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Research Article

Domenico Genchi*, Francesca Dodici, Tiziana Cesca and Giovanni Mattei Design of optical Kerr effect in multilayer hyperbolic metamaterials

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Abstract: The design of optical materials in nonlinear devices represents a fundamental step for their optimization and miniaturization, that would significantly contribute to the progress of advanced nanophotonics and quantum technologies. In this work, the effect of geometry and composition of multilayer hyperbolic metamaterials on their third-order nonlinear optical properties, i.e. the optical Kerr effect, is investigated. One figure of merit is provided to be used as a predictive tool to design and best exploit the local intensity enhancement in low-loss metamaterials to be used for various applications in nonlinear nanophotonics.

Keywords: hyperbolic metamaterials; epsilon-near-zero; multilayers; optical Kerr effect; nonlinear optical properties

1 Introduction

In recent years, great interest of scientific research has grown in novel materials whose optical properties can be tailored as desired and actively controlled by external parameters in order to manipulate light at the nanoscale [1]. This is highly relevant for advanced applications in nonlinear nanophotonics, such as ultrafast all-optical switching [2], optical limiting [3] and frequency conversion [4], representing the heart of quantum technologies. One of the main challenges is the design of materials and devices properly featured to achieve an enhanced ultrafast nonlinear response with high efficiency at low-intensity excitation. To

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this scope, predicting, controlling and enhancing the nonlinear optical response becomes essential for the development of efficient and reconfigurable nonlinear optical devices with optimized properties to be used in integrated optical circuits [5].

In this framework, near-zero-index (NZI) materials offer new interesting features that can be used for light control in photonic devices [6]. Indeed, their refractive index reaches a very low value (close to zero), thus producing an increase in the normal field component at the boundary with an isotropic medium $(E_{\text{NZI}} = n_i^2 E_i / n_{\text{NZI}}^2)$, a diverging phase velocity, and a very small phase advance of the propagating wave, i.e. the group velocity tends to zero. These result in a large energy confinement inside the material, and in its ability to support slow light with long interaction lengths [7]. The peculiar properties of NZI materials favour a strong light-matter interaction which is able to trigger enhanced optical nonlinearities owing to the inverse proportionality between the complex nonlinear refractive index and the linear one [8].

Epsilon-near-zero (ENZ) materials are one of the most promising class of non-magnetic NZI media potentially employable as platforms for these applications [9], and several intriguing phenomena useful for their operation and miniaturization have been predicted and observed, such as radiation phase pattern tailoring [10], electromagnetic energy squeezing [11], subwavelength light guiding [12], and time refraction [13]. At the ENZ wavelength ($\lambda_{\rm ENZ}$) of these materials, where the real part of the dielectric permittivity crosses the zero, a strong electric field enhancement is obtained, which is able to induce important ultrafast nonlinear optical responses under light excitation, thus making them very appealing for several nonlinear optical applications [8], [14]. Among these, all-optical switching is one of the most important to fulfill the ON/OFF function needed for ultrafast optical signal processing [15]-[18]. One method to achieve it is based on the optical Kerr effect (OKE), namely the dependence of the nonlinear refractive index and the nonlinear absorption coefficient of a medium on the intensity of the pump field, which permits the modulation of the local refractive index and absorption coefficient on the picosecond and femtosecond time scale [19]-[21].

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The OKE can be significantly enhanced in ENZ media near λ_{ENZ} [22]–[24]. However, their performances are typically limited by their intrinsic losses, that can suppress the NZI effects regardless of how close to zero is the refractive index. Indeed, the ENZ regime typically coincides with the NZI regime in the absence of losses. Differently, with non-negligible losses, the NZI regime is distinguished from the ENZ regime, with the former generally redshifted in the spectrum compared to the latter, and shown to be more beneficial for the efficiency of nonlinear optical effects [25]. Thus, in nonlinear ENZ materials the near-zero permittivity is not the only sufficient condition for the nonlinear response enhancement, but also the intrinsic losses have to be reduced.

Hyperbolic metamaterials (HMs) emerge among the highest-potential ENZ materials. As a matter of fact, unlike plasmonic metals (Au, Ag, Cu, ...) and transparent conductive oxides (ITO, AZO, ...) showing natural ENZ properties at their plasma frequencies respectively fixed in the ultraviolet or infrared ranges of the electromagnetic spectrum, HMs offer a broadband spectral tunability of their λ_{ENZ} at visible wavelengths thanks to the tailorability of their compositional and morphological features. The combination of a metal and a dielectric in a multilayered geometry or their arrangement as metal nanorods in a dielectric matrix allows to obtain an extremely anisotropic uniaxial dielectric tensor whose components are opposite in sign [26]. This gives rise to a topological transition from elliptical to hyperbolic dispersion, and adjustable optical properties, which can be engineered by changing the metal filling fraction and controlled by external parameters [27], [28]. Recently, in the field of nonlinear nanophotonics, enhanced third-order nonlinearities in HMs have been reported [24], [29], and their angle and polarization tunability has been demonstrated [30], [31]. Then, HMs have been proposed as platforms for OKE-based all-optical modulators [15], [20], [32].

Multilayer HMs are particularly attractive for the simplicity of their fabrication through lithography-free nanofabrication methods, and the capability of integrating them within optical devices using current technologies. Moreover, they behave as effective waveguides made up of coupled optical nanocavities where the spatial mode matching between the surface plasmon polaritons (SPPs) supported at the metal-dielectric interfaces gives rise to highly confined plasmonic modes with very large wavevectors and slow light nature that can be excited by free-space light [33]. In the literature, several names are used to call these modes, such as bulk plasmon polaritons [34], volume plasmon polaritons [35], high-k modes [36], Ferrell–Berreman modes [37], Bloch plasmon polaritons [38], and multilayer

plasmons [39]. Due to the heterogeneous denomination, here we prefer to call them with a more general definition of plasmon polaritons (PPs). Such PPs permit to obtain large field enhancement and a huge increase of the optical density of states, so producing resonances with narrow linewidths that can be exploited in subwavelength optics for achieving large optical nonlinearities [31], enhanced photoluminescence [40] and high-sensitivity optical sensing [41].

