Bioinspired micrograting arrays mimicking the reverse color diffraction elements evolved by the butterfly Pierella luna

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Recently, diffraction elements that reverse the color sequence normally observed in planar diffraction gratings have been found in the wing scales of the butterfly Pierella luna. Here, we describe the creation of an artificial photonic material mimicking this reverse color-order diffraction effect. The bioinspired system consists of ordered arrays of vertically oriented microdiffraction gratings. We present a detailed analysis and modeling of the coupling of diffraction resulting from individual structural components and demonstrate its strong dependence on the orientation of the individual microgratings. This photonic material could provide a basis for novel developments inbiosensing, anti-counterfeiting, and efficient light management in photovoltaic systems and light-emitting diodes.

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Three-dimensional photonic crystals (1–5), materials with two-dimensional micro- or nanosized periodic morphologies (6–8), and one-dimensional multilayer configurations (9) have been identified as the primary cause of structural coloration in a wide variety of nonrelated biological organisms. In contrast, surface-confined diffraction elements for the separation of incident light into specific colors are less abundant in nature and have only been discovered in a handful of organisms (10), including a fossil polychaete (7), sea mouse Aphrodita sp. (6), and some flowering plants (11). Recently, diffraction elements that reverse the color sequence normally observed in planar diffraction gratings have been found in the scales of the butterfly Pierella luna (12).

Inspired by this biological light manipulation strategy, we devised an artificial material morphology mimicking the butterfly’s diffraction effect by creating periodic arrays of vertically oriented individual microdiffraction gratings. In addition to the butterfly-inspired reverse color-order diffraction arising from each individual micrograting, the periodicity between the individual gratings causes diffraction on a different length scale, leading to complex intensity distributions in experimentally measured angularly resolved reflection spectra. An in-depth analysis of the observed diffraction phenomenon complemented by optical modeling revealed a strong dependence of the optical signature on the orientation of the gratings. Such an effect can only be seen because of the hierarchical nature of the superposed, orthogonal grating features. To further elucidate the role of the different structural components for the emerging reflection spectra, the initially vertically oriented individual microgratings were subjected to a tilt, resulting in a predictable change of the surface’s optical signature.

The dorsal side of the fore- and hind wings of P. luna males are dull brown in diffuse ambient illumination (Fig. 1A, Left). When exposed to directional illumination at grazing incidence, a coin-sized spot on each fore wing displays an angle-dependent color variation across the whole visible spectrum (Fig. 1A, Right). The color changes from red to blue with increasing observation angle, unlike the variation from blue to red normally observed in conventional diffraction gratings (13). This reverse color diffraction effect results from the local morphology of individual scales within the colored spot on the fore wings (12). The top parts of the scales are curled upward, orienting lines of periodically arranged cross-ribs perpendicular to the wing surface. (Fig. 1C and D). Light incident at an angle onto the curled parts of the scales is diffracted by the cross-rib structure acting as a diffraction grating, with a periodicity of ∼400 nm. The alignment of the grating perpendicular to the surface results in the reverse color sequence that can be observed in angularly resolved reflection spectra and in the diffraction pattern (Fig. 1 E and F).

The identification of this unusual diffraction effect on the wings of P. luna (12) provided inspiration for the development of a bioinspired photonic system that incorporates vertically oriented microdiffraction gratings with submicrometer periodicity analogous to the key features observed in the natural structure. In addition, the artificial system displays a periodic arrangement of the individual vertical gratings in large arrays with 2D micromscale periodicity. This structural feature, which is not found in the natural organism, enriches the optical signature of the artificial system via coupling of the diffractive modes of the two present hierarchical morphologies. In the following, we discuss the optical properties of the artificial system and demonstrate that the modification of either one of the grating morphologies changes the diffraction signature in predictable ways.

Significance

In the course of evolution, many organisms have developed unique light manipulation strategies that rely on intriguing combinations of a broad range of optical effects generated by materials with sophisticated multiscale hierarchical structural arrangements. By exploiting the optical principles underlying natural structural color, we can generate new photonic materials. Researchers have only just begun to match nature’s morphological and compositional complexity in man-made materials using nanofabrication. We present a bioinspired photonic material that mimics the reverse color-order diffraction found in the butterfly Pierella luna. Exploiting and improving the butterfly’s strategy, we create photonic materials that increase our basic understanding of the optical interplay of hierarchical structures and provide a platform for the development of novel photonic devices.

