

# Graphene–Silver Hybrid Metamaterial for Tunable and High Absorption at Mid-Infrared Waveband

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**Abstract**—We proposed a hybrid metamaterial structure based on graphene-silver, which has tunable and efficient light-absorbing performance at the mid-infrared regime. It is shown that this model can absorb light effectively with a maximum of 100% at the absorption peak wavelength. In addition, the absorption peak wavelength can be adjusted flexibly by changing parameters, including the chemical potential applied to graphene or other structural parameters of the structure. This nearly perfect absorption capability is attributed to the synergetic effects of the localized surface plasmon enhancement of Ag nanoribbon, graphene plasmonics, metal–insulator–graphene mode and Fabry–Perot resonance. This kind of tunable absorber can be used in mid-infrared photodetectors or spectroscopy fields.

**Index Terms**—Graphene, metamaterial, absorber, localized surface plasmon resonance enhancement, tunability.

## I. INTRODUCTION

OVER the past decades, metamaterials based perfect absorbers (MPAs) have been widely exploited for optoelectronic applications, including plasmonic biosensors [1]–[3], infrared thermal detection [4], [5], energy-harvesting devices [6]–[10]. Many MPAs with periodic nanostructures were proposed, which show a super absorptive property from visible to terahertz domain [11]–[14]. Noble metals, such as gold and silver, are widely used for the MPAs. Localized surface plasmon resonance (LSPR) excited by the metallic nanostructures is one of the main factors to enhance the interaction between the incident wave and the structure.

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However, the optical response of noble metals is passive and not easy flexible in tuning the absorption and resonating properties of the devices. This limits MPAs in applications such as spatial light modulators [15], [16].

In middle infrared and terahertz regime, graphene has prominent electric, optical properties and can behave like metals when it interacts with the incidence light and supports surface plasmons (SPs) [17]–[20]. This makes it useful in applications in transformation optics, tunable metamaterials, super-resolution and absorption enhancement [21]–[30]. Comparing with conventional metal SPs, graphene SPs can be tuned by electric gating or chemical doping and thus makes dynamically tunable resonant and active devices possible [31]–[33]. However, the absorption of graphene MPAs is relatively weak in the near-infrared range in that it is normally used as a lossy material without supporting SPs [34].

In view of the merits of noble metals and graphene, we proposed a graphene-based metamaterial absorber (GMA), which can support ultra-high absorption at the mid-infrared band. The finite-difference time domain (FDTD) method was employed to analyze GMA and the absorption of GMA is as high as 100% at wavelength of 4.55  $\mu\text{m}$ . Additionally, the absorption mechanism of GMA is numerically studied and the effect of different parameters of GMA on the absorption was also discussed in detail.

## II. METHODS AND STRUCTURE

Figure 1 show the proposed GMA structure which consists of Ag/SiO<sub>2</sub>/graphene/Polyethylene oxide (PEO)/Ag nanoribbon and the geometric parameters of the GMA. Due to realize the extremely high value for the chemical potential in graphene, we chose the PEO, as dielectric material bewtten the top Ag nanoribbon and graphene. An ion gel, which consists an ionic liquid (e.g. Li<sup>+</sup>) and a polymer (like PEO), can be realized in the experiments [35], [36]. The period of unit cells is 500 nm period in both  $x$  and  $y$  directions.

The model of graphene sheet was similar to and can be found in our previous work [25]. The graphene thickness was also assumed as 1 nm [37]–[40] in simulations, to make the numerical calculations reach proper convergence. In the simulation, the parameters of graphene are: Kelvin temperature  $T = 300$  K, Fermi velocity  $v_F = 10^5$  m/s, DC mobility  $\mu = 10000$  cm<sup>2</sup>V<sup>-1</sup>s<sup>-1</sup>, and the value of the chemical potential  $\mu_c$  was tuned from 1.0 eV to 2.0 eV.

