Independent Amplitude Control of Arbitrary Orthogonal States

of Polarization via Dielectric Metasurfaces

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Abstract

Exquisite polarization control using optical metasurfaces has attracted considerable attention thanks to their ability to manipulate multichannel independent wavefronts with subwavelength resolution. Here we present a new class of metasurface polarization optics, which enables imposition of two arbitrary and independent amplitude profiles on any pair of orthogonal states of polarization. The implementation method involves a polarizationdependent interference mechanism achieved by constructing a metasurface composed of an array of nanoscale birefringent waveplates. Based on this principle, we experimentally demonstrate chiral grayscale metasurface and chiral shadow rendering of structured light. These results illustrate a general approach interlinking amplitude profiles and orthogonal states of polarization and expands the scope of metasurface polarization shaping optics.

An optical metasurface, composed of two-dimensional (2D) arrays of subwavelength meta-atoms, is a compact and planar nanophotonic platform being utilized for manipulating the various fundamental properties of an incoming electromagnetic wave [1-6]. Depending on the meta-atom design, metasurfaces are able to modify the phase profile, polarization ellipse, scattering angular, intensity distribution of incident light wave. One of the attractive features that distinguishes them from conventional optics is their ability to simultaneously perform multiple different optical operations since meta-atoms can be designed to respond differently to the wavelength [7-10], polarization [11-14], and/or the illumination angle of incident light [15]. For example, metasurfacebased polarization optics has recently attracted much attention as a breakthrough concept for engineering multichannel independent wavefront encoding [16-27]. The implementation of this function requires the individual meta-atom to behave as a waveplate-like birefringent element. Typically, there are two wavefront regulation mechanisms in a polarization-dependent dielectric metasurface design—propagation phase and geometric (or Pancharatnam-Berry, PB) phase [28-29]. Recently, a strategy that combines both geometric phase and propagation phase is proposed to control independent phase profiles using a pair of orthogonal polarizations [21-27]. This exciting capability allows for a new class of polarization-dependent switchable photonic functionalities, such as chiral hologram [21], transformation of orbital angular momentum [22], manipulation of accelerating light beams [23], polarization imaging [24-26], and mode-selective waveguide couplers [27].

However, up until now, much of the effort in metasurface optics has primarily focused on establishing the relationship between polarization and phase profile. Usually it is expected that the amplitude profile of an incident polarized electromagnetic wave is maintained upon transmission through a metasurface. As a result, such metasurfaces exhibit high transmission efficiency, but they are limited to only manipulate the phase profile but not the amplitude profile of incident light [16, 18, 30-31]. Although the polarization conversion efficiency of individual meta-atom can be utilized to control the amplitude for a specific polarization state, these metasurfaces require additional free-space optical elements to fully perform the desired amplitude control [32, 33]. Furthermore, these devices exhibit the same amplitude response when the chirality of incident circularly polarized light is flipped, and cannot achieve the completely decoupled output amplitudes for two arbitrary, orthogonal incident polarization states [32-36]. To the best of our knowledge, the mapping from

orthogonal polarization states to completely independent amplitudes has yet to be fully established in metasurface optics.

Here, we propose a new class of metasurface polarization optics where two arbitrary and independent amplitude profiles can be imposed on a pair of orthogonal polarization states (linear, circular, or elliptical) by a singlet metasurface, conceptually shown in Fig. 1(a). This is fundamentally different from previous works where the amplitude and phase control is only achieved for one specific incident polarization state [32-33]. Orthogonally polarized light incident on the metasurface experiences polarization-dependent constructive or destructive interference with contributions from both the propagation phase and the geometric phase. Here, the phase modulation act as an intermediary in associating the input polarization states with the output amplitudes of light. As the transmission amplitude control originates from the interference within each meta-molecule in our method, a single layer metasurface can perform the complete operation without requiring any additional optical elements, which is beneficial for their use in ultracompact integrated optical systems. To demonstrate the ability of this approach, we design and fabricate chiral grayscale metasurface and show nanoprinting with chiral shadow rendering of structured light. This method significantly expands the scope of metasurface polarization optics, paving the way towards engineering of novel polarization switchable meta-devices.

