Optically induced metal-to-dielectric transition in Epsilon-Near-Zero metamaterials

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Epsilon-Near-Zero materials exhibit a transition in the real part of the dielectric permittivity from positive to negative value as a function of wavelength. Here we study metal-dielectric layered metamaterials in the homogenised regime (each layer has strongly subwavelength thickness) with zero real part of the permittivity in the near-infrared region. By optically pumping the metamaterial we experimentally show that close to the Epsilon-Near-Zero (ENZ) wavelength the permittivity exhibits a marked transition from metallic (negative permittivity) to dielectric (positive permittivity) as a function of the optical power. Remarkably, this transition is linear as a function of pump power and occurs on time scales of the order of the 100 fs pump pulse that need not be tuned to a specific wavelength. The linearity of the permittivity increase allows us to express the response of the metamaterial in terms of a standard third order optical nonlinearity: this shows a clear inversion of the roles of the real and imaginary parts in crossing the ENZ wavelength, further supporting an optically induced change in the physical behaviour of the metamaterial.

Recent advances in metamaterial science have opened routes to unprecedented control over the optical properties of matter, with a wide array of applications and implications for novel light-matter interactions. Examples are the demonstration of negative index materials and, more recently, significant attention has been devoted to the behaviour of light in a medium with zero dielectric permittivity. We will refer to these materials as Epsilon-Near-Zero (ENZ) materials with the implicit assumption that in all passive materials, the ENZ condition will only be met for the real part of the permittivity, \( \varepsilon' \), (as a result of absorption that will always imply that the imaginary part is greater than zero) and at one single wavelength (due to dispersion). Such ENZ materials may either occur naturally at the plasma frequency or may result from engineering the propagation medium, for example so that light propagates in a waveguide near the cutoff frequency. Another option, investigated here, is to create a metamaterial made of deeply subwavelength alternating layers of dielectric and metal with thicknesses that are chosen such that \( \varepsilon' \) is zero at a chosen wavelength. The linear properties of ENZ metamaterials have been investigated in depth with a range of applications for example in novel waveguiding regimes and for controlling the radiation pattern of electromagnetic sources. The ENZ condition has also been predicted to have far-reaching consequences in terms of the effective optical nonlinearity of the metamaterial, but with limited experimental evidence. A compelling experimental evidence of the role of ENZ properties affecting the optical nonlinearity is the recent demonstration of efficient third harmonic generation due to the enhancement of the pump electric field longitudinal component in a uniform film of Indium-Tin-Oxide (ITO).

A different, yet related area of study, is the search for materials that can be optically controlled so as to exhibit a sharp and rapid transition from metallic to dielectric (or vice versa), thus implying a fundamental change in the material properties. Examples that have been investigated and observed in literature are optically induced phase transitions (for example in Vanadium Oxide and other compounds) or sudden increase in conductivity in glass when optically pumped with single cycle pulses, close to the breakdown damage threshold. A metal-to-dielectric transition has also been theoretically proposed in metallo-dielectric stacks, obtained by

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optically pumping close to the ENZ wavelength with relatively fast ~1 ps response times. Here we experimentally investigate the optical behaviour of MMs made of deeply subwavelength alternating layers of fused silica glass and silver with thicknesses that are chosen such that $\varepsilon' = 0$ in the near-infrared region, as shown in Fig. 1.

We show that by optically pumping the MM far from the ENZ wavelength, it is possible to induce a marked and rapid (on the same time scale of the 100 fs pump pulse) transition from metallic ($\varepsilon' < 0$) to dielectric ($\varepsilon' > 0$). By changing the layer thickness of the MM we can also control the wavelength at which this transition occurs. Remarkably we find that the metal-dielectric transition occurs linearly as a function of the pump intensity. We show that this fact allows to describe the optical behaviour in terms of a standard third order nonlinear susceptibility. Our measurement technique provides the full complex amplitude across a wide spectral range centred at the ENZ wavelength.

Results

We fabricated the metamaterial samples by electron beam deposition of alternating layers of Ag and SiO$_2$ on a thick (1 mm) SiO$_2$ substrate with a total of 10 layers. The Ag thickness in each layer is kept at 5 nm whilst the SiO$_2$ thickness is the same in each layer and tuned to 80–70 nm in order to provide the ENZ condition around 820–890 nm, in the centre of our laser tuning region. To obtain smooth continuous layers of silver on silica below the standard percolation limit, we seeded the deposition of each metal layer with 0.7 nm of Germanium.

Figure 1 shows a photograph of one of the samples and an SEM image of the multilayer structure, respectively. More details on the fabrication process are provided in the Methods section.