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The artificial system consists of an array of individual 10-μm-long, 2-μm-wide, and 18-μm-high plates arranged in rows with an interplate spacing of 10 μm and a separation of 5 μm between individual rows of plates (Fig. 2A). These parameters result in an overall periodicity of 12 μm in the direction perpendicular to the plates and 15 μm in the direction collinear with the plane of the plates and the sample surface. A periodic wave pattern—termed “scallops”—of ∼500-nm pitch runs along the sides of each individual plate (Fig. 2B).

The bioinspired diffraction elements are fabricated in a double-molding procedure. Starting from a silicon master, a periodic array of scalloped microplates is first cast into polydimethylsiloxane (PDMS) to form a negative mold (15), which is then replicated with a UV-curable epoxy to produce a positive replica of the master silicon structure with the original scallops on the individual plates well preserved. The silicon master is formed using the Bosch process (16), in which multiple etching and passivation steps give rise to the periodic undulations on the microplate surface. The pitch and height of these grating structures can be controlled by adjusting the etching parameters (17, 18). Here, they are chosen to be comparable to the spacings and dimensions of the diffraction-inducing microrots on the P. luna scales (Fig. 2B), and hence are expected to cause a similar diffraction effect.

Notice that in the biological system, the periodicity of the diffraction-grating-supporting scales is of the order of 80 ± 10 μm along their length and 60 ± 10 μm perpendicular to the scale axis (Fig. 1B). Due to these large distances between diffraction elements and a nonnegligible amount of irregularity in the location of individual scales, no coherence is observed for light diffracted from adjacent scales. The overall color splitting only results from the diffraction caused by the cross-rib gratings on the individual scales, which is confirmed by variable-angle spectroscopy and by diffraction microscopy measurements (Fig. 1 D–F). Unlike in the biological system, the individual microdiffraction gratings in the artificial system are intentionally arranged in a highly periodic manner, which is expected to result in a richer diffraction signature and provide additional possibilities of tailoring the interaction of light beyond the diffraction induced by the scallops on the plates.

Variable observation-angle spectroscopy performed on the artificial system serves to spectrally and angularly resolve parts of the complex diffraction pattern (Fig. 2C). For each measurement the plane of light incidence is chosen to be perpendicular to the surface of the individual microplates. The light incidence angle θl is fixed and the observation angle θD is varied in the plane of light incidence to capture light reflected in an angular range of ±75° around the sample surface normal. Two main features are observed in these measurements:

i) Straight lines of higher intensity resulting from diffraction caused by the interplate periodicity represent the individual diffraction orders; the experimentally observed locations of these diffraction orders (shown in Fig. 2 C and H) can be directly calculated using the grating equation (13). An example of such a calculation can be seen in Fig. S1. Due to the large interplate pitch of 12 μm, the angular separation between adjacent diffraction orders and the free spectral range of each individual order are small. For light incident at an angle θl = 45°, eight positive propagating diffraction orders [left of the zeroth order at sin(θ) = −0.71 in Fig. 2C] and 53 negative propagating orders are captured with the highest intensity in the direct reflection (zeroth order) and in the adjacent orders.

ii) There is an arc-shaped distribution of intensity maxima across different diffraction orders (emphasized by the red shaded region in Fig. 2C).

This anomalous redistribution of light in the diffraction orders is caused by the scallops on each individual plate. To properly describe the diffraction resulting from the microdiffraction gratings oriented normally to the substrate, the grating equation (13) has to be reformulated taking into account their vertical orientation for diffraction in the plane of the scallops:

\[ \lambda = \frac{d}{m} \left( \cos \theta_l + \cos \theta_D \right) \]  

where \( d \) is the grating periodicity, \( m \) is the diffraction order, \( \theta_l \) is the light incidence angle, and \( \theta_D \) is the diffraction angle. This equation describes the arc-shaped pattern observed in the
experiment under the assumption that the plates are vertical (Fig. 2C, red shaded region).

Imaging of the diffraction patterns by diffraction microscopy provides a direct visualization of the effects observed in the variable-angle spectroscopy measurements. The angle range that can be visualized in these measurements is determined by the numerical aperture of the microscope objective. For a given light incidence angle \( \theta \), the diffraction caused by the periodic ensemble of plates is most clearly observed when collecting light with the objective’s axis aligned with the specular reflection direction \( \theta_p = -\theta \) (Fig. 2D, with the signal in the blue box corresponding to the blue shaded spectral range in Fig. 2C). Imaging of the sample’s diffraction in the Littrow mounting (13), where light is incident on the sample through the microscope objective, allows for the capturing of the diffraction component induced by the scallops on the individual plates (Fig. 2E, with the signal in the yellow box corresponding to the yellow shaded spectral range in Fig. 2C). This feature is easily distinguished from the diffraction of the plate ensemble by the wider color spread.