Interference, as an important fundamental optical effect, corresponds to the coherent interaction of two or more optical waves yielding a resultant intensity distribution. In accordance with the principle of superposition, the complex electric field \vec{E} , at a point in space, arising from separate fields \vec{E}_a and \vec{E}_b of two contributing sources is given by $\vec{E} = \vec{E}_a + \vec{E}_b$. Dielectric elements can be considered as individual sources because the polarization ellipse and phase of the optical field \vec{E}_{out} transmitted through each element can be arbitrarily controlled by the element design. The implementation of the proposed concept requires at least two different meta-atoms to act as interference sources. In order to make the meta-molecule be with standard square shape which ensures the identical amplitude modulation capability in both the horizontal and vertical directions, here we employ a pair of staggered twin-meta-atoms to design a meta-molecule and form a sub-micron interference system, as shown in Fig. 1(b). The anisotropy of the meta-atoms ensures that the system has the capability of providing polarization-dependent complex amplitude modulation. For each meta-molecule (*x*, *y*), the two components of output electric fields in the scalar form can be

expressed as:

$$P^{\pm}(x, y) = p(x, y)e^{i\varphi_{p\pm}(x, y)},$$
(1)

$$Q^{\pm}(x,y) = q(x,y)e^{i\varphi_{q\pm}(x,y)},$$
 (2)

where + and - denote a pair of orthogonal states of polarization. The interference occurs near the exit surface of the meta-atoms due to their subwavelength separation. Then the total complex amplitude of the interference system is given by $U^{\pm}(x, y) = P^{\pm}(x, y) + Q^{\pm}(x, y)$ and the resultant intensity is as:

$$|U^{\pm}(x,y)|^{2} = p^{2}(x,y) + q^{2}(x,y) + 2p(x,y)q(x,y)cos[\varphi^{\pm}(x,y)] , \qquad (3)$$

where $\varphi^{\pm}(x, y)$ is the phase difference of the complex amplitudes between two pairs of metaatoms and expressed as: $\varphi^{\pm}(x, y) = \varphi_{p\pm}(x, y) - \varphi_{q\pm}(x, y)$. The resultant intensity $|U^{\pm}(x, y)|^2$ therefore depends on the well-known interference term $2p(x, y)q(x, y)cos[\varphi^{\pm}(x, y)]$. Eq. (3) can be expressed as a composite function $|U^{\pm}(x, y)| = f[\varphi^{\pm}(x, y)]$, and thus its inverse function has the form $\varphi^{\pm}(x, y) = f^{-1}(|U^{\pm}(x, y)|)$. If it is possible to generate arbitrary value of $\varphi^{\pm}(x, y)$, it would directly contribute to the modulation of resultant intensity in transmission space. Therefore, $\varphi^{\pm}(x, y)$ act as an intermediary in associating the input polarization states with output amplitudes of light field.

Supposing that a metasurface is designed to impose two independent amplitude profiles $|U^+(x,y)|$ and $|U^-(x,y)|$ on a pair of orthogonal polarization states $\{\vec{\alpha}^+, \vec{\alpha}^-\}$, as shown in Fig. 1. Concurrently, the metasurface is also required to consistently transform the input polarization states to output polarization states $\{\vec{\beta}^+, \vec{\beta}^-\}$ as $\vec{\alpha}^+ \rightarrow \vec{\beta}^+$ and $\vec{\alpha}^- \rightarrow \vec{\beta}^-$ during transmission. Considering each constituent meta-atom is a linearly birefringent waveplate, then the relationship between the input polarization and output polarization is $\vec{\alpha}^+ = (\vec{\beta}^+)^*$, where * denotes complex conjugate [16, 23]. As mentioned above, using Eq. (3) one can convert the required amplitude profiles $|U^{\pm}(x,y)|$ into corresponding phase difference $\varphi^{\pm}(x,y)$. Hence, the metasurface can be described by a Jones matrix J(x, y) that simultaneously satisfies $J(x, y)\vec{\alpha}^+ = e^{if^{-1}(|U^+(x,y)|)}\vec{\beta}^+$ and $J(x, y)\vec{\alpha}^- = e^{if^{-1}(|U^-(x,y)|)}\vec{\beta}^-$. An arbitrary pair of orthogonal polarization states in the linear polarization basis has the form:

$$\vec{\alpha}^{+} = \begin{bmatrix} \cos\chi\\ e^{i\delta}\sin\chi \end{bmatrix} \qquad \vec{\alpha}^{-} = \begin{bmatrix} -\sin\chi\\ e^{i\delta}\cos\chi \end{bmatrix}, \qquad (4)$$

where χ and δ represent the azimuth and ellipticity of the polarization ellipse, respectively.

