning electron microscopy assisted by focused ion beam milling (see the detailed fabrication process in part 3 of the ESI†). The working principle of the resonant metafilm device can be described as an effective tunable metafilm with direct and indirect resonant scattering for two radiation channels (Fig. 2(b)). The effective metafilm is constructed from a composite of air and VO<sub>2</sub> nanobeams. The design and analysis of the device are conducted based on the 0th and 2nd order EMTs<sup>55,56</sup> and TCMT formalism<sup>38–42</sup> as follows. As shown in the left 2-dimensional plot of Fig. 2(c), when  $t$  and  $w$  are fixed, variation of the unit cell pitch ( $p$ ) critically affects the optical resonance spectra as the filling factor ( $f_{\text{VO}_2} = w/p$ ) of VO<sub>2</sub> in the metafilm layer and its effective refractive index ( $\epsilon_{\text{TM}}$  and  $\epsilon_{\text{TE}}$ ) change. Such effective medium analysis of the metafilm can be further studied *via* the 0th and 2nd order effective medium theories. According to the 0th order EMT of such a 2-dimensional periodic nanobeam structure,<sup>55</sup> the effective dielectric functions for TE and TM polarizations are described as follows.

$$\epsilon_{\text{TE},0} = f_{\text{VO}_2} \epsilon_{\text{VO}_2} + (1 - f_{\text{VO}_2}) \epsilon_{\text{air}}, \quad (1)$$

$$\epsilon_{\text{TM},0} = \left( \frac{f_{\text{VO}_2}}{\epsilon_{\text{VO}_2}} + \frac{(1 - f_{\text{VO}_2})}{\epsilon_{\text{air}}} \right)^{-1}. \quad (2)$$

However, it is well known that the 2nd order EMT is much more accurate than the 0th order one when the pitch of the unit cell is not much smaller than the wavelength, as is the case with photonic crystals and general metasurfaces exhibiting Fano resonances.<sup>51–56</sup>

$$\epsilon_{\text{TM},2} =$$

$$\epsilon_{\text{TM},0} \left\{ 1 + \frac{\pi^2}{3} \left( \frac{p}{\lambda} \right)^2 f_{\text{VO}_2}^2 (1 - f_{\text{VO}_2})^2 \frac{(\epsilon_{\text{VO}_2} - \epsilon_{\text{air}})^2 \epsilon_{\text{TE},0}}{\epsilon_0} \left( \frac{\epsilon_{\text{TM},0}}{\epsilon_{\text{air}} \epsilon_{\text{VO}_2}} \right)^2 \right\}. \quad (3)$$

As shown in eqn (3), the second order EMT for TM polarization is determined by adding an additional 2nd order compensation term proportional to  $(p/\lambda)^2$  based on the 0th order formulas in eqn (1) and (2). We analytically calculated the reflectance resonance of the effective metafilm at a wavelength of 650 nm using Airy's formula for various effective complex refractive indices of the metafilm. The two distinct 0th and 2nd order EMT calculation curves in the 2-dimensional reflectance map

show that the 2nd order curves accurately account for the reflectance at the wavelength of 650 nm both in the insulating and metallic phases of VO<sub>2</sub> in comparison to the full-field simulation results. While the 2nd order EMT is useful to estimate effective metafilm characteristic more accurately, the 0th order EMT is highly advantageous to explain the Fano resonance mechanism. The reflection and transmission beams are generated by interference between the direct background scattering (bright mode) and indirect resonant scattering (dark mode) of the metafilm.<sup>37–44</sup> Since the 0th order EMT is based on the assumption of the deep subwavelength limit and it cannot explain subwavelength Mie or plasmonic resonances in metamaterials,  $r_d$  and  $t_d$  are the reflection and transmission coefficients of the thin metafilm with dielectric functions of the 0th order EMT. On the other hand,  $r_{id}$  and  $t_{id}$  originate from local subwavelength optical resonances with Lorentzian characteristics. Thus, using temporal coupled mode theory with the 0th order EMT, two port radiation (reflection and transmission) of a single Fano resonance mode can be modelled analytically as follows (part 4 of the ESI†).<sup>38–42</sup>

$$r_{\text{tot}} = r_d + r_{id} = r_d + \frac{ae^{i\phi}}{i(\omega - \omega_0) + \gamma_{\text{tot}}}, \quad (4)$$

$$t_{\text{tot}} = t_d + t_{id} = t_d + \frac{be^{i\phi}}{i(\omega - \omega_0) + \gamma_{\text{tot}}}, \quad (5)$$

$$|r_{id}|^2 = \frac{a^2}{(\omega - \omega_0)^2 + \gamma_{\text{tot}}^2}, |t_{id}|^2 = \frac{b^2}{(\omega - \omega_0)^2 + \gamma_{\text{tot}}^2}. \quad (6)$$