In the present work, we show how to finely tailor the nonlinear optical properties in multilayer hyperbolic metamaterials by geometry and composition. We demonstrate the enhancement of the optical Kerr effect favoured by PPs at λ_{ENZ} in a spectral range far from the interband transitions of the metal component and where its optical losses are minimum. We introduce one figure of merit to be used for designing ENZ multilayer metamaterials with strong third-order optical nonlinearities, best exploiting the local intensity enhancement generated by their ENZ resonances in different angle and polarization configurations, and to optimize them for nonlinear optical applications.

2 Results and discussion

The optimization of the geometrical and compositional features of HMs represents a crucial step for their implementation in nonlinear photonic devices whose properties and performances have to be customizable and accurately controllable. In this work we engineer HMs by changing the metal filling fraction f_m , which is changed by varying the thickness of the dielectric layers (t_d) while maintaining that of the metal layers constant ($t_m = 15$ nm) (see Section 4). In such a way, λ_{ENZ} is tuned across the optical spectrum [28]. In particular, we want $\lambda_{\rm ENZ}$ beyond the region of the metal interband absorption. For instance, the real part of the parallel component of the effective permittivity (Re[ε_{\parallel}]) calculated by effective medium theory (EMT, see Section 4) for Au/Al₂O₃ HMs is shown in Figure 1(a): λ_{ENZ} is redshifted by decreasing f_m , that is by increasing t_d . The absorptance calculated by EMUstack at normal incidence for Au/Al₂O₃ HMs is depicted in Figure 1(b) as a function of t_d . Up to $t_d = 30$ nm the metamaterial exhibits a wide absorption band between 300 nm and 500 nm produced by the interband transitions of gold [42], and an abrupt change of character at the ENZ wavelength owing to the topological transition from elliptic to hyperbolic dispersion occurring at this wavelength [27], [43]. For $t_d > 40$ nm additional peaks appear, and the optical response can be importantly modulated, even in the interband region. The origin of these peaks stems from the spatial mode matching between the SPPs sustained at the metaldielectric interfaces in the HMs, whose coupling gives rise to



Figure 1: Optical and morphological properties of multilayer hyperbolic metamaterials. (a) Effective permittivity (parallel, real part) and (b) absorptance of Au/Al_2O_3 HMs with different metal filling fraction, varied by changing the thickness of the dielectric layers while maintaining that of the metal layers equal to 15 nm. (c) Cross-sectional SEM image (acquired using backscattered electrons), (d) AFM surface topography, (e) RMS roughness profile of the sample AuAlu16. (f–h) Experimental (open symbols) and simulated (solid lines) optical spectra of the samples AuAlu16 (red) and AgAlu16 (blue). The vertical dashed lines in panels (a–b) and (f–h) indicate the λ_{ENZ} of each sample.

PPs [36]. Indeed, the HMs act like photonic crystals made up of multiple stacked optical nanocavities where a very high electric field confinement occurs [31], [44]. This effect is evidenced by the field distribution in the stack, reported as an example in Figure S1(b) of Supplementary Material for the metamaterial with $t_d = 85$ nm at its λ_{ENZ} (666 nm, red profile). For comparison, the field distribution is also depicted at $\lambda = 500$ nm (green profile), namely close to the gold interband transitions region, where the plasmonic effect is weaker due to metal losses. With sufficiently thin metal layers and thick dielectric layers, a standing wave is formed inside the nanocavities, so that the coupling effect becomes more significant and multiple plasmonic resonances can occur. This effect has been also explained in terms of optical tunneling experienced by the photons travelling within each nanocavity composing the multilayers, which globally behave as gratings so that the PP modes can be excited without special momentum matching techniques [44]-[46]. In Figure 1(b), the increase of the dielectric thickness corresponds to a decrease in the metal filling fraction of the HMs. This lowers the losses and the effective refractive index of the metamaterial, so promoting the SPP coupling inside it and allowing the activation of a larger number of PPs, whose existence is corroborated by the multiple dips appearing in the reflectance spectra plotted in Figure S1(c) of Supplementary Material (connected with the absorptance peaks in Figure 1(b)), which are the distinctive features of PP modes [47]. The properties of these resonances (e.g. their number, spectral distance, and quality factor) depend on the geometrical features of the multilayer (i.e. layers thickness and number of cavities) [44], [46], [47]. The condition for these resonances to be sustained is given by the ratio between the real part and the imaginary part of the metal refractive index, and a critical value of this ratio $(n_m/\kappa_m = 0.2)$ has been found by *Caligiuri* et al. [45], thus defining the so-called "hermiticity limit". The n_m/κ_m ratio for Au is plotted in Figure S1(d) of Supplementary Material. In Aubased systems the cavity resonances can be sustained in the Hermitian region at $\lambda > 550$ nm where $n_m/\kappa_m < 0.2$, and cannot exist in the non-Hermitian interband region ($\lambda < 550$ nm, $n_m/\kappa_m > 0.2$).

Experimentally, we fabricate hyperbolic multilayers corresponding to those considered in Figure 1(a), (b). The constituent materials (gold and alumina) are chosen owing to their high chemical stability. Details about the synthesis and characterization are reported in Section 4. As an example, Figure 1(c) displays the cross-sectional SEM image acquired with backscattered electrons for the 16/85 Au/Al₂O₃ metamaterial with $f_m = 16$ % (AuAlu16): the metal layers (light gray) and dielectric layers (dark gray) are clearly distinguished thus evidencing the four-period structure of the stack. The AFM topography and the RMS roughness (R_{RMS}) profile of the sample are shown in Figure 1(d), (e), revealing a continuous unvoided surface with an average