Overall, the measurements provide clear evidence of coupling between the first-order mode of the scallop diffraction and the interplate periodicity-based diffraction modes. The diffraction coupling can be controlled by adjusting the pitch of the scallops on individual plates and the interplate geometry in the manufacturing procedure. Likewise, a change in inclination of the microdiffraction gratings with respect to the substrate is expected to be reflected in a variation of the diffraction patterns. This change can indeed be observed by imposing a controlled tilt of \( \sim 20^\circ \) on the microplates by shearing the sample (Fig. 2F and G).

The bending leaves the locations of the diffraction modes resulting from the interplate periodicity largely unaffected. Only minor wavelength-dependent intensity variations in each diffraction mode are apparent (Fig. 2H and I). At large angles \( \theta_D \), measured from the sample surface normal, a region of higher reflection intensity appears, which is related to the inclination of the individual plates, i.e., the blaze of the grating array. By contrast, the diffraction caused by the scallops on the individual plates is strongly influenced by the variation in plate tilt angle shifting the observed arc-shaped pattern in wavelength and angular position (Fig. 2H and K). For micrometer-sized diffusion gratings, which have a tilt angle \( \beta \) relative to the surface normal, the diffraction grating equation can be reformulated as

\[
\frac{\lambda}{m} = d \left( \cos(\theta_1 - \beta) + \cos(\theta_D - \beta) \right).
\]

By calculating a fit for the arc-shaped intensity distribution across the different diffraction orders in the variable-angle spectroscopic data (Fig. 2H, red shaded region), the tilt angle of the plates relative to the surface normal is found to be \( \beta \sim 23^\circ \), in agreement with the SEM image analysis.

These findings are further supported by finite-difference time-domain simulations (FDTD) (19). By comparing the plates with and without scalloping, the arc-shaped intensity distribution across diffraction orders is clearly identified as the diffraction resulting from the scallops (Fig. 3). Simulations of 23° tilted plates with 45° incident illumination show that this arc-shaped diffraction pattern gets skewed and spectrally shifted as predicted by Eq. 2 and observed in the experiments; the influence of the blaze of
the regular array of individual gratings, separate from the effect of the scallops, can also be seen by closely examining Fig. 3 and noting that the two tilted plate simulations have regions of high-intensity reflection at \( \sin(\theta) \sim 0 \) whereas the upright plates do not. Furthermore, the simulations serve to predict and optimize the influence of variations in interplate geometry, scallop grating pitch and shape, and plate tilt angle before the manufacturing of the system.

Whereas the angular positions of the diffraction modes resulting from the interplate periodicity only vary with the angle of light incidence \( \theta_i \), for a given \( \theta_f \) the diffraction pattern caused by the regular scallops on individual plates is strongly affected by a change in tilt of the plates relative to the sample normal. Consequently, a means of reversibly varying the plate tilt angle can provide the possibility to dynamically tune the diffraction pattern independent of light incidence. Possible means of achieving such reversible actuation include embedding the plates in a stimuli-responsive hydrogel (20–23), using a soft material negative of the structure and applying shear force to bend the microgratings, modifying the tips of the plates with ferromagnetic particles to allow for dynamic reconfiguration of the plate geometry using magnetic fields (24), or implementing tuning mechanisms shown for simple planar diffraction gratings that rely on electric fields or mechanical deformation (25–28).

Biological strategies for light manipulation have already been successfully implemented in nanophotonic devices for applications in chemically selective vapor sensing, pH determination, infrared imaging, surface-enhanced Raman spectroscopy-based chemical analysis, and localized heating from infrared absorption (29–34). The rich and tunable optical signature of our hierarchical bioinspired diffraction-based photonic material platform could provide a basis for novel developments in biosensing (35–37), efficient light management in photovoltaic systems (38–40), enhanced light extraction and radiation profile shaping in light-emitting diodes (41–43), and optically variable devices in consumer product design and anticounterfeiting (44–46).