$r_{\text{tot}}$ ,  $t_{\text{tot}}$ ,  $r_d$ ,  $r_{id}$ ,  $t_d$ , and  $t_{id}$  in the above equations are the complex coefficients of total reflection, total transmission, direct reflection, indirect reflection, direct transmission, and indirect transmission, respectively. It is known that direct scatterings  $r_d$  and  $t_d$  are analytically calculated as Fabry-Pérot scatterings of reflection and transmission of a dielectric film with a dielectric function of the zeroth order EMT in eqn (2). Then, the  $r_{id}$  and  $t_{id}$  indirect resonant pathways account for Lorentzian type optical resonance in the system. Thus, we conducted numerical fitting of indirect scattering intensities  $|r_{id}|^2$  and  $|t_{id}|^2$  by determining  $a$ ,  $b$ ,  $\omega_0$ , and  $\gamma_{\text{tot}}$  (part 5 of the ESI†). However, the formulation of TCMT does not perfectly account for the low  $Q$ -factor resonance and multi-resonant case.<sup>41</sup> As depicted in Fig. 2(d), dual indirect resonances are observed in the subwavelength regime. The inset figures of the magnetic field intensity profiles show two resonances with large field enhancement inside the VO<sub>2</sub> nanobeams. We conducted numerical fitting in the insulating phase based on an approximated single resonance formulation near the resonances. By approximately fitting two different Lorentzian resonances independently and adding them, the full spectrum of the reflection is fitted. It is also worth noting that the  $|t_{id}|^2$  spectrum is very similar to  $|r_{id}|^2$  in terms of the resonance location, which matches well with the TCMT formulation (see part 5 of the ESI† for detailed fitting results).

The simulation results considering the fabricated device with tapering walls in Fig. 2(e) and (f) exhibit gradually tunable reflection and transmission *via* the phase change of VO<sub>2</sub> similar to the non-tapered ideal design (part 6 of the ESI†). The gradual



phase change is analytically approximated as the Maxwell-Garnett EMT for intermediate phases between the insulating and metallic phases (part 7 of the ESI†). The measurement results in the heating process in Fig. 2(g) and (h) also match well with the simulation results (see part 8 of the ESI† for reconfigurable and repetitive thermo-optic measurement results). It is remarkable that a large modulation depth reaching 0.8 in a broad bandwidth is achieved both in the visible reflection and transmission intensities (part 8 of the ESI†). Here, the modulation depth ( $\eta$ ) is defined as  $\eta = |I_i - I_m|/\max(I_i, I_m)$ , which is a positive value between 0 and 1.  $I_i$  and  $I_m$  denote the signal intensities in the insulating and metallic phases, respectively.

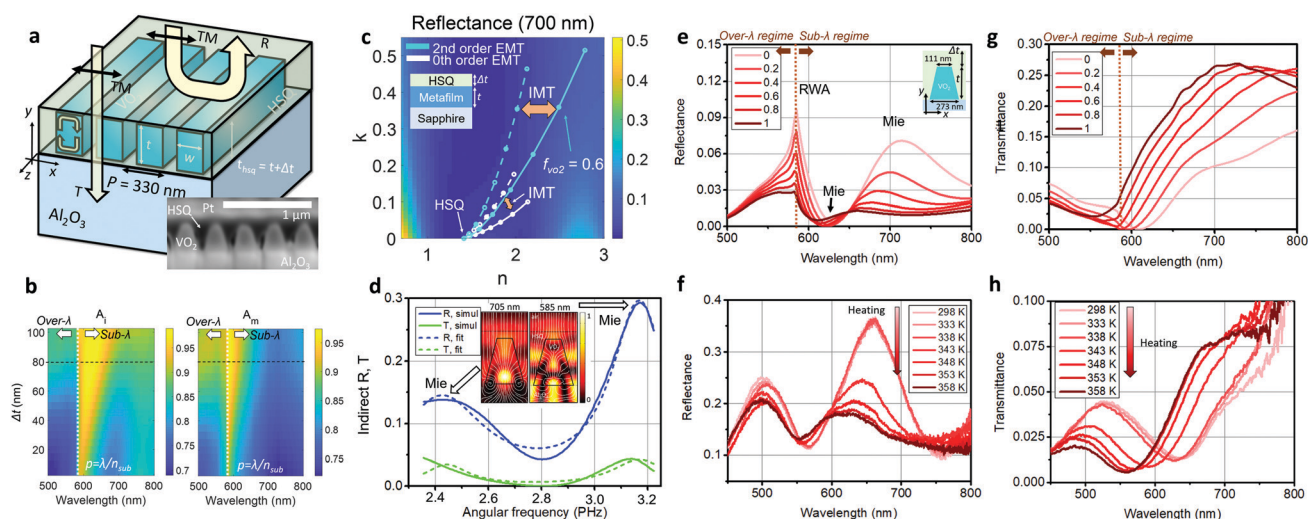
### 2.3 Fano-resonant modulation of a VO<sub>2</sub> metafilm with a transparent dielectric coating