According to [25, eqs. (1)–(3)], the relative permittivity of graphene with different  $\mu_c$  were calculated and shown

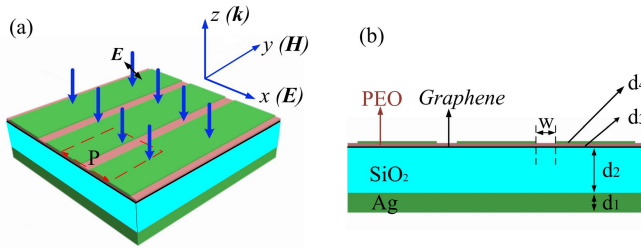


Fig. 1. The schematic diagram of the graphene-based metamaterial absorber with geometric parameters:  $P = 500$  nm,  $w = 100$  nm,  $d_1 = 100$  nm,  $d_2 = 230$  nm,  $d_3 = 20$  nm,  $d_4 = 10$  nm. (a) The perspective view and (b) the cross sectional view of the GMA.

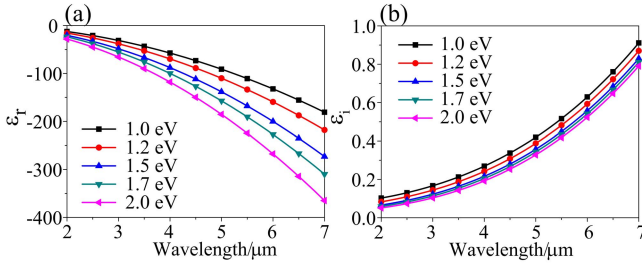


Fig. 2. Optical properties of graphene as a function of chemical potential in 2-7  $\mu\text{m}$  waveband. (a) The real and (b) imaginary part of permittivity of graphene, respectively.

in Figs. 2(a) and 2(b), respectively. As shown in Fig. 2(a), a smaller  $\mu_c$  results in a greater real part  $\epsilon_r$ . However, the imaginary part changes slightly when the  $\mu_c$  is increased, as depicted in Fig. 2(b).

By using the rigorous FDTD (Lumerical FDTD Solutions), the GMA structure was analyzed. The TM polarized plane wave at 4-6  $\mu\text{m}$  waveband incident in  $z$  direction. The inset of Fig. 1(a) shows the direction of incident waves with respect of the perspective of the proposed absorber. Periodic boundary conditions (PBC) was employed in both  $x$  and  $y$  directions and perfectly matched layer (PML) boundary condition in  $z$  direction. The optical constants of Ag are taken from literature [41]. The refractive index of PEO can be obtained from [42]. We assumed the chemical potential  $\mu_c$  of graphene is 1.7 eV and the relative permittivity was showed in Fig. 2.

The absorption was calculated by the equation  $A(\lambda) = 1 - R(\lambda) - T(\lambda)$ ,  $R(\lambda)$  is the reflection and  $T(\lambda)$  is the transmission. Due to the thickness of the bottom Ag layer (100 nm) is much larger than its skin depth, there is nearly no transmittance in the waveband, i.e.  $T(\lambda) = 0$ . Under this condition, the absorptivity can be simplified into  $A(\lambda) = 1 - R(\lambda)$ .

### III. RESULTS AND DISCUSSION

Frequency-domain field and power monitors were used to investigate the S-parameters of GMA. Figure 3 shows the absorption spectra of GMA (the red curve) for the normal incident light. The absorption has one peak at wavelength of 4.55  $\mu\text{m}$ , the corresponding absorption is 100%. In order to show the effect on absorption of graphene in GMA, we also calculated the structure without the graphene. The absorption of this structure (the green curve) was also shown in Fig. 3.

