Experimental demonstration of tunable graphene-polaritonic hyperbolic metamaterial

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Abstract: Tuning the macroscopic dielectric response on demand holds potential for actively tunable metaphotonics and optical devices. In recent years, graphene has been extensively investigated as a tunable element in nanophotonics. Significant theoretical work has been devoted on the tuning the hyperbolic properties of graphene/dielectric heterostructures; however, until now, such a motif has not been demonstrated experimentally. Here we focus on a graphene/polaritonic dielectric metamaterial, with strong optical resonances arising from the polar response of the dielectric, which are, in general, difficult to actively control. By controlling the doping level of graphene via external bias we experimentally demonstrate a wide range of tunability from a Fermi level of $E_F = 0$ eV to $E_F = 0.5$ eV, which yields an effective epsilon-near-zero crossing and tunable dielectric properties, verified through spectroscopic ellipsometry and transmission measurements.

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1. Introduction

Spectral tunability is key for controlling light-matter interactions, critical for many applications including emission control, surface enhanced spectroscopy, sensing, and thermal control. Particularly in the subwavelength range, tuning plasmonic resonances has been essential in controlling color, typically achieved by controlling the size of plasmonic nanoparticles, antennas and metamaterials [1–4]. In obtaining a large range of spectral tunability, it is preferable to operate near an optical resonance rather than a broadband plasmonic response. Nevertheless, it is in general easier to tune a broadband optical response rather than a resonant one since resonances in nanophotonics typically entail subwavelength-scale geometrical features.

From a very wide range of recently investigated metamaterials and heterostructures for spectral control, particular emphasis has been given to hyperbolic media, due to enhanced light-matter interactions arising from a larger range of wavenumbers available for propagating modes [5]. This property makes hyperbolic media attractive as hyperlenses, broadband thermal emitters, perfect absorbers, among others. These media are in generally uniaxial and support a hyperbolic frequency dispersion given by the equation [3,6–8]

$$\frac{k_x^2 + k_y^2}{\epsilon_\text{o}} + \frac{k_z^2}{\epsilon_\text{e}} = \frac{\omega^2}{c^2}$$

where $\epsilon_\text{e}$ and $\epsilon_\text{o}$ refer to the ordinary (in-plane) and extraordinary (out-of-plane) dielectric permittivity, respectively. Due to the different sign in $\epsilon_\text{e}$ and $\epsilon_\text{o}$, upon fixing the frequency $\omega$, the isofrequency diagram of the relevant electromagnetic modes opens up into a hyperbola, giving rise to a very large density of optical states, promising for waveguiding [9], emission engineering and Purcell enhancement [1,2,10] thermal photonics [11], lasing [12], and imaging [13,14]. Particularly, near the epsilon-near-zero frequency crossing of either $\epsilon_\text{e}$ or $\epsilon_\text{o},$ many
exciting phenomena can be supported, the most prominent of which is light propagation with near-zero phase advance [15–17].

There has been significant effort in frequency-tuning of the optical response of hyperbolic metamaterials [6,18–21]. For this, particular interest holds the case of graphene, a well-studied monolayer material for electronics [22] and in infrared photonics [23]. Namely, the optical properties of graphene can be dynamically tuned via optical pumping [24], or with electrostatic modulation of its carrier concentration with field-effect gating [25,26], often targeting tunable plasmonic properties [27–29]. The high degree of localization of graphene plasmons, together with the dielectric tunability of graphene provides a promising platform for investigating tunable graphene-based hyperbolic metamaterials. There has already been considerable theoretical effort in the past decade to understand the properties of tunable graphene metamaterials [30–34], with significant focus on the potential of tuning hyperbolic properties of graphene/dielectric planar heterostructures [6,35–38]. There have previously been experimental demonstrations of graphene-based hyperbolic media [39,40], nevertheless, the reported properties have remained fixed at the time of fabrication. No post-fabrication way to control the dielectric permittivity tensor (\(\epsilon_o\) and \(\epsilon_e\) in Eq. (1)) has been reported until now. In this work, we experimentally demonstrate large dynamical tuning ratio of the dielectric response of a graphene-based hyperbolic metamaterial.