In the previous section, it is verified that nano-structuring of a VO<sub>2</sub> thick film as a nanobeam type metafilm is simple and effective to modulate full-space visible intensities. In this section, we propose a simple additional method of a transparent optical coating on the metafilm to improve the *Q*-factor (spectral bandwidth) and to make use of a different Fano-resonance mechanism, RWA, in addition to coupled Mie resonance. By spin-coating a transparent optical film (hydrogen silsesquioxane, HSQ, with a refractive index of 1.41) on the VO<sub>2</sub> metafilm device, an additional sharp Fano resonance is generated and the tunability of the coupled Mie resonance can be enhanced. Fig. 3(a) shows a

schematic illustration of the transparent dielectric-coated metafilm device, and the inset figure shows the fabricated device image captured by a scanning electron micrograph (see the detailed fabrication process in part 3 of the ESI†). A full-field simulation is conducted to investigate the effect of the HSQ spin coating thickness on the absorption resonance spectra (Fig. 3(b)). Here,  $\Delta t$  is the difference between the HSQ thickness ( $t_{\text{hsq}}$ ) and VO<sub>2</sub> thickness ( $t = 320$  nm). As shown in the plots of Fig. 3(b), two absorption resonances are generated. One is the strong near-unity absorption resonance with coupled Mie scatterings excited near the wavelength of the RWA condition of phase-matching as described in eqn (7).<sup>57–61</sup>

$$\lambda_{\text{RWA}} = \frac{pn_{\text{sub}}(-\sin\theta_i \pm 1)}{m}, \quad m = \pm 1, 2, 3, \dots \quad (7)$$

Here,  $n_{\text{sub}}$  and  $\theta_i$  are the refractive index of the substrate (sapphire) and the incident illumination angle. In addition another Mie resonance is generated with relatively weak resonance strength in the red–near infrared wavelength region. In the case of the weaker one, EMT is an effective way to analyse the resonance (Fig. 3(c)). The 2nd order EMT accurately accounts for sub-wavelength range resonance switching of reflectance compared to the 0th order EMT. The inset figure of Fig. 3(c) of the HSQ coated metafilm describes the reflectance calculation scheme using EMT. The 320 nm-thick metafilm layer made of VO<sub>2</sub> nano-beams and HSQ nanobeams is capped between the 80 nm-thick



**Fig. 3** (a) Schematic diagram of the transparent dielectric-covered VO<sub>2</sub> metafilm ( $p = 330$  nm,  $t = 320$  nm, and  $w = 180$  nm). The subplot shows a cross-sectional scanning electron micrograph of the fabricated device. (b) Design of the dynamically tunable metafilm. The 2-dimensional plots show the absorbance spectra according to varying pitch ( $p$ ) of the unit cell in the insulating (left) and metallic (right) phases, respectively. The horizontal black dashed and vertical white dashed lines refer to the selected HSQ deposition condition ( $t_{\text{hsq}} = t + 80$  nm) and diffraction phase-matching condition, respectively. (c) Reflectance map of the approximated metafilm (thickness = 320 nm) effective medium according to varying refractive index and extinction coefficient at a wavelength of 700 nm. The white and sky blue coloured lines denote the 1st and 2nd order effective medium theory for TM polarization as the filling factor of VO<sub>2</sub> ( $f_{\text{VO}_2}$ ) varies from 0 to 0.7. The solid and dashed lines exhibit the location of the effective index coordinate in the insulating and metallic phases of VO<sub>2</sub>, respectively. (d) Indirect scattering intensity of transmission and reflection. The bright green and blue legends refer to indirect transmittance and indirect reflectance, respectively. The doubly resonant Lorentzian fitting results of indirect scatterings correspond to the dashed lines. The subplots of (d) show the resonant magnetic field intensity profiles in the insulating phase at a wavelength of 585 nm (left) and 705 nm (right) normalized equally. Phase-change induced gradually tunable reflectance spectra from (e) simulations and (f) measurements. The inset figure of (e) describes the simulation unit cell considering tapering walls. Phase-change induced gradually tunable transmittance spectra from (g) simulations and (h) measurements. The experimental results of (f) reflectance and (h) transmittance spectra are measured in the heating process. The legends in (e) and (g) denote the filling factors of metallic VO<sub>2</sub>, and thus the 0 and 1 values correspond to the insulating and metallic phases, respectively. The legends in (f) and (h) denote device temperatures in the heating process.