In this case, the required Jones matrix J(x, y) is:

$$J(x,y) = \begin{bmatrix} \cos\chi \cdot e^{if^{-1}(|U^{+}(x,y)|)} & -\sin\chi \cdot e^{if^{-1}(|U^{-}(x,y)|)} \\ e^{-i\delta}\sin\chi \cdot e^{if^{-1}(|U^{+}(x,y)|)} & e^{-i\delta}\cos\chi \cdot e^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} \cos\chi & -\sin\chi \\ e^{i\delta}\sin\chi & e^{i\delta}\cos\chi \end{bmatrix}^{-1},$$
(5)

This matrix provides a general mapping from orthogonal polarization states to independent amplitudes, which can be translated into the design of polarization-controlled amplitude metadevices. The eigenvalues and eigenvectors of J(x, y) determine the birefringence phase shifts and fast-axis orientation angle of a meta-atom, respectively corresponding to propagation phase and geometric phase. Therefore, in principle, independent amplitude profiles can be imposed on any set of orthogonal polarization states by simultaneously modifying a meta-atom's shape and angular orientation.

To demonstrate the amplitude control for any pair of orthogonal polarization states, including linear, circular, and elliptical polarization, we design and numerically simulate three kinds of meta-molecule arrays encoding normalized polarization-dependent amplitude values 0 and 1, as shown in Fig. 2. The meta-atoms in the designed meta-molecule are made of rectangular titanium dioxide (TiO₂) nanopillars on a fused-silica substrate. The TiO₂ nanopillars are designed to have identical heights of *H*=600 nm and organized in a square array with a lattice constant of *P* = 360 nm. Detailed information about these designs are described in the Supplemental Material, Sec. I. Based on Eq.(5), for linear polarization state, one can easily obtain the required phase shifts along fast and slow axes and rotation angles of meta-atom as $\varphi_x(x, y) = f^{-1}(|U^+(x, y)|)$, $\varphi_y(x, y) = f^{-1}(|U^-(x, y)|)$, $\theta(x, y) = 0$. It means that the constructed meta-molecules only need propagation phase design, which relies on the shape of the meta-atom. In contrast, for circular polarization state, one can find analytical solutions for the required phase shifts and rotation angle as a function of amplitude value given by (see Supplemental Material, Sec. II):

$$\varphi_x(x,y) = \frac{1}{2} [f^{-1}(|U^+(x,y)|) + f^{-1}(|U^-(x,y)|)], \tag{6}$$

$$\varphi_{y}(x,y) = \frac{1}{2} [f^{-1}(|U^{+}(x,y)|) + f^{-1}(|U^{-}(x,y)|)] - \pi,$$
(7)

$$\theta(x,y) = \frac{1}{4} [f^{-1}(|U^+(x,y)|) - f^{-1}(|U^-(x,y)|)].$$
(8)

Different from linear polarization state, a unification of propagation phase and geometric phase is employed to achieve the polarization-dependent constructive or destructive interference. In the more general case of an elliptical polarization state, the analytical solutions do not yield simple and explicit expressions and thus need to be calculated for specific amplitude values and ellipticity of incident polarization states. The specific designs of meta-molecule for two orthogonal linear ($\delta = \pi/2$ and $\chi = \pi/2$), circular ($\delta = \pi/2$ and $\chi = \pi/4$) and elliptical ($\delta = \pi/3$ and $\chi = \pi/6$) polarizations are shown in Fig. 2(a), 2(d) and 2(g), respectively.