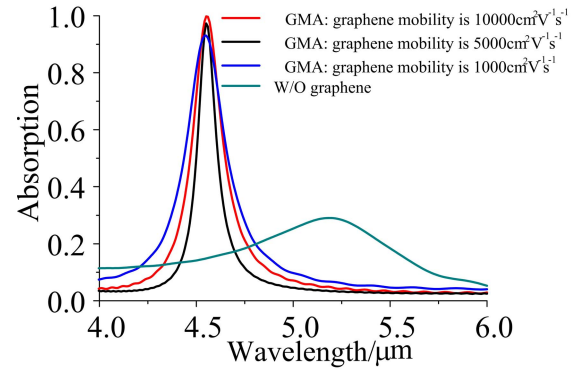


Fig. 3. Absorption spectra of GMA with graphene mobility  $10000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  (red), GMA with graphene mobility  $5000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  (black), GMA with graphene mobility  $1000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  (blue) and GMA without graphene (green).

of 5.18  $\mu\text{m}$ , however, the corresponding absorption is only 28.1%. As a result, it is confirmed that the graphene layer plays an important role in the enhancement of the absorption at this waveband. Electrolyte gates can be efficient at modulating the carrier concentration in graphene but also introduce high amount of scattering and the mobility in the layer is significantly reduced even down to  $1000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  [43]. Due to the scattering effect of PEO, we also showed the absorption curves of graphene with a mobility of  $1000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  and  $5000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  (the black and blue line), respectively. From Fig. 3, the resonant wavelength of absorption peak is almost unchanged and the maximal absorption just decreased to 97.3% and 93.0% when the mobility of graphene is  $5000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$  and  $1000\text{cm}^2\text{V}^{-1}\text{s}^{-1}$ , respectively. Therefore, the scattering, due to the PEO, does not have much effect on absorption.

The electric field distributions at different planes were calculated at wavelength of 4.55  $\mu\text{m}$  (the absorption peak) and the results were shown in Figs. 4(a)-4(d). Figure 4(a) shows the distribution of the electric field at  $y = 0$   $\mu\text{m}$  plane. From the color bar, one can observe clearly that the electric field, located at the edges of upper Ag ribbon/PEO interface, is enhanced greatly with more than 50 factors. The detail of electric field distribution of Ag ribbon at  $z = 124$  nm (3 nm above the Ag nano-ribbon) is shown in Fig. 4(b). One can find that the electric field is concentrated mainly at the edges of the Ag ribbon. This phenomenon can be attributed to the localized surface plasmon resonance (LSPR) effect [44]. Figure 4(c) shows the distribution of real  $E_z$  at  $y = 0$   $\mu\text{m}$  plane. As depicted in Fig. 2, at the waveband of 2-7  $\mu\text{m}$ , the graphene shows a metal-like properties. Therefore, from Fig. 4(c), it can be found the SPs exist in the graphene layer. The PEO film, sandwiched by the Ag nano-ribbon and graphene, forms a sandwich nanocavity. The standing wave (SW) pattern of SPs exists in the nanocavity as shown in Figs. 4(a) and 4(c). Since the nanocavity functions as a cavity resonator for SPs, the electric field is enhanced greatly with more than 30 factors. This can be attributed to the hybrid Metal-Insulator-Graphene (MIG) plasmon mode, like MIM (Metal-Insulator-Metal) mode, due to the proximity of the graphene layer to the Ag grating. The incident wave could penetrate the graphene layer, which is a one-atom-thick material. The detail of electric