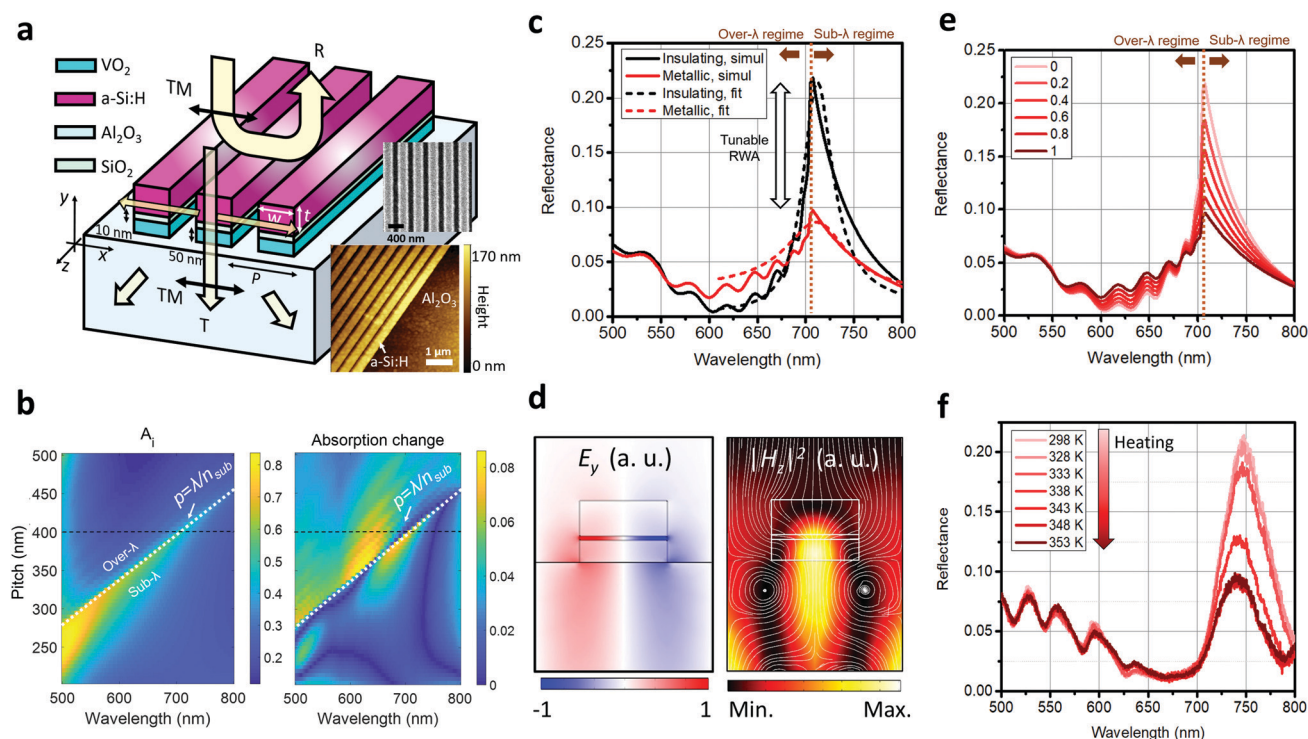
HSQ coating and half-infinite sapphire substrate, and the effective dielectric function of the metafilm layer is described by EMT.

As discussed in the previous section, the 0th order EMT explains direct scatterings while indirect scatterings can be modelled *via* Lorentzian resonance. The plots in Fig. 3(d) show dual resonances of indirect reflection and transmission Lorentzian resonance (see detailed fitting results in part 9 of the ESI†). The inset figures of the magnetic field intensity show that two Lorentzian resonances are excited as coupled Mie-like resonances with large field enhancement and different field profiles. The resonance at higher frequencies shows higher coupling between neighbouring meta-atoms, while the other one shows higher confinement only in the VO<sub>2</sub> region rather than the HSQ region. This abrupt evanescent coupling induced RWA resonance occurs since the metafilm layer is composed of HSQ (refractive index of about 1.41) and VO<sub>2</sub> so that the index contrast is lowered and the nearest-neighbour coupling effect of meta-atoms is increased compared to the air-VO<sub>2</sub> composite metafilm.<sup>57–61</sup> As the indirect absorption resonance at higher frequency is well overlapped with the RWA wavelength in the spectrum, a strong RWA resonance is formed. Such two different resonances contribute to large tunability in reflectance and transmittance (Fig. 3(e)–(h)). Fig. 3(e) and (f) show the reflectance tuning results of numerical calculations and measurements, respectively. Fig. 3(g) and (h) present