RMS roughness equal to 0.9 ± 0.1 nm, indicating a good guality of the films growth during the depositions. The spectroscopic ellipsometry (SE) data measured for AuAlu16 are reported in Figure S2 of Supplementary Material, which are used to evaluate the quality of the multilayers in terms of thickness homogeneity, surface roughness and interfaces definition. The morphological parameters extracted by AFM and SE for all the metamaterials fabricated in the present work are tabulated in Table S1 of Supplementary Material. The optical transmittance of AuAlu16 is illustrated in Figure 1(f) (red curve), showing a good agreement between the experimental data and the simulations by EMUstack. The simulated reflectance and absorptance are plotted in Figure 1(g), (h). The existence of PP modes supported by the metamaterials is evidenced by the reflectance dips, corresponding to the transmittance and absorptance peaks [37], [47], [48]. Since the propagation and coupling of SPPs within HMs strongly depend on the thickness of the layers constituting the coupled nanocavities [44], we study the evolution of the optical properties in Au/Al₂O₃ HMs with different geometrical features (see Figure S4 in Supplementary Material). The results show that the transparency window for light propagation through the metamaterial can be broadened by increasing the thickness of the dielectric layers, that is lowering the metal filling fraction, so that a higher number of PP modes are sustained. Moreover, the SPP coupling plays a more important role in metamaterials with low filling fraction, significantly modifying the spectral shape of their optical features, strongly modulated by increasing the thickness of the layers. As shown in Figure S1(d) of Supplementary Material, the Hermitian spectral range where the cavity resonances are sustained can be extended by using Ag due to its lower optical damping and a blueshifted interband region at $\lambda < 350$ nm [45]. Then, we fabricate an Ag/Al₂O₃ HM (named as AgAlu16), equivalent to AuAlu16 (i.e. having $t_m = 16 \text{ nm}, t_d = 85 \text{ nm},$ $f_m =$ 16 %) in order to compare their optical properties and give insights into the role of composition on the plasmonic response of HMs. A very thin (4 nm) alumina capping layer is also deposited on its top to increase the chemical stability of the sample, preventing oxidation of the upper Ag layer without affecting the optical properties of the metamaterial. The SE spectra and a SEM image of the sample are shown in Figure S2(c), (d) of Supplementary Material, while the measured morphological parameters are reported in Table S1(b) of Supplementary Material. A good agreement is obtained between the experimental and simulated transmittance in Figure 1(f), showing a higher number of optical resonances associated with PP modes, whose number

corresponds to the number of optical nanocavities constituting the metamaterial. The ENZ wavelength determined for AgAlu16 is $\lambda_{ENZ} = 632$ nm.

Considering the main points highlighted so far, we introduce a figure of merit (FOM) to be used as a tool for tailoring the geometry and composition of HMs and better take advantage of the contribution given by the ENZ and PP modes in order to obtain enhanced third-order optical nonlinearities, which are enabling properties for nonlinear optical applications, like all-optical switching, optical limiting and mode-locking. The basic idea of this FOM is to maximize the local intensity promoted by the confinement of PPs and minimize the optical damping given by the metallic component or by the reflection losses in order to favour the nonlinear optical response in a high-transmission metamaterial. In such a way, the FOM allows to find the optimal trade-off between these effects, and devices with high nonlinear efficiency and high output signal can be designed. The FOM is defined as:

$$FOM = \frac{T}{R} \frac{\phi_m}{Im[\varepsilon_m]}$$
(1)

Here, T and R are, respectively, the transmittance and reflectance of the metamaterial, $\text{Im}[\epsilon_m]$ is the imaginary part of the permittivity of the metal constituent, and ϕ_m is an effective filling fraction (see Section 4) which accounts for the local intensity enhancement in the metamaterials and plays the main role in controlling their optical nonlinearities, as demonstrated in Ref. [31]. It is worth to underline that all the parameters included in the FOM are wavelengthdependent. Besides, *T* , *R* and ϕ_m are angle-dependent. Consequently, the FOM depends on both wavelength and angle, and thus it can be used for any spectral and angular configuration. Hereafter we make use of the FOM to study the effect of the dielectric thickness, metal thickness, and composition on the third-order nonlinear optical properties of HMs. However, it is worth to point out that the FOM includes important parameters that can be useful for a broader range of applications, such as in optical sensing [41] and radiative decay engineering [40].

Figure 2(a) shows the FOM calculated at normal incidence in the visible spectrum for a four-period Au/Al₂O₃ HM with 15 nm metal thickness and varying dielectric thickness (10–200 nm). The white dashed line and green solid line are the locus of points of the ENZ wavelengths calculated respectively by EMT [43] and inverse scattering matrix method (ISMM) [49] for each geometrical combination. In the contour plot, three claw-like bands appear for multilayers with $t_d > 30$ nm above $\lambda = 500$ nm. These correspond to PP modes existing in the Hermitian region of the spectrum (see Figure S1(d) of Supplementary Material)



Figure 2: Contour plots of (a, d) the FOM, (b, e) nonlinear absorption coefficient β , and (c, f) nonlinear refractive index n_2 calculated for HMs with 15-nm-thick metal layers and different dielectric layers thickness, arranged in four-period stacks. Panels (a – c) refer to Au/Al₂O₃ HMs, whereas panels (d–f) to Ag/Al₂O₃ HMs. The FOM (a, d) is calculated by Eq. (1), whereas β and n_2 are computed by our model using the effective filling fraction [31]. The dashed green curves correspond to the ENZ wavelengths extracted by EMT, whereas the solid green curves represent those extracted by ISMM. The stars mark a metamaterial structure made up of 15 nm metal layers and 85 nm metal layers.

where ENZ resonances can be sustained by the coupled nanocavities in the HMs [45]. In particular, the rightmost band corresponds to PP modes occurring near the ENZ wavelength of the metamaterials. The PP band and the ENZ curves intersect at a point corresponding to a structure having 85-nm-thick dielectric layers (AuAlu16, marked with the star). This crossing can represent the limit of validity of EMT against ISMM for the considered four-period multilayer structure, and it can be used as a benchmark for multilayer periodic structures possessing the optical properties of coupled nano-waveguides but still describable as ENZ metamaterials in a quite good approximation. Indeed, the values of $\lambda_{\rm ENZ}$ calculated by the two methods (white = EMT, green = ISMM) coincide within a 1 % error up to $t_d = 85$ nm, whereas the two curves start to diverge for larger dielectric thickness. The percent deviation between $\lambda_{\text{ENZ}}^{\text{ISMM}}$ and $\lambda_{\text{ENZ}}^{\text{EMT}}$, calculated as $\Delta \lambda_{\text{ENZ}}(\%) = [\lambda_{\text{ENZ}}^{\text{ISMM}} - \lambda_{\text{ENZ}}^{\text{EMT}}] / \lambda_{\text{ENZ}}^{\text{ISMM}}$ for $t_m = 15$ nm as a function of t_d is shown in Figure S5(a) of Supplementary Material. Besides, a comparison between the effective permittivity calculated by EMT and ISMM for AuAlu16 is reported in Figure S6(a) of Supplementary Material.