Gating graphene, when integrated with dielectric layers, is difficult due to graphene’s two-dimensional nature with weak out-of-plane Van der Waals bonds that yield poor adhesion to most dielectric substrates. Furthermore, large-area graphene sheets on the order of few mm\(^2\) with gate-induced tunability are needed to perform metamaterial optical measurements at infrared frequencies. Exfoliated flakes are generally limited to sizes of tens of \(\mu m^2\), so large-area graphene samples grown by chemical vapor deposition and subsequently transferred from their growth substrates, are necessary. Additionally, deposition of large-area thin dielectric layers on graphene is challenging. Films prepared by electron-beam evaporation exhibit thermal stress-induced delamination [41]. Films grown by atomic layer deposition (ALD) with an H\(_2\)O precursors exhibit difficulty in bonding to chemically-inert hydrophobic graphene [42], whereas ozone-based ALD processes oxidize graphene.

Here, we discuss how we overcome these challenges and are, thus, able to tune a graphene-based hyperbolic metamaterial unit cell for a wide range of doping levels in graphene translating to a Fermi level that ranges from \(E_F = 0\ eV\) to \(E_F = 0.5\ eV\), without dielectric breakdown. Previous theoretical proposals have considered non-dispersive dielectric materials [6,35–37], thereby yielding a broadband hyperbolic response. By contrast, here, we consider a polaritonic dielectric material, namely SiO\(_2\). The polaritonic resonances that all polar materials exhibit at infrared frequencies, at their Reststrahlen band, are typically not tunable, as they constitute a fundamental material property. Furthermore, resonances in nanophotonics are typically induced via subwavelength photonic structures, which are also quite difficult to dynamically tune. We show here that, upon the integration of graphene, it is feasible to actively tune the polaritonic resonances of naturally occurring dielectrics. Graphene provides a tunable character to the in-plane response of the composite graphene/SiO\(_2\) heterostructure, and its plasmonic nature assigns a hyperbolic frequency region near the polar resonance of SiO\(_2\), at a free-space wavelength of 20 \(\mu m\). We are therefore able to experimentally observe, through multi-angle spectroscopic ellipsometry and transmittance measurements, a tunable epsilon-near zero permittivity along the in-plane direction near the surface phonon polaritonic resonance while leaving the out-of-plane response unchanged (due to the two-dimensional nature of graphene), thereby yielding a widely tunable hyperbolic response.
2. Metamaterial motif and experimental realization

The metamaterial under consideration is depicted in Fig. 1, and is composed of a graphene monolayer sandwiched between two SiO$_2$ layers of thickness 300 nm. The alumina (Al$_2$O$_3$) layers depicted in Fig. 1 have thickness 0.5 nm and are placed to prevent poor graphene adhesion. Particularly, a viable dielectric deposition method was developed consisting of functionalization of the surface by deposition of trimethylaluminium (TMA) [43] or an aluminum nucleation layer [44] to create a seed layer for additional deposition. A suitably thin layer of aluminum is needed so that it can fully oxidize and not compromise the electrical gating of the graphene. We found that deposition of Al$_2$O$_3$ via plasma-enhanced chemical vapor deposition (PECVD) resulted in reduced thermal stress and avoided delamination. The graphene is grown by chemical vapor deposition (CVD) and transferred onto the thermal oxide, whereas the top SiO$_2$ film is deposited by plasma-enhanced chemical vapor deposition (PECVD). The thickness of the film layers were measured by both a thin film analyzer and visible ellipsometry with a qualitative agreement of 2nm. Lithographically-defined patterns were used to deposit 3nm/100nm of Cr/Au contacts on the graphene layer, and were used to gate the graphene monolayer against the silicon substrate, which serves as the back-side contact for field-effect tuning.

![Fig. 1. Left: Schematic of a theoretical metamaterial stack. Right: Schematic of the fabricated individual device. The layers: Lightly-doped silicon substrate, thermally-grown SiO2, Al$_2$O$_3$, transferred chemical-vapor deposited (CVD) graphene, Al$_2$O$_3$, and plasma-enhance chemical vapor deposition SiO$_2$. The thin layers of Al$_2$O$_3$ are necessary for the feasibility of the fabrication. The thick SiO$_2$ contribute to the majority of the dielectric response. Contacts are added to gate and measure the resistance of the graphene. The graphene is tuned by gating against the back silicon substrate.](image)

3. Theory and modeling

Since the composite in Fig. 1 is extremely subwavelength to infrared light, one can homogenize it and assign an effective in-plane and out-of-plane dielectric response, namely $\epsilon_o$ and $\epsilon_e$ [8]. The two-dimensional nature of graphene leaves the out-of-plane response unaffected, therefore in the out-of-plane direction, this metamaterial behaves to far-field radiation effectively as bulk SiO$_2$. By striking contrast, by electrostatically tuning the graphene carrier we can shift the
epsilon-near-zero point of $\epsilon_0$, and therefore control the hyperbolicity of the heterostructure as shown in Fig. 2.