Our results demonstrate the versatility of a bioinspired approach toward the creation of novel photonic systems. The unique diffraction-inducing nano- and microscale architecture previously discovered in the scales of the male butterfly *P. luna* served as inspiration for artificial microdiffraction grating arrays. Our photonic system mimics the reverse diffraction color sequence found in the butterfly’s scales and also provides additional complexity in the diffraction patterns due to a periodic arrangement of the diffraction elements not found in the natural structure. Such arrays provide a platform for hierarchical photonic systems displaying unique diffraction coupling. A detailed optical analysis and modeling of the diffraction patterns allowed us to observe, understand, and decouple diffraction effects induced by the plate ensemble and by the regular scalloping of individual plates. A variation of the light incidence angle results in the expected shifting of the plate ensemble diffraction modes but does not affect the diffraction resulting from individual plates. On the other hand, a variation in plate inclination leaves the interplate geometry diffraction modes untouched but has a strong influence on the scallop-induced diffraction.

The intensity distribution of the diffraction induced by the scallops is modulated by the diffraction induced by the plate ensemble. Where the scallop diffraction arc coincides with a plate ensemble diffraction order a peak in intensity is observed. A decrease in the plate ensemble period would result in fewer propagating plate ensemble diffraction orders with a larger free angular range in between. This would lead to a discretization of the scallop diffraction pattern (seen in the FDTD simulation in Fig. 4) inducing a discrete and easily perceivable color variation with potential applications in the development of novel optically variable devices in security printing and consumer product labeling. Currently, efforts are underway to fabricate the diffraction structures in different material combinations that provide a higher refractive index contrast, thereby strengthening the diffractive signal.

**Materials and Methods**

**Manufacture of Artificial Diffraction Structures.** When Bosch etching (16) is used in conjunction with photolithography, the multiple etching and passivation steps give rise to a periodicity in the sidewall of the structures due to the repeated underetching. Although typically the goal of this etching method is to create vertical sidewalls, these undulations in the sidewall can form a diffraction grating similar to that on the *P. luna* scales if the periodicities are chosen to fall within the range of optical wavelengths.

Once structures with the correct geometry are created in silicon, they can be replicated in other materials with better optical properties by using soft lithographic methods (15). By using soft, transparent materials instead of silicon, the structures can be bent much more easily and the optical properties can be changed by applying metal coatings or doping the material with pigments or other materials with interesting optical properties. The replication is achieved by using a PDMS mold of the silicon master and curing an epoxy via UV light before removal from the mold.

**Structural and Optical Analysis.** Once scalloped plate structures are made, optical characterization is performed for comparison of the structures observed on *P. luna* and the fabricated systems. Several different types of
characterization were used, including variable-angle spectroscopy, diffraction microscopy, and SEM.

For the variable-angle spectrometry setup, an Ocean Optics DH-2000 ultra-violet-visible-near infrared light source was used to illuminate a small spot (~1 mm) of the sample at a given incidence angle i. For each angle of illumination, light was collected at half-degree increments for –75° to +75° relative to the sample normal and spectrally analyzed using an Ocean Optics Maya Pro-2000 spectrometer.

For the diffraction microscopy images, a Bertrand lens was used to focus on the back focal plane of the objective of an upright BXFM Olympus optical microscope. The sample was illuminated either from an external source oriented at a 90° angle to the microscope objective with a spot size larger than the sample field of view, or with a small-diameter optical fiber coupled into the microscope light path (Littrow mounting).

SEM images were taken on a Zeiss Ultra55 or a Zeiss Supra55VP SEM after coating the polymer structures with a thin layer of gold to prevent charging.

Optical Modeling. FDTD (47) methods were used to numerically simulate the results obtained from the variable-angle spectroscopy measurements. A commercial-grade simulator based on the FDTD method was used to perform the calculations (19). Perfectly matched layers (48) were used to prevent reflections from the top and bottom of the simulation cell, whereas Bloch boundary conditions were used for the sides of the simulation cell to model light scattering from a periodic array at nonnormal incidence. To simulate broadband angled scattering, each simulation was repeated for a range of Bloch wavevectors, which were combined to produce scattered field profiles corresponding to an incidence angle of 45° ± 1° for wave-lengths of 320–800 nm. Far-field projections of the calculated local fields resulted in diffraction patterns (Fig. 3) analogous to the variable-angle spectroscopic data.

The diffraction discretization FDTD simulation was similarly performed on MEEP, a free FDTD software (49). The use of normal incidence illumination for this simulation allowed all frequencies and angles to be calculated without using a sweep of incidence angles.

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