The highly confined PPs produce a local intensity enhancement (represented by ϕ_m in Figure S7(a), (b) of Supplementary Material) that potentially favours the nonlinear response of the metamaterials. Figure 2(b), (c) show the nonlinear absorption coefficient β and the nonlinear refractive index n₂ calculated by following the approach described in our previous work [31] (see Section 4). A strong nonlinear response is obtained around $\lambda = 500$ nm due to the interband contribution of gold, producing a large peak of β and a sign inversion from negative to positive n_2 . A change of sign is also obtained in the β coefficient at $\lambda < \beta$ 500 nm. The sign reversal of both the nonlinear parameters corresponds to a sign variation of the third-order susceptibility of gold ($\chi^{(3)}_{Au}$) in the interband region [50]. The predicted nonlinear response is maximum when λ_{ENZ} falls in the interband region. However, non-negligible values of the nonlinear parameters are also obtained along the λ_{ENZ} curve. By increasing the thickness of the dielectric layers in the HMs, the ENZ wavelength is spectrally redshifted as a result of the metal filling fraction decrease, and so does the nonlinear optical response in the ENZ regime, favoured by the strong field enhancement produced by the plasmonic modes associated with the first branch of the FOM. In such a way, non-null optical nonlinearities can be obtained even far from the metal interband absorption region exploiting the plasmonic effect near λ_{ENZ} . In particular, positive β and n_2 are predicted in the Au-based HMs taken as an example, and they tend to diminish in absolute value by moving λ_{ENZ} towards the NIR region owing to a raise of the optical damping of gold (i.e. its $Im[\epsilon]$ increases, see Figure S3(b) in Supplementary Material) in a spectral range where also $\chi^{(3)}_{Au}$ is low [50]. This concurrently underlines the relevance of low losses in order to enhance the nonlinear optical response in a nonlinear material, and the intrinsic contribution given by the metallic component. It is worth to note that not all the PP modes represented by the FOM bands correspond to an enhanced nonlinear optical response. Indeed, the nonlinear parameters predicted in correspondence of the other PP branches (not close to λ_{EN7}) are quite low. This demonstrates that the ENZ regime is essential to significantly increase the nonlinear response favoured by the plasmonic resonance because it enables a higher local intensity enhancement, as demonstrated by the intense peak of ϕ_m in Figure S7(a), (b) of Supplementary Material.

The details of the FOM features depend on the compositional properties of the HMs. As a matter of fact, the field confinement within the stack can be varied by changing the dielectric contrast between the metal and dielectric constituents. Besides, the metal mostly affects the optical damping and the capability to support ENZ resonances depending on the spectral position of the interband absorption. The FOM can be increased with Ag-based HMs, in which the optical losses are largely dropped because the interband absorption of silver occurs in the near-UV spectral range, thus also widening the Hermitian region supporting ENZ resonances (see Figure S1(d) of Supplementary Material). For example, as shown in Figure 2(d), in the case of the Ag/Al₂O₂ HM a 3-fold increase of the FOM and a larger number of PP modes exist in a wider spectral region compared to Au-based HM. The greater prominence of the PP bands in the Ag/Al_2O_3 stacks is related to the better decoupling of these modes from the interband region. As a consequence, the nonlinear optical response in the visible range is predicted to be larger than in Au-based HM. Indeed, as shown in Figure 2(e), (f), the nonlinear parameters β and n_2 are predicted to be 10 times higher: the highest optical nonlinearities still occur near λ_{ENZ} , where eta changes its sign from positive to negative and n_2 exhibits a maximum. Even in this case the λ_{ENZ} curves calculated by EMT (dashed green) and ISMM (solid green) overlap up to $t_d = 85$ nm ($\Delta \lambda_{ENZ} = \pm 2$ %, see Figure S5(a) in Supplementary Material), and they intersect the rightmost resonance band of the FOM with a 15/85 nm multilayer (AgAlu16, marked with the star). The effective permittivity of AgAlu16 calculated by EMT and ISMM is shown in Figure S6(b) of Supplementary Material.

It is worth to stress that the presented calculations predict the highest third-order nonlinearities near the ENZ wavelength, and not at the NZI condition where the metamaterial has the minimum refractive index (n), that would occur at $\lambda > \lambda_{ENZ}$ as shown in other works [6], [25]. For example, according to EMT, AuAlu16 and AgAlu16 respectively exhibit the lowest *n* at 837 nm and 830 nm, as shown in Figure S8(a), (b) of Supplementary Material where the effective optical functions of the metamaterials are plotted. Therefore, a high nonlinear response might be expected near these wavelengths due to NZI effects. Nevertheless, in this spectral region the metamaterials exhibit high values of their extinction coefficient κ and reflectance, so that losses play an important role in the definition of the NZI regime, and limit the nonlinear response. Thus, since κ cannot be neglected in the case of hyperbolic metamaterials with effective parameters, we considered the modulus of the complex effective refractive index $(|\tilde{n}| = \sqrt{n^2 + \kappa^2})$ to identify the NZI condition $(|\tilde{n}| < 1)$, as done in Ref. [51]. This is plotted for AuAlu16 and AgAlu16 in Figure S8(c) of Supplementary Material, where the curves show that the minimum value of $|\tilde{n}|$ coincides with the ENZ wavelength. This is consistent with the prediction of the highest Kerr nonlinear response in the ENZ regime, as also stated in refs. [14], [24], [30].