### Fig. 2. Ellipsometrically derived effective in-plane dielectric permittivity, $\epsilon_0$, for the graphene/SiO$_2$ metamaterial of Fig. 1, under applied bias, for three different Fermi levels $E_F = 0$ eV, $E_F = 0.3$ eV, $E_F = 0.5$ eV. Grey and black curves correspond to the homogeneous dielectric permittivity of the bottom and top SiO$_2$ films, respectively. (a) Imaginary part, and (b) real part. (c) Inset showing the epsilon-near-zero regime of $\epsilon_0$ at different $E_F$’s.

In estimating the Fermi level to which we can actively tune the doping level in graphene, we use a capacitor model based on the materials between the gate and the applied voltage [45].

$$E_F = 0.031\sqrt{V - V_{\text{Dirac}}}.$$  \hspace{1cm} (2)

Experimentally, the location of the Dirac peak was determined via measuring change in sheet resistance. Furthermore, we use the Kubo formula [46] to calculate the sheet conductance $\sigma$ from the $E_F$ of graphene. This value can be used to compute the transfer matrix for graphene [47].

$$\vec{G} = \begin{bmatrix} 1 & 0 \\ 4\pi\sigma/c & 0 \end{bmatrix}$$ \hspace{1cm} (3)

We utilize the transfer matrix approach [48], accounting for graphene via $\vec{G}$, and obtain the complex scattering amplitudes of the fields at different Fermi levels $E_F$. In these calculations, fabrication and material imperfections are removed by having, a priori, measured experimentally the individual layer thicknesses and optical constants of all thin films in the metamaterial, with ellipsometry. For example, in Fig. 2(a) and (b) we show the experimentally determined dielectric permittivity of the top and bottom SiO$_2$ films shown in Fig. 1, where their small differences are expected since the top SiO$_2$ is deposited via PECVD whereas the bottom one is thermally grown. The scattering amplitudes are fed into previously developed parameter retrieval approaches [8], from which we obtain an effective uniaxial tensorial dielectric permittivity $\epsilon = \text{diag}(\epsilon_0, \epsilon_0, \epsilon_e)$ that characterizes the metamaterial composite. This process is repeated at different gating voltages $V$, in other words for different Fermi levels $E_F$. 
4. Results

By taking spectroscopic ellipsometry measurements of the full metamaterial stack of Fig. 1, we perform an ellipsometric fitting where we use the effective dielectric permittivity \( \epsilon = \text{diag}(\epsilon_o, \epsilon_o, \epsilon_e) \) as a model to fit to the experimental data, namely the ellipsometric observables \( \Psi \) and \( \Delta \). In Fig. 2(a) and (b) we show the imaginary and real part of the ellipsometrically-derived in-plane permittivity \( \epsilon_o \), at different Fermi levels \( E_F \). We note that the out-of-plane effective permittivity \( \epsilon_e \) is not tunable as described above, and therefore is omitted. There resonant character of \( \epsilon_o \) near the regime of 20 \( \mu \)m is attributed to the surface phonon polaritonic resonance of SiO\(_2\) at this wavelength, nevertheless this resonance has now become tunable via incorporation of a monolayer-thick graphene sheet in between SiO\(_2\) films. As can be clearly seen in 2(c), by gradually tuning the Fermi level of graphene from \( E_F = 0 \) eV (blue curves) to \( E_F = 0.3 \) eV (green curves) to \( E_F = 0.5 \) eV, we redshift the infrared response of the metamaterial by approximately a micron, i.e. from a near-zero crossing at 20 \( \mu \)m under no bias to 19 \( \mu \)m under large applied bias. Redshifting is expected as a response of applied bias because the electrostatic doping induces

![Graphs showing transmission measurements](image)