**Linear response.** The linear response (real and imaginary part of $\varepsilon$, $\varepsilon'$ and $\varepsilon''$, respectively) was measured by a standard reflection/transmission measurement (see Methods for details) and is shown as the dashed lines in Fig. 1(c,d) for two different samples with SiO$_2$ thickness equal to 80 nm (referred to as “sample A”) and 70 nm (“sample B”), respectively. $\varepsilon'$ is measured to be zero at 885 nm (sample A) and 820 nm (sample B).

Under the approximation of deeply subwavelength films, light polarised parallel to the film does not interact with each individual layer of the multilayer structure but rather with an effective homogenised medium whose complex dielectric susceptibility is given by $\chi_{\text{eff}} = (l_d \chi_d + l_m \chi_m)/(l_d + l_m)$, where $\chi$ may represent either the linear
that also increase with pump intensity. Inset (b) shows material damage. In the inset (a) to Fig. 3 we also show the corresponding imaginary parts of the permittivity over which the optically-induced metal-dielectric transition occurs for the maximum pump power (limited by on sample A for a pump power of 17 GW/cm2: the rise time is of the order of the 100 fs pump pulse duration (followed by a decay time of a few ps that is typical for Ag). We see that by tuning the delay, it is possibly to tune the Δε to positive, thus indicating a transition of the medium from metallic to dielectric. The total variation between the pumped and not pumped case (within the noise limit of our detectors, a few percent). This can be explained by a simultaneous change in absorption that eventually balance a variation in the light transmitted into the sample. We thus consider ΔT = 0.

While the sudden change in reflectivity is clearer for sample A, we still observe a significant variation also for sample B, yet with a broader response. As we will show in the following, different effects contribute to the change in reflectivity ΔR/Rlin, including the dispersion of the χ(3) and the transition through the ENZ wavelength. A thorough analysis of the variation of the permittivity with the pump intensity and probe wavelength is therefore required to unveil the underlying processes. We thus extend the same method followed to extract the linear permittivity ε′ from Rlin and Tlin, to also extract ε in the presence of the pump: the values of the reflectivity and transmissivity in the nonlinear (pumped) case allow to retrieve the nonlinear (pumped) value of the permittivity. In particular, here we are interested in the behaviour just above the ENZ wavelength: as can be seen in Fig. 1, here the unpumped ε′ is negative. In the presence of a positive and sufficiently large increase in ε′ due to the optical pump we may predict that the permittivity will transition from below to above zero. Figure 3 shows ε′ as a function of pump intensity for sample A and sample B, measured at 890 nm and 825 nm, respectively (in both cases, 5 nm above the ENZ wavelength). As can be seen, in both cases the permittivity transitions from negative to positive, thus indicating a transition of the medium from metallic to dielectric. The total variation Δε′ ∼ 0.05 is of the same order of the absolute value of the permittivity itself, implying a relatively large bandwidth of ∼ 10 nm over which the optically-induced metal-dielectric transition occurs for the maximum pump power (limited by material damage). In the inset (a) to Fig. 3 we also show the corresponding imaginary parts of the permittivity that also increase with pump intensity. Inset (b) shows ε′′ as a function of the relative pump-probe delay measured on sample A for a pump power of 17 GW/cm2; the rise time is of the order of the 100 fs pump pulse duration (followed by a decay time of a few ps that is typical for Ag). We see that by tuning the delay, it is possibly to tune the precise value of ε′′, crossing from metal to dielectric and back again.

A notable feature of this data is the clear linear dependence of both ε′ and ε″ with pump power (in disagreement for example with the theoretical predictions of Husakou et al.32): the dashed lines in Fig. 3 represent linear fits to the data, which are seen to pass through the value measured in the absence of the pump (and reported in Fig. 1, as expected). We also note that this linear behaviour was observed over a wide range of wavelengths (700 nm to 1000 nm, data not shown). This feature is remarkable as it allows us to relate the behaviour of the MM and the transition from metal to dielectric in terms of a standard third order nonlinear susceptibility,
Indeed, because of this linear behaviour we can extract the nonlinear susceptibility tensor element as (see Methods for details):

\[
\chi^{(3)}(\omega_p, \omega_p) = \frac{n_p \varepsilon_0 c}{3} \frac{\partial \varepsilon(\omega_{pr}, I_p)}{\partial I_p},
\]

where \(\omega_{pr}\) and \(\omega_p\) are the probe and pump frequencies, respectively, and \(n_p\) is the real part of the medium refractive index at the pump frequency.

The complex values of the permittivity, \(\varepsilon(\omega_{pr}, I_p)\), were retrieved from the reflectivity and transmissivity measurements at different probe wavelengths and pump intensities. We used the transfer matrix approach to determine the value of permittivity that results in the measured reflectivity and transmissivity\(^35\) (see Methods for further details). The nonlinear susceptibility is then calculated from these values using Eq. (1).