The loss properties of the metamaterials and the ability of SPPs to propagate and couple at their inside drastically depend on the thickness of the metal layers [44]. For a comprehensive investigation, we study the evolution of the FOM and the nonlinear parameters as a function of the thickness of the metal layers in the metamaterials. The contour plots in Figure 3 display the spectrally-resolved FOM (panels (a, d)) and nonlinear OKE parameters β (panels (b, e)) and n_2 (panels (c, f)) as a function of t_m in four-period metamaterials with fixed thickness of the dielectric layers ($t_d = 85$ nm). As described above, this dielectric thickness corresponds to the value at which the establishment of λ_{ENZ} by EMT is at its limit (as seen in Figure 2), and $\lambda_{\rm ENZ}$ matches a PP mode (see Figure 1(b)). The plots in Figure 3(a)–(c) correspond to Au/Al₂O₃ multilayers. The FOM contour plot in Figure 3(a) shows the presence of local maxima at $\lambda > 500$ nm (namely over the gold interband absorption region) related to the SPP modes coupling for $t_m < 30$ nm. The bands become progressively less intense and less separated as t_m increases, and completely disappear with $t_m > 30$ nm. The decrease in the absolute value of the FOM is due to the fact that, by increasing t_m with fixed t_d , the metamaterial has a gradually larger metal filling fraction, thus leading to larger absorption and reflection. Furthermore, thicker metal layers provide a smaller penetration depth bringing to the breakdown of long-range SPPs [52], then to a low T/R ratio and low output signal, thus making the metamaterials of less practical use in applications like all-optical switching where high



Figure 3: Contour plots of (a, d) the FOM, (b, e) nonlinear absorption coefficient β , and (c, f) nonlinear refractive index n_2 calculated for HMs with 85-nm-thick dielectric layers and different metal layers thickness, arranged in four-period stacks. Panels (a–c) refer to Au/Al₂O₃ HMs, whereas panels (d–f) to Ag/Al₂O₃ HMs. The FOM (a, d) is calculated by Eq. (1), whereas β and n_2 are computed by our model using the effective filling fraction [31]. The dashed green curves correspond to the ENZ wavelengths extracted by EMT, whereas the solid green curves represent those extracted by ISMM. The horizontal dashed gray line indicate $t_m = 15$ nm.

transmitted signals are needed. On the other hand, the decrease in the bands separation is due to the fact that, when t_m is increased, the coupling between the cavities is weakened by the increasingly strong decay of the field in the metal layers. As the coupling decreases, so does the splitting between the cavity modes. The highest FOM values are predicted for the band around $\lambda = 650 \text{ nm}$ occurring near the minimum of Im[$\epsilon_{
m Au}$]. Furthermore, it is worth to notice that the FOM peak at larger wavelengths is located in proximity of λ_{ENZ} , which corresponds to a higher local field enhancement in the metamaterial as demonstrated by the strongest peak of ϕ_m in Figure S7(c), (d) of Supplementary Material, and the spectral position of the FOM peak follows the blueshift of λ_{ENZ} with increasing t_m . The curves of the ENZ wavelengths calculated by EMT (dashed green) and ISMM (solid green) nearly follows the same trend with a small divergence (within 2%, see Figure S5(b) in Supplementary Material) over $t_m = 15 \text{ nm}$ (marked with the horizontal dashed gray line). As already discussed above, the ENZ condition is essential to enhance the OKE in the metamaterials far from the spectral region associated with the interband contribution of the metal component. This is evidenced in Figure 3(b), (c), where we plot the calculated nonlinear parameters as a function

of the metal layers thickness in Au/Al₂O₃ metamaterials with $t_d = 85$ nm. The larger nonlinear response is predicted around $\lambda=500~\text{nm}$ owing to the high $\chi^{(3)}_{\rm Au}$ deriving from the contribution of interband transitions [50]. $\chi^{(3)}_{Au}$ exhibits a sign inversion in the same spectral region, and so do the nonlinear parameters [50]. The nonlinear response descending from this contribution clearly tends to grow proportionally to t_m (that is f_m) due to an increasing metallic character of the metamaterial. The nonlinear response becomes very similar to that of a single Au thin film since the incident light has significant interaction with the first layer only [31], [53]. However, with thin metal layers (and so low metal filling fraction) the ENZ regime allows to obtain a peak of β and n_2 at longer wavelengths (see insets in Figure 3(b), (c)) due to the field enhancement at λ_{ENZ} . As the metal thickness t_m is increased (and so f_m), the effective behavior of the metamaterial becomes progressively more metallic [54], thus $\lambda_{\rm ENZ}$ and the associated resonance blueshift. The peak of the nonlinear parameters follows the same trend until it merges with the peak related to the interband contribution, which dominates at lower wavelengths. However, with large t_m the high intrinsic losses of the metal and the poor transmittance of the metamaterials make them unpractical for transmission-based all-optical modulators, indeed the FOM

is very low. This evidences the importance of tailoring a metamaterial with thin metal layers and low metal filling fraction.

The FOM and the nonlinear parameters calculated for Ag/Al_2O_3 metamaterials are shown in Figure 3(d)–(f) for comparison. As discussed above, the resonances observed in the FOM are sharper and higher in absolute value compared to those observed for the Au/Al_2O_3 stacks, as a consequence of the better decoupling of the PP modes from the interband region of Ag and a larger Hermitian spectral region where they are supported (see Figure S1(d) of Supplementary Material). The bands become progressively less intense and less separated at higher values of t_m due to the increasing metallic character of the metamaterials and their larger optical losses. The resonance bands related to the PP modes exist for $t_m < 40$ nm, and the one around $\lambda = 500$ nm exhibits the highest FOM values in a region where $\mathrm{Im}[arepsilon_{\mathrm{Ag}}]$ is very low. The modes at larger wavelengths (around $\lambda = 650$ nm) are associated with the ENZ regime of the metamaterial, and the band follows the blueshift of λ_{ENZ} with increasing t_m . Even in this case the ENZ curves calculated by EMT (dashed green) and ISMM (solid green) approximately overlap ($\Delta \lambda_{ENZ} = 2$ %) up to $t_m = 15$ nm (marked with the horizontal dashed gray line), over which the divergence starts to become more significant, even larger than in the case of Au-based metamaterials (see Figure S5 in Supplementary Material). By comparing the contour plots of the OKE parameters with those of the FOM, local maxima of the nonlinear parameters are found at the same wavelengths where the FOM bands related to the PP modes appear. Compared to the case of Au/Al₂O₂ metamaterials, these peaks are more distinct due to their better decoupling from the interband absorption region of Ag, which is shifted below $\lambda = 300$ nm. Together with the lower optical damping of Ag, this leads to a stronger nonlinear response compared to the Au-based metamaterials. The highest nonlinear parameters are obtained in correspondence of the ENZ wavelength. The peak band of both β and n_2 follows the same trend of the λ_{ENZ} curve, thus blueshifting with growing t_m due to the increasing metal filling fraction.