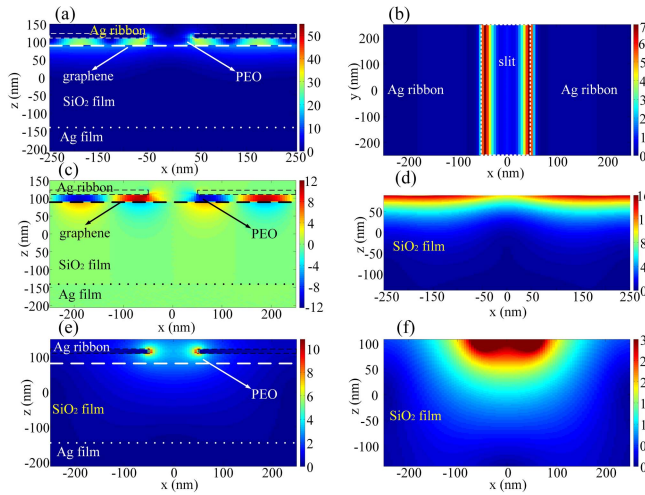


Fig. 4. (a) The distribution of the electric field of GMA in  $y = 0$  plane. (b) The distribution of the electrical field at  $z = 124$  nm plane. (c) The distribution of real  $z$  component of the electric field of GMA in  $y = 0$  plane. (d) The detail of the distribution of the electric field in the  $\text{SiO}_2$  film. (e) The distribution of the electric field of the simple Ag ribbon-PEO- $\text{SiO}_2$ -Ag structure in  $y = 0$  plane, with the same parameters of GMA. (f) The detail of the distribution of the electric field of the simple Ag ribbon-PEO- $\text{SiO}_2$ -Ag structure in  $\text{SiO}_2$  film.

field distribution of  $\text{SiO}_2$  film at  $y = 0 \mu\text{m}$  in Fig. 4(a) is shown in Fig. 4(d). As shown in Fig. 4(d), the intensity of the electric field in the  $\text{SiO}_2$  film is nearly zero close to the interface of the bottom Ag and  $\text{SiO}_2$  film.

As a comparison, the electric field distribution of the GMA without graphene layer at  $y = 0 \mu\text{m}$  plane at wavelength of  $5.18 \mu\text{m}$  was also calculated and shown in Figs. 4(e) and 4(f). From Fig. 4(e), the electric field is also concentrated mainly at the edges of the Ag nano-ribbon and the enhancement is only about 10 factors, which is much weaker than that of GMA with graphene. As shown in Fig. 4(f), due to lack of graphene plasmonics, there is no electric field enhancement in the  $\text{SiO}_2$  film. As a result, the absorption of GMA without graphene is much lower than that of GMA with graphene.

Therefore, considering the results shown in Fig. 4 and the discussion mentioned above, one can conclude that the physical mechanism for the strong absorption of GMA is the synergistic effect of localized surface plasmon enhancement, graphene plasmonics and MIG mode. Firstly, localized surface plasmon enhancement was excited by the Ag nano-ribbon. And graphene SPs was generated by this electric field enhancement. Due to the very large imaginary part of the refractive index, the Ag and graphene absorbs the incident wave greatly [20]. Then Fabry–Pérot formed between the Ag grating and the Ag back reflector contributes to the absorption of remaining energy as well.

To further understand the absorption performance of GMA, the effect of different parameters including the chemical potential  $\mu_c$  of graphene, the period ( $P$ ) and the thickness of the PEO ( $d_3$ ) and  $\text{SiO}_2$  ( $d_2$ ) on the absorption were studied and the results are shown in Fig. 5. Figure 5(a) shows the absorption versus wavelength for chemical potential  $\mu_c$  of graphene. From Fig. 5(a), it can be observed that the change of  $\mu_c$  has different effects on absorption and resonant wavelength.