We note that the linearity observed in the variation of \(\varepsilon\) with pump intensity (as shown in Fig. 3) implies that Eq. (1) is consistent, as it provides us with \(\chi^{(3)}\) values that are constants (do not depend on \(I_p\)). We remark that this method differs from the simple retrieval of the permittivity in the pumped case, as it exploits the found linear behaviour of \(\varepsilon\) versus intensity to interpret the nonlinear mechanism in term of a third-order nonlinearity, also allowing to extract the complex value of the \(\chi^{(3)}\) tensor at different pump and probe wavelengths.

This derivation neglects the variation of the pump intensity inside the sample along the propagation direction (due to absorption). Averaging the intensity over the sample thickness would lead to a small correction factor \(\sim 2\) for the values of \(\chi^{(3)}\), thus here we chose the simplified formulation.

We use Eq. (1), applied separately to the real and imaginary part of \(\varepsilon\) to plot the real and imaginary parts of \(\chi^{(3)}\). Figure 4(a,c) show the real (solid blue line) and imaginary (dashed red line) third order nonlinear coefficients \(\chi^{(3)}\) and \(\chi^{(3)}\) for samples A and B, respectively. Figure 4(b,d) show the same data but plotted as the absolute value \(|\chi^{(3)}|\) (solid blue line) and phase, \(\phi\) (dashed red line). The notable feature of these results is that as the probe wavelength crosses from the dielectric-like region (\(\varepsilon' > 0\)) to the metallic-like region (\(\varepsilon' < 0\)), the \(|\chi^{(3)}|\) changes in nature from predominantly real to predominantly imaginary.

**Discussion**

The maximum measured \(|\chi^{(3)}|\) is of the same order or slightly larger than that measured by other means in bulk silver \(|\chi^{(3)}| = 2.810^{-19} \text{m}^2/\text{V}^2\)\(^34,36\). However, it is interesting to note that whereas silver cannot be used in transmission in samples thicker than \(\sim 40\) nm (all light is reflected from the surface at these thicknesses), the metamaterial is significantly thicker and more than two orders of magnitude thicker than the total thickness of silver present in the sample. This therefore provides a much longer effective interaction length for light with the high nonlinearity of the metal, as compared to the bare material. Note that the increased transmission is a known result for alternating layers of dielectric and metals, and it is for example exploited in one-dimensional metal-dielectric photonic crystal\(^37,38\).

We also underline that the large imaginary component of the \(\chi^{(3)}\) tensor measured in the "metallic" wavelength region does not imply larger nonlinear losses. Indeed, third order nonlinearities lead to nonlinear phase shifts and nonlinear absorption that are determined by the real and imaginary part of nonlinear refractive index, respectively (and not of the susceptibility). These are given by (in the degenerate case)\(^39\):

\[
\beta_1 = 3/(4\pi \varepsilon_0 c) |\chi^{(3)}|/D \quad \text{and} \quad \beta_2 = 3/(4\pi \varepsilon_0 c) [n_\lambda \chi^{(3)} - n_\chi^{(3)}]/D \quad \text{where} \quad D = n_\chi (n_r^2 + n_i^2). \]

The imaginary part of \(n_\lambda\) is usually associated to what is known as the nonlinear absorption coefficient, \(\beta_1 = 4\pi n_\lambda /\lambda\), where \(\lambda\) is the vacuum wavelength. We underline that \(n_\lambda\) and \(\beta_1\) depend respectively, on the sums and differences of the real and imaginary parts of of the \(\chi^{(3)}\) tensor. This implies for example that a large \(\chi^{(3)}\) will enhance Kerr effects such as phase modulation that are associated to the \(n_\lambda\) coefficient whilst simultaneously minimising nonlinear absorption (the data in Fig. 4 shows that actually \(\beta_2 \sim 0\) close to the ENZ wavelength).

In conclusion, ENZ metamaterials allow to tailor and access novel optical propagation regimes. The interplay between the linear and nonlinear, real and imaginary propagation constants leads to metamaterials that exhibit nonlinearities with amplitudes similar to those of metals yet spread over material thicknesses two orders of
magnitude larger and with substantially reduced losses. We have also shown that it is possible to optically induce a metal-dielectric transition over a wide bandwidth that can support ~100 fs laser pulses. Further optimisation by e.g. reducing losses (and hence the linear refractive index) by introducing a gain medium can enable new forms of efficient switches for light and even a platform for non-perturbative nonlinear optics at low light intensities.

Finally we would like to highlight that our results are not specific to the sample used in our experiments, rather they can be generally applied to all ENZ media. Indeed, for these materials, close to the ENZ wavelength we can expect to observe a region where $\varepsilon_r$ is slightly negative, and provided that the nonlinear response is fast enough and the $\chi^{(3)}$ is positive, we might expect similar results (alternatively for materials with negative $\chi^{(3)}$, similar results can be obtained at a wavelength displaying a slightly positive real permittivity). For example, non-structured materials exhibiting an accessible ENZ region in the near infrared, such as transparent conductive oxides (TCOs), are expected to display a similar behaviour. In aluminium-doped zinc oxide (AZO) thin films ultrafast switching close to the ENZ wavelength at 1300 nm has been recently observed, and this could be a suitable candidate to observe a similar dynamics.