Considering the insights emerged from the use of the FOM, and in order to stress its validity for the design of optimized nonlinear multilayer metamaterials, we experimentally investigate the nonlinear optical response of the produced four-period Au/Al₂O₃ and Ag/Al₂O₃ ENZ metamaterials with 15/85 nm layers structure in the visible range around their λ_{ENZ} by the z-scan technique [55] (see Section 4). The measurements are performed at an average intensity of 350 MW/cm², chosen to be sufficiently high to activate the nonlinear response of the HM while being

below the damage threshold of Au and Ag. In Figure 4 we plot (full symbols) the nonlinear refractive index n_2 (panel a) and nonlinear absorption coefficient β (panel b) measured at normal incidence for AuAlu16 (red) and AgAlu16 (blue). The error bars are determined on the basis of repeated measurements, as well as considering the variability of the parameters in the function used for fitting the experimental z-scan curves with good agreement. In agreement with the simulations shown in Figures 2 and 3, AuAlu16 exhibits a peak of positive n_2 and β around its $\lambda_{\rm FNZ} = 666$ nm, respectively associated with self-focusing (SF) and reverse saturable absorption (RSA). The open aperture (OA) and closed aperture (CA) z-scan curves measured for AuAlu16 at $\lambda_{FNZ} = 666$ nm are reported in Figure S9(a), (b) of Supplementary Material as examples, from which β = 4 × 10⁻⁵ cm/W and n_2 = 3 × 10⁻¹⁰ cm²/W are extracted. The largest nonlinear response (positive β and negative n_2) is observed at $\lambda < 550$ nm (red area), namely in



Figure 4: Nonlinear refractive index (a) and nonlinear absorption coefficient (b) of AuAlu16 (red) and AgAlu16 (blue). The experimental data (full symbols) are compared to the curves calculated by our model using the effective filling fraction [31] (solid lines). The measurements are taken at an intensity of 350 MW/cm². The vertical dashed lines indicate the λ_{ENZ} of each sample, namely 666 nm for AuAlu16 and 632 nm for AgAlu16. The red area indicates the interband absorption region of gold.

the region of gold interband absorption, as a consequence of the large $\chi^{(3)}_{Au}$ [31], [50]. In Figure S9(c), (d) of Supplementary Material we report the *z*-scans taken at $\lambda = 500$ nm, where the highest values of the nonlinear parameters for the metamaterial are extracted ($\beta = 5.5 \times 10^{-4}$ cm/W and $n_2 = -6 \times 10^{-10} \text{ cm}^2/\text{W}$), in comparison to the corresponding z-scans taken from a 15-nm-thick Au film. The experimental data in Figure 4 are well described by the solid curves calculated by our model using the effective filling fraction [31]. For λ < 550 nm saturation effects in the nonlinear absorption process have to be considered, which are taken into account in the fitting procedure of the z-scan curves by using an intensity-dependent expression for the nonlinear absorption coefficient that becomes a saturated parameter according to $\beta_s(I) = \beta/(1 + I/I_s)$, where β is the unsaturated coefficient and I_S is the saturation intensity [31]. A saturation intensity of $I_s = 200 \text{ MW/cm}^2$ is estimated. No nonlinear response is observed in the 85 nm Al₂O₃ reference film. On the other hand, AgAlu16 exhibits SF ($n_2 > 0$) across all the explored spectrum with a peak in proximity of λ_{ENZ} (632 nm), where correspondingly a transition from saturable absorption (SA, $\beta > 0$) to RSA ($\beta < 0$) occurs. Near $\lambda_{\rm ENZ}$ the nonlinear refractive index has a peak 10 times higher than in AuAlu16. This is due to the concomitant effect of: (i) a lower optical damping of silver (Im[ε_{Ag}] = 1.0) compared to gold (Im[ϵ_{Au}] = 1.8), (ii) a more efficient propagation of SPPs in silver than in gold [56], and (iii) a larger index contrast between silver and alumina ($n_{Al_2O_3}/n_{Ag} = 11.7$) compared to gold and alumina ($n_{Al_2O_3}/n_{Au} = 6.6$). Moreover, the nonlinear response of AgAlu16 is enhanced as compared to that of a 15 nm silver film used as a reference. Indeed, as shown in Figure S6(e), (f) of Supplementary Material, no nonlinear response is observed in the Ag film i.e. a flat signal for the OA and CA normalized transmittance curves is measured. Near $\lambda_{\rm ENZ}$ AgAlu16 exhibits $\beta = -4 \times 10^{-4}$ cm/W and $n_2 = 2 \times 10^{-9} \text{ cm}^2/\text{W}$. For the sake of completeness, in Figure S10 of Supplementary Material we report the effective $\chi^{(3)}$ (parallel and perpendicular components) calculated by the model using the effective filling fraction [31] for the samples AuAlu16 and AgAlu16, from which the nonlinear parameters n_2 and β simulated in Figure 4 are extracted. It is worth to stress again that, consistently with the results demonstrated in Refs. [14], [24], [30], with both the metamaterials characterized experimentally a strong Kerr nonlinear response is observed near the ENZ wavelength, which coincides with the condition of minimum $|\tilde{n}|$ (see Figure S8(c) in Supplementary Material), namely the NZI point according to Ref. [51]. Owing to high reflectance, with our transmission-based z-scan setup we could not investigate the nonlinear parameters at λ >

800 nm where the metamaterials exhibit the minimum n (see Figure S8(a), (b) in Supplementary Material), that is at the NZI condition defined as in Ref. [25]. However, considering the calculations of the nonlinear parameters in Figures 2 and 3, and those of the effective $\chi^{(3)}$ in Figure S10 of Supplementary Material, we may expect null nonlinearities in this spectral region due to high losses (high κ) that importantly limit the nonlinear response.