the transmittance results of simulations and measurements, respectively. While the reflectance and transmittance resonances show blue and red shifts on heating and cooling of the device, the Mie resonance anti-reflection effect, RWA, and Mie resonances are tuned with large modulation depths reaching about 0.8 and 1 (see Fig. 3(e)–(h) and part 10 of the ESI†). The difference between the calculation and the experiment seen in Fig. 3(e) and (f) mainly originates from the inherent angle sensitivity of the sharp RWA resonance. It is numerically found that portions of slightly oblique incidence light due to the highly focusing objective lens (50×) in the microscopy setup critically suppress the sharp RWA resonance and induce the blue shifted resonance with a lowered *Q*-factor (see the detailed numerical analysis in parts 11 and 12 of the ESI†).<sup>57–61</sup>

## 2.4 Fano-resonant Rayleigh–Wood’s anomaly in a hybrid metagrating

In this section, we propose a Fano-resonant diffractive metagrating for enhanced *Q*-factor and efficiency. Since it is hard to make a high *Q*-factor even with conventional low loss dielectric metasurfaces and nanoantennas, achieving a high *Q*-factor with high signal efficiency and its reconfigurable modulation in lossy PCM metasurfaces is limited with conventional dielectric



**Fig. 4** (a) Schematic diagram of the hybrid metagrating ( $t = 80$  nm and  $w = 200$  nm). The subplots show top-view images of the scanning electron micrograph (up) and atomic force micrograph (down) of the fabricated device with a pitch of 400 nm. (b) Investigation of the pitch effect on the dynamically tunable metagrating. The 2-dimensional plots show the insulating phase absorptance (left) and phase-change-induced absolute difference of absorptance spectra (right) according to the varying pitch of the unit cell. The horizontal black dashed and oblique white dashed lines refer to the selected pitch condition ( $p = 400$  nm) and diffraction phase-matching condition, respectively. (c) Numerically calculated reflectance spectra in the insulating and metallic phases (solid lines) with Fano-resonance fitting results (dashed lines) (d) Normalized profiles of the  $E_y$  field (left) and magnetic field intensity (right) at the insulating phase resonance. The white streamlines in the left sub-figure denote the power flow of TM-polarized incident light toward the resonator. (e) Simulation and (f) measurement results of reflectance. The legends in (e) denote the filling factors of metallic VO<sub>2</sub>, and thus the 0 and 1 values correspond to the insulating and metallic phases, respectively. The legends in (f) denote the device temperature in the heating process.

resonances including the Mie resonant VO<sub>2</sub> metafilms proposed in the previous section.

Our idea to relieve these limitations is to combine the less lossy GPMR meta-atom (Fig. 1(a) and (d)) and RWA resonance phenomenon at the RWA diffraction condition ( $p = \lambda/n_{\text{sub}}$ ). Fig. 4(a) describes a schematic diagram of the metagrating device. The two inset figures of the scanning electron micrograph (upper) and atomic force micrograph (lower) of the top-view image show a fabricated metagrating with a 400 nm pitch produced by the conventional e-beam lithography technique using a thin sputtered VO<sub>2</sub> thin film (see the details in part 13 and 14 of the ESI†). In a diffractive regime ( $p < \lambda/n_{\text{sub}}$ ), as discussed in the previous section, the 1st and -1st diffraction channels are generated and RWA occurs with extraordinary resonance effects leading to sharp spectral change, which can be engineered by designing individual unit cell nanostructures.<sup>57–61</sup> In this case, to enhance the *Q*-factor, the low loss GPMR meta-atom is adopted. It is expected that the phase-change of VO<sub>2</sub> would only affect the strength of the sharp resonance peak at the RWA wavelength ( $\lambda_{\text{RWA}}$ ) but not the location of the resonance wavelength according to eqn (7). To investigate the principles of the hybrid metagrating, we calculated absorption spectra according to varying pitch length in the insulating phase as shown in Fig. 4(b) (left). With the fixed geometry of the unit cell structure, the pitch is chosen to be 400 nm for achieving a sharp resonance with large tunability within the visible regime (black dashed lines in Fig. 4(b)). With such an abrupt sharp absorption resonance peak, the reflectance peak is significantly modulated with a narrow and asymmetric Fano lineshape near resonance (Fig. 4(c)).<sup>57–61</sup> The numerically calculated and measured reflectance spectra in both VO<sub>2</sub> phases are successfully fitted to the conventional Fano resonance formula (Fig. 4(c) and part 15 of the ESI†),<sup>37</sup>

$$R = R_0 + \frac{H \left( \frac{(\omega - \omega_0)}{q} + \gamma \right)^2}{(\omega - \omega_0)^2 + \gamma^2}. \quad (8)$$