The finite-difference time-domain (FDTD) simulation results of three metasurfaces shown in Fig. 2(b, e, h) agree well with the theoretical design. One of the incident polarization states experiences constructive interference, while the other orthogonal state undergoes destructive interference, respectively leading to a complete transmission and a reflection back towards the source. In Fig. 2(c, f, i), the calculated transmittance spectra exhibit broadband characteristics and a strong dichroism, which implies that the designed amplitude modulation metasurfaces can be used as ultrathin and high-performance dichroic filters for arbitrary orthogonal polarization states.

Relying on this ability to arbitrarily control amplitude using the designed meta-molecules, next we experimentally demonstrate a grayscale metasurface with gradient amplitude modulation. Since circular polarization has been widely investigated in metasurface polarization optics, here we choose two orthogonal circular polarization states for incident light for our proof-of-concept experimental demonstration. This demonstration can also be applied to orthogonal linear and elliptical polarizations as they have the similar physical mechanism for amplitude modulation. A metasurface encoding grayscale information but with opposite variation tendency for LCP and RCP light is designed. Fig. 3(a-b) show the calculated phase shifts, $\varphi_x(\varphi_y = \varphi_x - \pi)$ and rotation angle, θ as a function of intensity $|U^{\pm}(x,y)|^2$ for the designed metasurface, which intuitively shows the contribution of propagation phase and geometric phase. By using atomic layer deposition and electron beam lithography techniques (see Supplemental Material, Sec. III for details), the TiO₂ grayscale metasurface is fabricated and the measurement results are shown in Fig. 3(c-f). As expected, for LCP and RCP light at the wavelength of 550 nm incident on the metasurface, the measured intensity distributions exhibit smooth grayscale variation from bright to dark and the variation tendency is opposite for the two orthogonal circular polarization states. Slight deviations between the design and measured results shown in Fig. 3(e-f) may originate from the fabrication imperfection of the nanostructures.

Besides the grayscale metasurface, we further demonstrate the amplitude modulation-induced chiral shadow rendering of structured light. Two metasurfaces (metasurface 1 and metasurface 2)

are designed to respectively generate nanoprinting patterns of a character string 'NJU' and a bird perching on the twig. Different intensity profiles $|U^{\pm}(x,y)|^2$ for two orthogonal circular polarization states are encoded on the metasurfaces. A collimated circularly polarized light at the wavelength of 550 nm is normally incident upon the fabricated metasurfaces and the schematic diagram of the measurement setup is shown in Fig. 4(a). As shown in Fig. 4(b), under the LCP light illumination, the 'NJU' characters on metasurface 1 present a stereoscopic convex effect, and the metasurface 2 exhibits a frontal portrait of the bird with a crooked head. When the incident light is switched from LCP to RCP, the pattern of 'NJU' characters on metasurface 1 present a totally different concave effect and the metasurface 2 shows a side portrait of bird. Further results illustrating polarization-controllable nanoprinting for orthogonal linear, circular and elliptical polarization states are described in the Supplemental Material, Sec. IV. These results clearly demonstrate that amplitude distributions imparted on two orthogonal polarization states can be completely decoupled by the metasurface, and thus it expands the degree of freedom of amplitude control in metasurface optics. This expansion of functionality enabled by the proposed metasurface has the potential to enable novel applications in the areas of optical data storage, information encryption and anti-counterfeiting.

In conclusion, we have proposed and demonstrated a new class of metasurface that imposes two arbitrary and independent amplitude profiles on a pair of orthogonal polarizations. The phase profile of metasurface acts as an intermediary in associating the input polarization state with output amplitude of light field. By combining the geometric phase and propagation phase, we provide a simple method for the design of polarization-dependent, amplitude-controlled meta-devices. This method significantly expands the scope of metasurface polarization optics and opens a new path towards achieving novel types of photonic functionalities in a compact platform.

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FIG. 1. (a) Conceptual schematic diagram. Two independent amplitude profiles $|U^{\pm}(x, y)|$ can be imposed on a pair of orthogonal input polarization states $\vec{\alpha}^{\pm}$ through the metasurface, and makes the output polarization states $\vec{\beta}^{\pm}$ flip handedness. The green, red, and blue wavy lines donate the input, output, and reflection of optical waves, respectively. (b) Schematic diagram of metasurface array. The amplitude controlled metasurface is formed by meta-molecules array. Each metamolecule (blue dashed box) consists of multiplexed waveplate-like birefringent meta-atoms. The right bottom inset shows a scanning electron microscope image of the fabricated metasurface (Oblique view). (c-d) Perspective and top view of a meta-atom. The dimension (L, W) and orientation angle θ of the meta-atom can be varied to respectively provide desired propagation phase and geometric phase.