Methods
Fabrication. The ENZ metamaterial was fabricated on 1 mm thick slides of SiO$_2$. The substrate was thoroughly cleaned in ultrasonic assisted baths of Acetone and Isopropanol (5 mins each) and blown dried with N$_2$ flow. The metallo-dielectric stack was deposited using an EDWARD auto 306 electron beam evaporator, with a base pressure below 3 $\times$ 10$^{-6}$ bar. The deposition rate was kept below 0.1 nm/s for the metals and below 0.3 nm/s for the SiO$_2$ to grant uniformity. The thickness of the seeding Ge layer (0.7 nm) was chosen after a thorough optimisation process, to minimise the losses of the thinnest achievable layer of Silver (5 nm). An excess of 50 independent evaporations was performed to determine the final values. The development process was complemented by a combination of SEM, STEM and AFM measurements on sacrificial test samples.

Linear characterisation. To characterise the linear properties of the MM we built a simple setup for the measurement of the reflection and the transmission. An Ocean Optics HL2000 Halogen Lamp covering the visible-NIR spectrum was collimated with a telescope to a beam radius of ≈5 mm, with controlled polarisation.
A thin film beam splitter was used to separate the reflected light from the incident one. The two beams were then focused on the tip of multimode fibres and analysed with two Ocean Optics spectrum analysers. Light was impinging on the sample on the side of the multilayer. The reference for the reflection was a silver mirror, for the transmission we recorded the collected light without any sample. The raw data were then used to retrieve the effective complex permittivity of the multilayer, using a standard least mean square fitting procedure: the experimental reflected and transmitted value vs $\lambda$ where compared with those calculated with a transmission matrix simulation, using a test value for the permittivity. Using the same method we also characterised the linear permittivity of SiO$_2$ and that of a single Ge/Ag bilayer, which were then used to calculate the effective index permittivity of the homogenised stack.

**Nonlinear characterisation.** The material polarisation at the probe wavelength is given by

$$P(\omega_{pr}) = \varepsilon_0 [\varepsilon(\omega_{pr}, I_p) - 1] E(\omega_{pr}),$$

(2)

where $\omega_{pr}$ and $\omega_p$ are the probe and pump frequencies, respectively, $E(\omega_{pr})$ is the probe electric field, and the relative nonlinear permittivity is defined as

$$\varepsilon(\omega_{pr}, I_p) = 1 + \chi^{(1)}(\omega_{pr}) + 6\chi^{(3)}(\omega_{pr}, \omega_p) \frac{I_p}{2n_p^2c^3},$$

(3)

being $n_p$ the real part of the medium refractive index at the pump frequency. It is thus clear that a linear scaling of the permittivity with the pump intensity allows an interpretation in terms of a third-order nonlinearity, where the the $\chi^{(3)}$ tensor can be retrieved by deriving the relative permittivity $\varepsilon(\omega_{pr}, I_p)$ with respect to $I_p$:

$$\chi^{(3)}(\omega_{pr}, \omega_p) = \frac{n_p^2c\partial\varepsilon(\omega_{pr}, I_p)}{3\partial I_p}.$$

(4)

The nonlinear (pumped) permittivity $\varepsilon(\omega_{pr}, I_p)$ has been evaluated from the reflectivity and transmissivity measurements in presence of the pump, for different pump intensities and probe wavelength. Starting from the linear $\varepsilon(\omega_p, 0)$ and exploiting a transfer matrix approach similar to the one used for the linear characterisation, we found the value $\varepsilon(\omega_{pr}, I_p)$ resulting in a $\Delta R/R_0$ and $\Delta T/T_0$ that best matched the experimental data at the given $I_p$. The dual condition on reflectivity and transmissivity allows one to retrieve both the real and imaginary part of $\varepsilon(\omega_{pr}, I_p)$. We then used Eq. (4), identical to Eq. (1) in the main text, to calculate $\chi^{(3)}(\omega_{pr}, \omega_p)$.

**Data availability.** All data relevant to this work may be obtained at doi: 10.17861/37615dc3-8a2d-4242-bdf8-3d06580e3571.

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Author Contributions

R.M.K. performed experiments with assistance from T.R., M.C. and M.P. fabricated and characterised linearly the samples, L.C. and M.P. performed data and theoretical analysis, C.R. and A.C. provided theoretical support, D.F. and A.D.F. directed the project and wrote the manuscript. All authors contributed to scientific discussions and to the manuscript.

Additional Information

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