Here,  $R$ ,  $R_0$ ,  $H$ ,  $q$ ,  $\gamma$ ,  $\omega$ , and  $\omega_0$  refer to the reflectance, baseline reflectance, resonance strength, asymmetry factor, decay rate, angular frequency, and resonant angular frequency, respectively. The electromagnetic field profiles in Fig. 4(d) at the resonance show that the GPMR phenomenon with large field confinement occurs and a large amount of absorption is generated at the VO<sub>2</sub> region rather than the SiO<sub>2</sub> and a-Si:H regions. Gradual full-space modulation performances with enhanced *Q*-factor and efficiency are verified both in simulations (Fig. 4(e)) and measurements (Fig. 4(f)). Moreover, the transmittance dip can be modulated significantly near the RWA resonance (part 16 of the ESI†). Similar to the case in the previous section, the little discrepancy between the calculations and experiments seen in Fig. 4(e) and (f) mainly originates from the angle sensitivity of the sharp RWA resonance. Owing to portions of oblique incidence light due to the focusing objective lens (10×) in the microscopy setup (part 12 of the ESI†), the *Q*-factor and resonance location of the RWA resonance are a little distorted (see the detailed numerical discussions in part 17 of the ESI†).

### 3. Conclusions

In conclusion, inspired by Fano resonances with EMT and RWA in periodic nanophotonic gratings, we demonstrated simple and effective metasurface devices to dynamically modulate the visible light intensity in both reflection and transmission beams for the first time. We proposed and studied three different types of Fano-resonant grating devices with different modulation characteristics. By simultaneously achieving high modulation depths of both reflection and transmission, we expect that the proposed methods of all dielectric Fano-resonant dynamic light modulation could be applied for practical use in a wide range of optoelectronics technologies such as advanced high-resolution spatial light modulators, structural colours, smart windows, photodetectors, tunable anti-reflection coatings, tunable smart windows, thermal light sources, and image sensors that essentially require strong and efficient visible range operation. Moreover, the universal intuitive design rules provided in this paper would be also fruitful to design reconfigurable nanophotonic devices based on other active material platforms. It seems to be very promising that a spatial light modulator with the proposed metasurface grating pixels can be also demonstrated with pixel-by-pixel electro-thermal control if transparent conductors like indium tin oxide and graphene are introduced for electrical heating.

### Conflicts of interest

There are no conflicts to declare.

### Acknowledgements

This research was supported by the National Research Foundation of Korea (NRF) funded by the Ministry of Science and ICT (2020R1A2B5B02002730) and the BK21 Plus Project in 2020 (21A20131612805). The authors thank Dr Sang-Eun Mun for fruitful discussions. I. K. and J. R. acknowledge the Global PhD fellowship (NRF-2016H1A2A1906519) and a NRF grant (CMM-2019M3A6B3030637), respectively funded by NRF-MSIT of the Korean government. H. Y. and J. S. acknowledge the support from the Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Science and ICT (2020R1A2C2006389). Parts of this study have been performed using facilities at the IBS Center for Correlated Electron Systems, Seoul National University and Electronics and Telecommunications Research Institute.

### References

- 1 N. Yu and F. Capasso, *Nat. Mater.*, 2014, **13**, 139–150.
- 2 N. I. Zheludev and Y. S. Kivshar, *Nat. Mater.*, 2012, **11**, 917–924.
- 3 Y. Lee, S.-J. Kim, H. Park and B. Lee, *Sensors*, 2017, **17**, 1726.
- 4 P. Genevet and F. Capasso, *Rep. Prog. Phys.*, 2015, **78**, 024401.
- 5 X. Luo, D. Tsai, M. Gu and M. Hong, *Chem. Soc. Rev.*, 2019, **48**, 2458–2494.
- 6 M. Khorasaninejad, W. T. Chen, R. C. Devlin, J. Oh, A. Y. Zhu and F. Capasso, *Science*, 2016, **352**, 1190–1194.