FIG. 2. Dichroism of orthogonal polarization states in a meta-molecule system, including (a-c) linear ($\delta = \pi/2$, $\chi = \pi/2$), (d-f) circular ($\delta = \pi/2$, $\chi = \pi/4$), and (g-i) elliptical polarizations ($\delta = \pi/3$, $\chi = \pi/6$). (a, d, g) Schematic diagrams of the meta-molecule design. (\vec{x}, \vec{y}), ($\vec{\sigma}$), ($\vec{\alpha}, \vec{\beta}$) denote the linear, circular and elliptical polarization, respectively. Here, the symbol * denotes complex conjugate. (b, e, h) The distribution of electric field in each meta-molecule under the illumination of corresponding orthogonal polarized light at the wavelength of 550 nm from the substrate side. (c, f, i) Broadband transmission characteristics of three meta-molecules.



FIG. 3. Grayscale metasurface and chiral transformation. (a, b) Required phase shifts, φ_x (top) and rotation angle, θ (bottom) as a function of the intensity $|U^{\pm}(x, y)|^2$ of incident wave. (c, d) Experimentally measured intensity distributions for LCP and RCP light at the wavelength of 550 nm. (e, f) The corresponding quantitative results along the horizontal direction of the device. The uncertainties are standard deviation of light intensity for repeated experimental measurements (five in total).



FIG. 4. Metasurface nanoprinting with chiral shadow rendering of structured light. (a) Experimental setup for measurement. A laser beam at the wavelength of 550 nm is collimated and incident on the metasurfaces though a linear polarizer (LP) and quarter wave plate (QWP). The images of metasurface nanoprinting can be captured by the standard microscope attached with a camera. (b) Experimentally captured optical images of the metasurface nanoprinting illuminated with LCP and RCP light.

Supplementary Materials for

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I .DESIGN AND SIMULATION OF PERIODIC ARRAYS OF META-MOLECULES

As mentioned in the main text, the meta-molecule designs are based on the Jones matrix J(x, y) in Eq. 5. We can extract the required phase shifts (φ_x , φ_y) and rotation angle θ of the dielectric unit from the eigenvalues and eigenvectors of the matrix. FIG. S1 and TABLE. S1 show the detailed structural parameters of the proposed meta-molecules. All the simulations shown in the main text are performed using a commercial finite-difference time-domain (FDTD) software with plane wave illumination from substrate for an infinitely periodic square arrays of the designed meta-molecules. In the simulations, periodic boundary conditions are employed at the *x* and *y* directions and perfectly matched layers (PML) at the *z* boundaries. The mesh size added on the meta-molecule is dx = dy = dz = 5 nm, which can ensure the accuracy of the calculated results. To obtain arbitrary polarized light, two orthogonal linearly polarized light sources with required phase difference and amplitude values are added in the simulation region.



FIG. S1. Details of the meta-molecule. Each meta-molecule is composed of two pairs of twin elements with different birefringent properties to form a submicron interference system. The proposed meta-molecule is composed of rectangular titanium dioxide (TiO₂) nanopillars on a fused-silica substrate. The TiO₂ nanopillars are designed to have identical heights of H=600 nm and organized in a square array with a lattice constant of P = 360 nm.

Table. S1. The meta-molecule designs for the demonstration of dichroism. LP: linearpolarization. CP: circular polarization. EP: elliptical polarization.