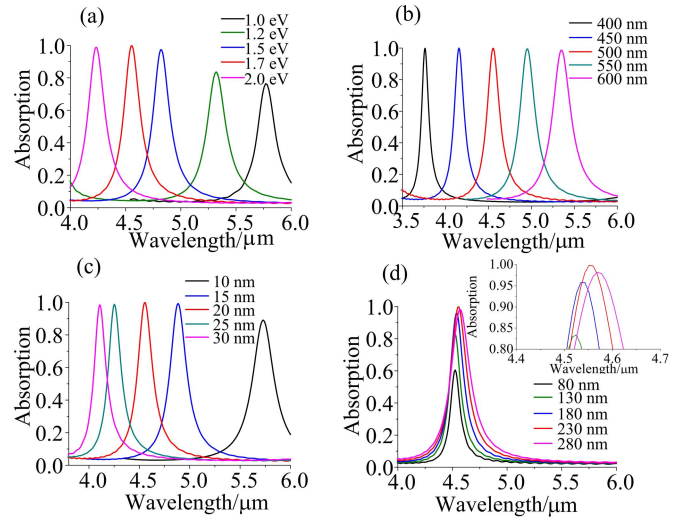


Fig. 5. The effect of different parameters of GMA on the absorption spectra: (a) chemical potential  $\mu_c$  of graphene, (b) the period ( $P$ ), (c) the thickness of the PEO ( $d_3$ ) and (d) the thickness of the  $\text{SiO}_2$  ( $d_2$ ). The chemical potential of graphene is 1.7 eV in (b), (c) and (d).

Specifically, the absorption peak wavelength has an obvious blue shift with the chemical potential  $\mu_c$  increases. For absorption, when  $\mu_c$  changes from 1.0 to 1.7 eV, the absorption increases from 76.4% to 100%. However, when  $\mu_c$  further increases to 2.0 eV, the absorption reduces significantly to 97.5%. Therefore, one can tune the absorption performance of GMA by changing the chemical potential of graphene. Figure 5(b) shows the effect of the period ( $P$ ) of GMA on the absorption. The period is changed from 400 to 600  $\mu\text{m}$  with a step size of  $\Delta P = 50 \mu\text{m}$ . As can be seen, there is almost no change for the absorption when the period changes. The absorption peak wavelength has an obvious red shift with the period increases. The wavelength of absorption peak changes with a direct dependence on periodicity of GMA. This is consistent with the reference [45]. Figure 5(c) shows the absorption versus wavelength for different thickness of PEO ( $d_3$ ), sandwiched by the Ag nano-ribbon and graphene. As shown in Fig. 5(c), it can be seen that the absorption peak wavelength has a blue shift with the thickness  $d_3$  increases. For absorption, when  $d_3$  changes from 10 to 20 nm, the absorption increases from 89.1% to 100%. However, when  $d_3$  further increases, the absorption has almost no change, only reduces slightly to 98.8% (25 nm) and 98.4% (30 nm). From Figs. 5(a)-5(c), one can find that the absorption peak wavelength can be tuned flexibly by simply changing of the parameters including chemical potential of graphene, the period and the thickness of the PEO film. Therefore, the Ag nano-ribbon, PEO film and graphene are the resonant layer of GMA. The effect of the thickness of the  $\text{SiO}_2$  film ( $d_2$ ) on the absorption is shown in Fig. 5(d). It can be seen that there is almost no change for the absorption peak wavelength when the  $d_2$  changes. To look into the details, the absorption curves at the peaks are magnified and shown in the inset of Fig. 5(d). One can find that the shifts of the resonant wavelength are no more than  $0.05 \mu\text{m}$ . The absorption of GMA increases from 60.5% ( $d_2 = 80$  nm) to 100% ( $d_2 = 230$  nm). However, the absorption just reduces



slightly to 98.2% when  $d_2$  further increases to 280 nm. The F–P vertical cavity are formed between the Ag grating and the bottom Ag film. Therefore, this effect originates from the vertical cavity between the top Ag nanoribbon and the Ag back reflector.

#### IV. CONCLUSION

In summary, we have proposed an efficient infrared graphene-based metamaterial absorber which can achieve an absorption of over 99%. The absorption property of GMA can be tuned easily by changing the chemical potential of graphene. The high absorptivity of the proposed hybrid absorber can be explained by the synergistic effect of the localized surface plasmon resonance enhancement, graphene plasmonics, MIG mode and F-P resonant. Our results can provide a valuable guide in applications including the design of plasmonic absorber and optical communications in the infrared range based on graphene.

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