- 7 M. A. Kats and F. Capasso, *Laser Photonics Rev.*, 2016, **10**, 735–749.
- 8 N. Yu, F. Aieta, P. Genevet, M. A. Kats, Z. Gaburro and F. Capasso, *Nano Lett.*, 2012, **12**, 6328–6333.
- 9 N. Yu, P. Genevet, M. A. Kats, F. Aieta, J. P. Tetienne, F. Capasso and Z. Gaburro, *Science*, 2011, **334**, 333–337.
- 10 Y. Yao, M. A. Kats, P. Genevet, N. Yu, Y. Song, J. Kong and F. Capasso, *Nano Lett.*, 2013, **13**, 1257–1264.
- 11 V. W. Brar, M. S. Jang, M. Sherrott, J. J. Lopez and H. A. Atwater, *Nano Lett.*, 2013, **13**, 2541–2547.
- 12 S. Kim, M. S. Jang, V. W. Brar, K. W. Mauser, L. Kim and H. A. Atwater, *Nano Lett.*, 2018, **18**, 971–979.
- 13 P. P. Iyer, N. A. Butakov and J. A. Schuller, *ACS Photonics*, 2015, **2**, 1077–1084.
- 14 Y. Horie, A. Arbabi, E. Arbabi, S. M. Kamali and A. Faraon, *ACS Photonics*, 2017, **5**, 1711–1717.
- 15 T. Lewi, P. P. Iyer, N. A. Butakov, A. A. Mikhailovsky and J. A. Schuller, *Nano Lett.*, 2015, **15**, 8188–8193.
- 16 J. Park, J. H. Kang, X. Liu and M. L. Brongersma, *Sci. Rep.*, 2015, **5**, 15754.
- 17 Y.-Y. Huang, H. W. Lee, R. Sokhoyan, R. A. Pala, K. Thyagarajan, S. Han, D. P. Tsai and H. A. Atwater, *Nano Lett.*, 2016, **16**, 5319–5325.
- 18 G. K. Shirmanesh, R. Sokhoyan, R. A. Pala and H. A. Atwater, *Nano Lett.*, 2018, **18**, 2957–2963.
- 19 S. Q. Li, X. Xu, R. M. Veetil, V. Valuckas, R. P. Domínguez and A. I. Kuznetsov, *Science*, 2019, **364**, 1087–1090.
- 20 M. J. Dicken, K. Aydin, I. M. Pryce, L. A. Sweatlock, E. M. Boyd, S. Walavalkar, J. Ma and H. A. Atwater, *Opt. Express*, 2009, **17**, 20975–20990.
- 21 S.-J. Kim, H. Yun, K. Park, J. Hong, J.-G. Yun, K. Lee, J. Kim, S. J. Jeong, S.-E. Mun, J. Sung, Y. W. Lee and B. Lee, *Sci. Rep.*, 2017, **7**, 43723.
- 22 M. Seo, J. Kyoung, H. Park, S. Koo, H. S. Kim, H. Bernien, B. J. Kim, J. H. Choe, Y. H. Ahn, H.-T. Kim, N. Park, Q.-H. Park, K. Ahn and D.-S. Kim, *Nano Lett.*, 2010, **10**, 2064–2068.
- 23 S.-J. Kim, S. Choi, C. Choi, Y. Lee, J. Sung, H. Yun, J. Jeong, S.-E. Mun, Y. W. Lee and B. Lee, *Opt. Express*, 2018, **26**, 34641–34654.
- 24 J. Rensberg, S. Zhang, Y. Zhou, A. S. McLeod, C. Schwarz, M. Goldflam, M. Liu, J. Kerbusch, R. Nawrodt, S. Ramanathan, D. N. Basov, F. Capasso, C. Ronning and M. A. Kats, *Nano Lett.*, 2016, **16**, 1237–1243.
- 25 M. A. Kats, D. Sharma, J. Lin, P. Genevet, R. Blanchard, Z. Yang, M. M. Qazilbash, D. N. Basov, S. Ramanathan and F. Capasso, *Appl. Phys. Lett.*, 2012, **101**, 221101.
- 26 F. Ligmajer, L. Kejik, U. Tiwari, M. Qiu, J. Nag, M. Konečný, T. Sikola, W. Jin, R. F. Haglund Jr, K. Appavoo and D. Y. Lei, *ACS Photonics*, 2018, **5**, 2561–2567.
- 27 N. A. Butakov, I. Valmianski, T. Lewi, C. Urban, Z. Ren, A. A. Mikhailovsky, S. D. Wilson, I. K. Schuller and J. A. Schuller, *ACS Photonics*, 2018, **5**, 371–377.
- 28 X. Yin, T. Steinle, L. Huang, T. Taubner, M. Wuttig, T. Zentgraf and H. Giessen, *Light: Sci. Appl.*, 2017, **6**, e17016.
- 29 A. Karvounis, B. Gholipour, K. F. MacDonald and N. I. Zheludev, *Appl. Phys. Lett.*, 2016, **109**, 051103.
- 30 N. R. Hosseini and J. Rho, *Appl. Sci.*, 2019, **9**, 564.