	LP	СР	EP	
Element 1	<i>l</i> _a =145 nm	<i>l</i> _a =100 nm	<i>l</i> _a =190 nm	
	w _a =145 nm	w _a =265 nm	w _a =100 nm	
Element 2	<i>l</i> _b =293 nm	<i>l</i> _b =125 nm	<i>l</i> _b =125 nm	
	wb=117 nm	w _b =305 nm	w _b =220 nm	

There are two possible arrangements (staggered and parallel) for a pair of twin-meta-atoms in a meta-molecule, as shown in Fig. S2. According to numerical simulations, these two arrangements offer similar amplitude response. The reason we choose staggered arrangement (Fig. S2 (a)) in our design is because compared to the parallel arrangement, the staggered case is easier to fabricate. For example, in some cases (as shown in Fig. S2(e)), the edge-to-edge distance between two neighboring meta-atoms in the parallel arrangement is much smaller than that can be achieved using state-of-the-art electron beam lithography techniques employed here for nanofabrication.



Figure. S2. Dichroism of orthogonal, elliptical polarization states ($\delta = \pi/3$, $\chi = \pi/6$) in a metamolecule system. The polarization conversion spectrum for the meta-molecule system with (a-b) staggered arrangement and (c-d) parallel arrangement. (e) Comparison of two possible nanostructures arrangements to form a square meta-molecule.

II : DERIVATION OF THE JONES MATRIX *J* AND ITS EIGENVALUES AND EIGENVECTORS FOR CIRCULAR POLARIZATION STATES

In this section, we emphasize only the relationship between amplitude and circular polarization states. Assume that the metasurface should impart two independent amplitude profiles, $|U^+(x,y)|$ and $|U^-(x,y)|$, on input circular polarization states $\{|R\rangle, |L\rangle\}$. As mentioned in the main text, the relationship between the input and output polarization states is complex conjugate. It means that the output polarization states have the same states as the input states with flipped handedness. Such meta-device can be expressed by Jones matrix J(x, y) that simultaneously satisfies:

$$J(x, y) |R\rangle = e^{if^{-1}(|U^+(x, y)|)} |L\rangle , \qquad (S1)$$

$$J(x,y) |L\rangle = e^{if^{-1}(|U^{-}(x,y)|)} |R\rangle .$$
(S2)

Upon matrix inversion of Eq. S1 and S2, the Jones matrix J(x, y) can be expressed as

$$J(x,y) = \begin{bmatrix} e^{if^{-1}(|U^{+}(x,y)|)} & e^{if^{-1}(|U^{-}(x,y)|)} \\ -ie^{if^{-1}(|U^{+}(x,y)|)} & ie^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} 1 & 1 \\ i & -i \end{bmatrix}^{-1}$$
(S3)

After calculation and simplification, the matrix J(x, y) has the form:

$$J(x,y) = \frac{1}{2} \begin{bmatrix} e^{if^{-1}(|U^{+}(x,y)|)} + e^{if^{-1}(|U^{-}(x,y)|)} & ie^{if^{-1}(|U^{-}(x,y)|)} - ie^{if^{-1}(|U^{+}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} - ie^{if^{-1}(|U^{+}(x,y)|)} & -e^{if^{-1}(|U^{+}(x,y)|)} - e^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix}$$
(S4)

This matrix provides a general form of switching between two independent amplitude profiles for circular polarization incident light. To understand the working principle of the metadevice more intuitively, we present the expression that relates output and input polarization light as $|E_{out}\rangle = J|E_{in}\rangle$. Here, incident light $|E_{in}\rangle$ is one of the two orthogonal, circular polarization states. The expansion of the output matrix has the form:

$$|E_{out}\rangle = \frac{1}{2} \begin{bmatrix} e^{if^{-1}(|U^{+}(x,y)|)} + e^{if^{-1}(|U^{-}(x,y)|)} & ie^{if^{-1}(|U^{-}(x,y)|)} - ie^{if^{-1}(|U^{+}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} - ie^{if^{-1}(|U^{+}(x,y)|)} & -e^{if^{-1}(|U^{+}(x,y)|)} - e^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} 1 \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} 1 \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} 1 \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\ ie^{if^{-1}(|U^{-}(x,y)|)} \end{bmatrix} \begin{bmatrix} 1 \\ ie^{if^{-1}(|U^{-}(x,y)|)} \\$$

$$= \frac{1}{2} \begin{bmatrix} 2e^{if^{-1}(|U^{+}(x,y)|)} \\ -2ie^{if^{-1}(|U^{+}(x,y)|)} \end{bmatrix}$$
$$= e^{if^{-1}(|U^{+}(x,y)|)} \begin{bmatrix} 1 \\ -i \end{bmatrix}$$
(S5)