**Fig. 3.** (a) Absolute FTIR transmission measurements over a range of Fermi levels from \( E_F = 0 \) eV to \( E_F = 0.6 \) eV. (b) Experimental data normalized to \( E_F=0 \), the Dirac point of graphene, in order to emphasize the gate-tunable response of the metamaterial. (c) Experiment compared with theory. The theoretical results were computed using the ellipsometrically derived dielectric properties of the metamaterial, and the experimentally measured thicknesses of each constituent layer. Data are normalized to the case of \( E_F=0 \), similar to panel (b). Deviations between theory and experiment arise due to hysteresis of the graphene induced by charge trapping.
additional charge carriers in the graphene sheet, hence making the composite medium more metallic.

In addition to spectroscopic ellipsometry, we perform Fourier-transform infrared spectroscopy (FTIR) to measure the sample transmission, and compare with the results of spectroscopic ellipsometry shown above, derived based on initial parameter retrieval-based derivation of $\epsilon = \text{diag}(\epsilon_o, \epsilon_o, \epsilon_e)$. Electrostatically gating the graphene induces changes in the transmission of the composite metamaterial, as shown in Fig. 3. Namely, as mentioned above, gating the graphene monolayer makes the composite metamaterial more metallic and, therefore, less transmissive, as shown with the colormap in Figs. 3(b) and (c). The dips near the wavelengths of 16 $\mu$m and 20 $\mu$m correspond to the two surface phonon polariton resonances of SiO$_2$, where the material absorbs resonantly, resulting in low transmittance. We note that, experimentally, graphene exhibits hysteresis, which is attributed to defects induced by the deposition of the aluminum layer, resulting in the discrepancies between experiment and theory. As the graphene is tuned, the Dirac peak shifts in the direction of applied bias, causing the sample to experience a reduced $E_F$, giving qualitative experimental agreement with theory without fitting parameters as can be seen in Fig. 3(c).

To further illustrate the epsilon-near-zero shifting and the resonant nature of the in-plane dielectric response of this metamaterial, i.e. $\epsilon_o$, in Fig. 4 we show the relative change in dielectric permittivity, i.e. $\Delta \epsilon = 100 \times (\epsilon_o,V=0 - \epsilon_o,V=V)/\epsilon_o,V=0$, for two different applied bias corresponding to $E_F=0.2$ eV and to $E_F=0.4$ eV, with blue and red color, respectively. These calculations were performed using the experimentally derived values for the optical properties and thicknesses of the constitutive components of the metamaterial, as described above. Near the surface phonon resonance of SiO$_2$ at 20 $\mu$m, significant tuning of the real part of $\epsilon_o$ is observed, coming from the epsilon-near-zero tuning, which shifts by approximately 1 micron. Bearing in mind that the out-of-plane response of this metamaterial ($\epsilon_e$ in Eq. (1)) is not tunable due to the two-dimensional nature of graphene, as explained above, the change in sign of $\epsilon_o$ on the left axis in Fig. 4 corresponds to a topological transition of the isofrequency surface of this metamaterial.

![Fig. 4. Relative change of the effective in-plane dielectric permittivity, $\epsilon_o$, of the metamaterial, for two different Fermi levels, namely $E_F = 0.2$ eV (blue curves) and $E_F = 0.4$ eV (red curves), with respect to the case $E_F = 0$ eV. Solid curves (left y-axis) correspond to real parts and dashed curves correspond to imaginary parts (right y-axis).](image-url)
5. Conclusions

In summary, we have experimentally demonstrated a graphene/polaritonic dielectric metamaterial with tunable epsilon-near-zero permittivity response. By tuning the Fermi level of graphene by 0.5 eV, we observe a shift of 1 µm in the near-zero response. Although previous theoretical proposals have focused on non-dispersive dielectric materials between graphene monolayers, here we showed that utilizing the polar response of dielectrics at infrared frequencies benefits tunability, and additionally provides means of tuning constitutive material properties of polar dielectrics and semiconductors, by incorporating graphene. Ellipsometry was used to determine the optical properties (dielectric response and thickness) of the constituent materials, and, based on effective parameter retrievals that homogenize the metamaterial, we experimentally characterized the full metamaterial stack. FTIR transmission measurements agree with our ellipsometric results, where transmission reduction is directly attributed to electrostatically induced charges in graphene and to epsilon-near-zero tuning.

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Disclosures
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