- 31 N. R. Hosseini and J. Rho, *Materials*, 2017, **10**, 1046.
- 32 A. L. Holsteen, A. F. Cihan and M. L. Brongersma, *Science*, 2019, **365**, 257–260.
- 33 E. Arbabi, A. Arbabi, S. M. Kamali, Y. Horie, M. F. Dana and A. Faraon, *Nat. Commun.*, 2018, **9**, 812.
- 34 A. L. Holsteen, S. Raza, P. Fan, P. G. Kik and M. L. Brongersma, *Science*, 2017, **358**, 1407–1410.
- 35 J. Sung, G.-Y. Lee, C. Choi, J. Hong and B. Lee, *Adv. Opt. Mater.*, 2019, **7**, 1801748.
- 36 M. A. Ansari, I. Kim, I. D. Rukhlenko, M. Zubair, S. Yerci, T. Tauqeer, M. Q. Mehmood and J. Rho, *Nanoscale Horiz.*, 2020, **5**, 57–64.
- 37 B. Luk'yanchuk, N. I. Zheludev, S. A. Maier, N. J. Halas, P. Nordlander, H. Giessen and C. T. Chong, *Nat. Mater.*, 2010, **9**, 707–715.
- 38 H. A. Haus, *Waves and fields in optoelectronics*, Prentice-Hall, 1984.
- 39 S. Fan, W. Suh and J. D. Joannopoulos, *J. Opt. Soc. Am. A*, 2003, **20**, 569–572.
- 40 A. Cordaro, J. V. D. Groep, S. Raza, E. F. Pecora, F. Priolo and M. L. Brongersma, *ACS Photonics*, 2019, **6**(2), 453–459.
- 41 J. W. Yoon and R. Magnusson, *Opt. Express*, 2013, **21**, 17751–17759.
- 42 J. Park, J. H. Kang, S. J. Kim, X. Liu and M. L. Brongersma, *Nano Lett.*, 2017, **17**(1), 407–413.
- 43 S.-E. Mun, H. Yun, C. Choi, S.-J. Kim and B. Lee, *Adv. Opt. Mater.*, 2018, **6**, 1800545.
- 44 M. Pu, C. Hu, C. Huang, C. Wang, Z. Zhao, Y. Wang and X. Luo, *Opt. Express*, 2013, **21**, 992–1001.
- 45 H.-T. Kim, B.-G. Chae, D.-H. Youn, S.-L. Maeng, G. Kim, K.-Y. Kang and Y.-S. Lim, *New J. Phys.*, 2004, **6**, 52–70.
- 46 H. Yoon, M. Choi, T.-W. Lim, H. Kwon, K. Ihm, J. K. Kim, S.-Y. Choi and J. Son, *Nat. Mater.*, 2016, **15**, 1113–1119.
- 47 I. H. Malitson, *J. Opt. Soc. Am.*, 1965, **55**, 1205–1208.
- 48 S. Kruk and Y. Kivshar, *ACS Photonics*, 2017, **4**, 2638–2649.
- 49 S. J. Kim and M. L. Brongersma, *Opt. Lett.*, 2017, **42**, 5–8.
- 50 M. G. Nielsen, A. Pors, O. Albrechtsen and S. I. Bozhevolnyi, *Opt. Express*, 2012, **20**, 13311–13319.
- 51 S. J. Kim, P. Fan, J. H. Kang and M. L. Brongersma, *Nat. Commun.*, 2015, **6**, 7591.
- 52 M. Esfandyarpour, E. C. Garnett, Y. Cui, M. D. McGehee and M. L. Brongersma, *Nat. Nanotechnol.*, 2014, **9**, 542–547.
- 53 S. J. Kim, J. Park, M. Esfandyarpour, E. F. Pecora, P. G. Kik and M. L. Brongersma, *Nano Lett.*, 2016, **16**, 3801–3808.
- 54 C. L. Holloway, M. A. Mohamed, E. F. Kuester and A. Dienstfrey, *IEEE Trans. Electromagn. Compat.*, 2005, **47**, 853–865.
- 55 W. Cai and V. Shalaev, *Optical metamaterials: fundamentals and applications*, Springer Science & Business Media, 2009.
- 56 D. H. Raguin and G. M. Morris, *Appl. Opt.*, 1993, **32**, 1154–1167.
- 57 M. Sarrazin and J.-P. Vigneron, *Europhys. News*, 2007, **38**, 27–31.
- 58 U. Fano, *J. Opt. Soc. Am. A*, 1941, **31**, 213–222.
- 59 M. Sarrazin, J.-P. Vigneron and J.-M. Vigoureux, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2003, **67**, 085415.
- 60 X. Zhang, S. Feng and T. Zhai, *Photonics Nanostruct.*, 2013, **11**, 109–114.
- 61 A. Maurel, S. Félix, J. F. Mercier, A. Ourir and Z. E. Djeflal, *J. Eur. Opt. Soc., Rapid Publ.*, 2014, **9**, 14001.