The device has generated the targeted amplitude profile and expected polarization state. We can come to the same conclusions for the orthogonal state. By calculating the matrix J(x, y), we can obtain the eigenvalues as

$$\xi_1 = e^{i\left[\frac{1}{2}(f^{-1}(|U^+(x,y)|) + f^{-1}(|U^-(x,y)|)\right]} , \qquad (S6)$$

$$\xi_2 = e^{i\left[\frac{1}{2}\left((f^{-1}(|U^+(x,y)|) + f^{-1}(|U^-(x,y)|)\right) - \pi\right]} , \qquad (S7)$$

and eigenvectors as

$$|r_{1}\rangle = \begin{bmatrix} \cos\frac{1}{4} [(f^{-1}(|U^{+}(x,y)|) - f^{-1}(|U^{-}(x,y)|)] \\ \sin\frac{1}{4} [(f^{-1}(|U^{+}(x,y)|) - f^{-1}(|U^{-}(x,y)|)] \end{bmatrix},$$
(S8)

$$|r_{2}\rangle = \begin{bmatrix} -\sin\frac{1}{4}[(f^{-1}(|U^{+}(x,y)|) + f^{-1}(|U^{-}(x,y)|)] \\ \cos\frac{1}{4}[(f^{-1}(|U^{+}(x,y)|) + f^{-1}(|U^{-}(x,y)|)] \end{bmatrix}.$$
 (S9)

Since $|r_1\rangle$ and $|r_2\rangle$ are linearly independent vectors, the Jones matrix J(x, y) can be transformed into canonical form $J = P\Lambda P^{-1}$, where P is an invertible matrix and Λ is a diagonal matrix. The expansion of the canonical form can be expressed as

$$J(x,y) = P\Lambda P^{-1} = \begin{bmatrix} \cos\frac{1}{4}[F(U^{+}) - F(U^{-})] & -\sin\frac{1}{4}[F(U^{+}) - F(U^{-})] \\ \sin\frac{1}{4}[F(U^{+}) - F(U^{-})] & \cos\frac{1}{4}[F(U^{+}) - F(U^{-})] \end{bmatrix} \cdots$$
$$\begin{bmatrix} e^{i[\frac{1}{2}(F(U^{+}) + F(U^{-}))]} & 0 \\ 0 & e^{i[\frac{1}{2}(F(U^{+}) + F(U^{-})) - \pi]} \end{bmatrix} \begin{bmatrix} \cos\frac{1}{4}[F(U^{+}) - F(U^{-})] & -\sin\frac{1}{4}[F(U^{+}) - F(U^{-})] \\ \sin\frac{1}{4}[F(U^{+}) - F(U^{-})] & \cos\frac{1}{4}[F(U^{+}) - F(U^{-})] \end{bmatrix} \end{bmatrix}$$
(S10)

Here, $F(U^+)$ and $F(U^-)$ represent $f^{-1}(|U^+(x,y)|)$ and $f^{-1}(|U^-(x,y)|)$, respectively. Since the Jones matrix of the meta-device operates on linear polarization basis, we can easily obtain the required phase shifts and rotation angles as a function of amplitude value

$$\varphi_x(x,y) = \frac{1}{2} [f^{-1}(|U^+(x,y)|) + f^{-1}(|U^-(x,y)|)],$$
(S11)

$$\varphi_{y}(x,y) = \frac{1}{2} [f^{-1}(|U^{+}(x,y)|) + f^{-1}(|U^{-}(x,y)|)] - \pi , \qquad (S12)$$

$$\theta(x,y) = \frac{1}{4} [f^{-1}(|U^+(x,y)|) - f^{-1}(|U^-(x,y)|)] .$$
(S13)