The third-order nonlinear optical properties of ENZ metamaterials has been recently demonstrated to be tunable by changing the polarization and the incidence angle of the exciting light [30], [31]. In particular, in our previous work [31] we demonstrated that in hyperbolic multilayers this effect is due to the modulation of the local field at their inside as a consequence of a change of interaction between the SPPs in the stacked nanocavities with changing angle, an effect being especially evident in the ENZ region far from the metal interband transitions region. Hence, we calculate the FOM for different incidence angles with TE and TM polarizations at $\lambda_{\rm ENZ}$ (extracted by ISMM), where the largest nonlinear response is expected. We map the computed FOM as a function of the layers thickness in order to evaluate the potential metamaterial geometries giving enhanced optical nonlinearities at their ENZ wavelength. For example, the FOM contour plots obtained for four-periods Au/Al₂O₃ HMs are illustrated in Figure 5 for the following configurations: (a) normal incidence, (b) 60° with TE polarization, and (c) 60° with TM polarization. These contour plots can be used as a control box for choosing multilayer configurations with optimized nonlinear optical properties at λ_{ENZ} . The stars reported with their labels indicate the samples fabricated in the present work. The labels next to the stars are the n_2 values measured experimentally for the considered multilayers. In particular, the yellow and green dashed lines respectively indicate HMs with $f_m = 16\%$ and $f_m = 33\%$, having the same $\lambda_{\rm ENZ}$, respectively 666 nm and 543 nm. The data show that at each angle and polarization configuration, the higher is the FOM, the higher is the n_2 . The trend of n_2 for these two series of HMs is also shown in the graphs of Figure S11 in Supplementary Material. The measured n_2 and β with their uncertainties for the HMs of this work are reported in Table S2 of Supplementary Material. However, it is worth to distinguish two regions in the countour plots of Figure 5, separated by the white dotted line (corresponding to HMs with $f_m \sim$ 25%). The HMs above this line have λ_{ENZ} far from the spectral region of gold interband absorption. Instead, for the HMs below the line, λ_{ENZ} falls inside this region, where the interband contribution and the large $\chi^{(3)}_{Au}$ dominate the nonlinear response. As a result, the nonlinear parameters are higher in HMs



Figure 5: Contour plots of the FOM calculated by Eq. (1) at (a) normal incidence, (b) 60° (TE polarization) and (c) 60° (TM polarization) at the ENZ wavelengths (extracted by ISMM) of four-period Au/Al₂O₃ HMs with variable thickness layers. The stars indicate the samples fabricated in the present work, and the labels indicate their n_2 (10⁻¹⁰ cm²/W) measured experimentally. The white dotted line separates the geometries having the ENZ regime far from the interband absorption of gold ($\lambda_{\rm ENZ} > 550$ nm, above the line), and close to it ($\lambda_{\rm ENZ} < 550$ nm, below the line). The yellow dashed line and the green dashed lines respectively are the series of HMs with $f_m = 16\%$ ($\lambda_{\rm ENZ} = 666$ nm) and $f_m = 33\%$ ($\lambda_{\rm ENZ} = 543$ nm).

with geometries below the white dotted line. Moreover, it is worth to note that comparable values of the nonlinear parameters are observed at obligue incidence with both TE and TM polarizations. A higher nonlinear response would be generally expected with TM polarization due to a more favourable excitation and coupling of PPs. Nevertheless, due to an interplay between TE and TM modes [33] and since the layers thickness importantly affects the subwavelength resonant behaviour within the metamaterial [45], the nonlinear effects can become important even with TE polarization due to the field enhancement produced by the cavity modes. This is reflected in the effective filling fraction representing the field enhancement generated by the plasmonic coupling inside the metamaterial, which has quite high values in all the configurations [31]. However, it results more stable with TM polarization, while it undergoes more important variations spectrally and in absolute values with TE polarization, thus producing an important variation of the nonlinear parameters, too [31]. Finally, the obtained results show HMs with very thin layers (e.g. $t_m = 10 \text{ nm}$ and $t_d = 20 \text{ nm}$) to have the best properties at all angles and polarizations. This can be explained as a consequence of quantum size effects starting to play an important role in ultrathin Au layers with thickness ≤ 10 nm, where a larger field confinement can be achieved with concomitant lower damping, so giving rise to stronger plasmonic resonances [57]-[59]. The increase in the nonlinear response of metamaterials with ultrathin metal layers is also due to the inverse proportionality of the $\chi^{(3)}_{Au}$ with the thickness of the Au film [52], [59]. These indications might be relevant for future nonlinear optics studies with ultrathin HMs.

3 Conclusions

We show how to finely tailor the nonlinear optical properties in multilayer hyperbolic metamaterials by changing their structural and compositional parameters. We evidence that tailoring the ENZ regime in a spectral range where plasmon polariton modes are supported and the optical damping of the metal component approaches its minimum (i.e. far from the interband absorption region) gives rise to high Kerr-like optical nonlinearities, which can be further enhanced in low-loss metamaterials with high index contrast between their metal and dielectric constituents. We introduce one figure of merit to optimize the geometry and composition of multilayer metamaterials, and take full advantage of the local intensity enhancement favoured by the plasmonic effects in the ENZ regime together with high transmission and low losses. The figure of merit provides a predictive tool for the design of ENZ multilayer metamaterials with high Kerr nonlinear optical response to be used at different angles and polarizations, which are enabling for the development of nonlinear optical devices, such as all-optical switches, optical limiters and passive mode-lockers. The outcomes suggest that further studies could be done in hyperbolic metamaterials with ultrathin layers, potentially having boosted nonlinear optical properties owing to quantum size effects. The features included in the FOM can be also useful for a broader range of applications, like optical sensing and radiative decay engineering.

4 Methods

4.1 Numerical methods

The complex anisotropic effective permittivity of multilayer hyperbolic metamaterials is calculated by effective medium theory (EMT) [43] and inverse scattering matrix method (ISMM) [49]. The values of λ_{ENZ} for the considered metamaterials are extracted by the zero-crossing of $\text{Re}[\varepsilon_{\parallel}]$, where $\varepsilon_{\parallel} = f_m \varepsilon_m + [1 - f_m] \varepsilon_d$ is the parallel component of the effective permittivity. Here f_m is the geometrical metal filling fraction calculated as $f_m = t_m/(t_m + t_d)$, where t_m and t_d are the thickness of the metal and dielectric layers, respectively.