III: NANOFABRICATION OF TiO2 METASURFACE



FIG. S3. Schematic illustrate of the metasurface fabrication. (a) Spin-coated photoresist. A doubleside polished fused silica substrate was prime-vapor-coated with a layer of hexamethyldisilazane (HMDS) and then spin-coated with a layer of 600 nm thick, positive-tone electron beam resist (ZEP520A). (b) The e-beam lithography (EBL) is used to obtain the exposed pattern. This process was performed at an accelerating voltage of 100 kV and beam current of 2 nA. Then, the sample was developed in hexyl-acetate for two minutes. (c) Atomic layer deposition (ALD) process. The patterned sample was carried out the deposition of TiO₂. To avoid deformation of the resist pattern, the ALD chamber is set to 90 °C. (d) The over-coated TiO₂ layer was etched by the inductivelycoupled-plasma reactive ion etching (ICP-RIE) in a mixture of Cl₂ and BCl₃ gas. The etching was

terminated when the over-coated TiO_2 had been completely removed and the resist was exposed. Finally, we removed the resist by soaking in n-methyl-2-pyrrolidone and produced the array of TiO_2 nanopillars with pre-designed geometries (e).

IV: SIMULATION OF METASURFACE NANOPRINTING FOR ANY PAIR OF ORTHOGONAL POLARIZATION STATES

In this section, we design and numerically simulate three metasurface arrays to demonstrate that our method can achieve two completely different nanoprinting patterns for any pair of orthogonal polarization states, including linear, circular, and elliptical polarization. Each metasurface array is designed to respectively generate two different character strings 'Meta' and 'Nano' for two orthogonal polarization states at the wavelength of 550 nm. Considering the calculation limit of the simulation workstation, the metasurface encoding these amplitude profiles is 36 μ m × 108 μ m in size and contain 50 × 150 meta-molecules. The simulation results are shown in Fig. S4. As expected, the metasurfaces present a pattern of 'Nano' characters with stereoscopic concave effect for one polarization state. When the incident light is switched to its orthogonal state, the metasurfaces exhibit a pattern of 'Meta' characters with totally different convex effect. These results are in good agreement with expectations and clearly demonstrate that independent amplitude control can be achieved for any pair of orthogonal polarization states. The rough image quality is mainly attributed to the small number of meta-molecules in the simulation. The corresponding metasurface arrays and feature sizes of TiO₂ nanopillars are shown in Fig. S5. and Table S2- S4, respectively.



FIG. S4. Polarization-controllable nanoprinting to achieve two completely different transmitted images for any pair of orthogonal polarization states, including linear ($\delta = \pi/2$, $\chi = \pi/2$), circular ($\delta = \pi/2$, $\chi = \pi/4$), and elliptical polarizations ($\delta = \pi/3$, $\chi = \pi/6$). The metasurface encoding these amplitude profiles was 36 × 108 µm in size and contained 50 × 150 meta-molecules.



FIG. S5. The designed metasurface arrays used for (a) linear, (b) circular, and (c) elliptical polarizations. The left and right insets show a magnified view of the metasurface arrays (Top view).

	Unit cell				
No.	Length(nm)	Width (nm)	No.	Length(nm)	Width (nm)
1	280	280	15	140	155
2	180	180	16	145	145
3	155	185	17	120	215
4	163	163	18	125	185
5	140	200	19	130	170
6	145	175	20	130	160
7	155	155	21	135	150
8	135	200	22	140	140
9	140	175	23	90	275
10	145	165	24	100	210
11	150	150	25	105	185
12	125	220	26	105	180
13	135	175	27	110	165
14	140	160	28	110	155

Table S2. Feature sizes of TiO₂ nanopillars used for linear polarization.

	Unit cell				
No.	Length(nm)	Width (nm)	No.	Length(nm)	Width (nm)
1	90	275	5	120	300
2	100	270	6	125	320
3	110	280	7	315	65
4	115	295	8	275	80

Table S3. Feature sizes of TiO_2 nanopillars used for circular polarization.

Table S4. Feature sizes of TiO_2 nanopillars used for elliptical polarization

	Unit cell				
No.	Length(nm)	Width (nm)	No.	Length(nm)	Width (nm)
1	230	135	9	80	235
2	240	145	10	300	155
3	245	155	11	295	165
4	155	295	12	295	185
5	290	130	13	110	195
6	290	145	14	105	255
7	280	160	15	105	290
8	290	175	16	105	335