The linear optical properties of the metamaterials, namely their transmittance (*T*), reflectance (*R*) and absorptance (A = 1 - R - T), and the electromagnetic field distribution within them is calculated by the scattering matrix method (SMM) via EMUstack [60].

The nonlinear refractive index and the nonlinear absorption coefficient are calculated as $n_2 = \text{Re}[\tilde{n}_2]$ and $\beta =$ $(4\pi/\lambda)$ Im $[\tilde{n}_2]$ respectively, where \tilde{n}_2 is the complex nonlinear refractive index of the metamaterial, defined as $\tilde{n}_2 =$ $[3/(4c\varepsilon_0\tilde{n}_0n_0)]\chi_{\rm eff}^{(3)}$, with c being the speed of light in vacuum, ε_0 the vacuum permittivity, \tilde{n}_0 the complex linear refractive index of the metamaterial, and n_0 its real part [24]. Owing to the optical anisotropy of the metamaterial, the effective third-order susceptibility $\chi^{(3)}_{\mathrm{eff}}$ is angle-dependent, and so is \tilde{n}_2 [30], [31]. The complex nonlinear refractive index is calculated following the approach proposed in our previous work [31] using an effective metal filling fraction that represents the local intensity enhancement within the metamaterial. The effective filling fraction is defined as $\phi_m(\lambda, \theta) =$ $f_m[\zeta_{\rm MM}(\lambda,\theta)/\zeta_{\rm film}(\lambda,\theta)]$ with $\zeta_{\rm MM}$ being the spatial average of the electric field square modulus $(|E|^2)$ in the metallic layers of the metamaterial, and $\zeta_{\rm film}$ being the same quantity integrated over a reference metal film corresponding to the metal layer in the structural unit of the metamaterial. These quantities are computed by simulating the electric field profile inside the multilayer and the single film with EMUstack [60].

All the parameters are calculated in the VIS-NIR wavelength range ($\lambda = 300-1000$ nm). A sketch of the metamaterial structure implemented for the numerical simulation is depicted in Figure S1(a) of Supplementary Material. The stack is made up of metal and dielectric layers arranged in a four-period structure with the metal on the top of each period. One period corresponds to one metal-dielectric bilayer, that is the unit cell of the metamaterial, which is repeated along the *z*-axis. The number of periods is chosen since it corresponds to the minimum value for the EMT applicability to be valid [61]. The semi-infinite half-spaces surrounding the stack are air and substrate (glass).

4.2 Samples synthesis and characterization

The synthesis of Au/Al₂O₃ and Ag/Al₂O₃ multilayer hyperbolic metamaterials is carried out by magnetron sputtering depositions. Glass substrates are cleaned beforehand in an acidic piranha solution (1:3 H₂O₂:H₂SO₄) at 80 °C for 1 h and then rinsed in ultra-pure Milli-Q water. The depositions are performed in an inert Ar atmosphere ($p = 5 \times 10^{-3}$ mbar) using a DC source for the metallic targets and a RF one for the dielectric ones. The targets are equipped with mechanical shutters to allow the alternate deposition of the different materials. The sample holder is kept in continuous rotation during the process to promote homogeneity and thickness uniformity of the layers. Thin films of Au, Ag and Al₂O₃ with the same thickness of the metal and oxide layers in the structural unit (one bilayer) of the metamaterials are also deposited as reference samples.

The structure of the produced multilayers is checked in cross-section with a field-emission scanning electron microscopy (SEM) using a *Zeiss Sigma HD FE-SEM*. The surface roughness and the total thickness are evaluated by atomic force microscopy (AFM, *NT-MDT Solver Pro*). The layers thickness is determined by variable-angle spectroscopic ellipsometry (SE, *J.A. Wollam V-VASE*). Ellipsometric measurements on the reference samples are also carried out to measure the dielectric functions of the constituent materials in the multilayers. The transmittance of the samples is measured with a UV-VIS spectrophotometer (*JASCO V670*).

4.3 Nonlinear optical characterization

The z-scan technique [55] is used to measure the OKE parameters of the samples, i.e. the nonlinear refractive index n_2 and the nonlinear absorption coefficient β . The third harmonic ($\lambda = 355$ nm) of a mode-locked Nd:YAG laser

(Leopard, by Continuum) emitting 18 ps pulses with a repetition rate of 10 Hz is used as the pump source. An optical parametric amplifier (TOPAS, by Light Conversion) is employed to spectrally tune the laser beam in the wavelength range 420–2400 nm. Optical density (OD) filters are used to vary the beam intensity. The beam is spatially filtered to obtain a TEM₀₀ Gaussian mode. The beam is focused by a 200 nm converging lens onto the sample, which is mounted on a motorized stage and translated along the focal axis of the lens. The beam waist at the lens focus is $w_0 = 19 \,\mu\text{m}$ at $\lambda = 600 \,\text{nm}$ measured by the knife-edge method [62]. A beam splitter is positioned at the end of the motorized stage to send the light transmitted by the sample to two Si photodetectors, allowing simultaneous open aperture (OA) and closed aperture (CA) measurements. The CA detector is preceded by an iris diaphragm with linear transmittance S = 0.3 to make the measurement sensitive to phase distortions. Another beam splitter is positioned before the focusing lens to send half of the input beam to another Si-photodetector, to be used as a reference. This allows the normalization of the CA and OA signals to the reference one, compensating for fluctuations of the input laser intensity. To further minimize signal noise, each point of the scans is collected by averaging at least 150 laser pulses. The measurements are taken at intensities around $I = 350 \text{ MW/cm}^2$. To account for the fraction of intensity lost by reflection, we estimate $I = (1 - R)I_0$ as the intensity actually entering the sample and causing its nonlinear response, where I_0 is the intensity of the laser beam at the focus of the lens, and R is the reflectance of the sample at each considered wavelength. The OKE parameters β and n_2 are evaluated by fitting the normalized transmittance curves from the OA and CA lines (T_{OA} and T_{CA}), respectively, using the method described in Ref. [31]. Z-scan measurements are also performed on the reference samples in order to evaluate the nonlinear response of the individual components of the multilayers.

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TC and GM; supervision – TC and GM; project administration – TC and GM. All authors have accepted responsibility for the entire content of this manuscript and approved its submission.

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Ethical approval: The conducted research is not related to either human or animals use.

Data availability: The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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