Structural colors in metasurfaces: principle, design and applications

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Structural colors resulting from the interference between incident light and structures are ubiquitous in nature and daily life. In contrast to conventional chemical pigments and dyes, structural colors have the advantages of low consumption, environmental protection and high durability. To reach higher resolutions beyond diffraction limits, structural colors based on metasurfaces consisting of artificially designed nanoresonators at subwavelength scale have been proposed in recent years. In this review, we classify structural colors into plasmonic colors and dielectric colors according to different mechanisms, summarizing the development and applications for a variety of color filters. Moreover, four significant methods of dynamically tunable colors are discussed, including mechanical stretching, chemical reaction, electrical control, and photon doping.

1. Introduction

Colors as important visual carriers in our daily lives have been widely studied since prehistoric times. 140 years ago, the discovery of mauveine by W. H. Perkin promoted development of the extensive synthetic dyes and pigments industry.1 So far, synthetic colorants have been used in decorations, synthetic fibers, printing inks and identification, for example.2–4 In particular, organic pigments are finding increasing use in high-technology industries, such as optoelectronic displays, optical data storage and photo-reprographics (Fig. 1a).4 However, from the perspective of recycling, chemical colorants are costly and environmentally damaging. In addition, their insufficient resolution and lack of durability invariably limit their practical applications in displays and images.5,6

To overcome these limitations, considerable attention has been paid to structural colors, which are an interference phenomenon between incident light and structures. Structural colors are ubiquitous in nature and daily life, such as in beetles, Pollia fruits, peacocks, hummingbirds, morpho butterflies

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and so on. For instance, reflected light from the complex nanoridges and lamellae in the morpho rhetenor wings exhibits bright metallic blue, as shown in Fig. 1b. Compared with chemical pigments, structural colors have the advantages of (1) showing brighter colors under sunlight, (2) higher resolutions, (3) the ability of dynamic tuning by changing surface structures, and (4) better durability under high-temperature and high-intensity illumination. In general, structural colors are generated from different bulk materials including light scattering, photonic crystals and multilayer materials. Nevertheless, owing to bulk effects, high-quality colors always require more particles, larger periods and added layers, indicating that the thickness of color structures is much greater than the wavelength. As a result, there is still considerable room to further improve the resolution of structural colors beyond the diffraction limit.

In this context, a metasurface as a two-dimensional (2D) artificial structure at a subwavelength scale is a good candidate for the generation of high-resolution structural colors. Controlled optical properties can be introduced by engineering the geometric parameters of the periodic resonators array. According to the type of material of the resonators, metasurfaces can be classified into plasmonic and dielectric metasurfaces. For plasmonic metasurfaces, optical responses are mainly driven by plasmonic resonance based on a variety of metallic nanostructures. In contrast, for dielectric metasurfaces, electric and magnetic-dipole resonances generated from high-index dielectric materials simultaneously simulate novel optical phenomena. So far, with the development of micro-nano fabricated techniques, a number of unprecedented optical properties based on metasurfaces have been demonstrated to control and guide electromagnetic waves from microwaves to the visible range, such as the spin Hall effect, anomalous refraction, nonlinear optics and, in particular, structural colors.

In this review article, we focus on the development of structural colors generated from plasmonic and dielectric metasurfaces on the basis of principle, design and applications. A variety of color filters with different nanostructures and materials are systematically analyzed and summarized from the static state to the dynamic state. The methods of dynamically tunable colors are discussed, including mechanical stretching, chemical reaction, electrical control and photon doping. In addition, we also introduce practical applications of color filters, such as high-end displays, encryption and colorimetric sensing, indicating the wide prospects in real life.

2. Color generation from metasurfaces

2.1 Colors from plasmonic metasurfaces

Surface plasmons (SPs), the collective oscillation of free electrons and photons at the interfaces of metal and dielectric materials, have shown the unprecedented properties of strongly enhanced local field and in-plane momentum. SP generated from plasmonic metasurfaces plays a vital role in many applications, such as optical holograms, waveguides, flat lenses, directional couplers, waveplates and so on. In particular, one of the most intriguing areas of research in recent years is the study of plasmonic structural colors. The plasmonic resonance by coupling between SP and incident light can selectively filter visible light in transmission or reflection mode. To date, many plasmonic additive and subtractive color filters beyond the subwavelength diffraction limit have been extensively reported due to high resolution, ease of fabrication and high flexibility at the subwavelength scale. In this section, some typical plasmonic nanostructures composed of nanoratings, nanoholes, disk-hole coupling nanostructures and metal–insulator–metal (MIM) structures are briefly discussed.
In its infancy, the study of plasmonic colors mainly concentrated on the transmission mode of the resonant spectra. Extraordinary optical transmission (EOT)\(^4\)\(^6\) based on SP in an optically-thick opaque film can filter light at a specific resonant wavelength and lead to additive colors. In contrast, the counter-intuitive extraordinary low transmission (ELT) phenomenon, \(^5\)\(^0\) realized by optically-thin semi-transparent metallic nanostructures, can selectively eliminate light at the resonant wavelength to generate subtractive colors. As shown in Fig. 2a(ii), a subtractive color filter based on 30 nm-thick Ag film deposited on glass slide, showing the word of “OPTICS”. a(ii) Schematic diagram of the designed nanograting. a(iii) Measured transmission spectra of yellow, magenta and cyan grating with different periods. Insets show the corresponding SEM images of nanogratings. Reproduced from ref. 51, with permission from Nature Publishing Group, Copyright 2013. b(i)–(iii) b(i) SEM images of color filter composing Al nanoholes deposited on quartz substrate. b(ii) Measured transmission spectra of red, green and blue filters at 0° and 30° polarization angle. Insets show the corresponding top-view schematic of plasmonic nanoholes. b(iii) Optical microscope images of color filters with round holes and triangular holes on hexagonal lattice and round holes on square lattice. Reproduced from ref. 55, with permission from American Institute of Physics, Copyright 2011. c(i)–(iv) c(i) Schematic drawing of MIM tandem nanodisk array. Al\(_2\)O\(_3\) layer is sandwiched between identical Ag layers in the unit cell. c(ii) Calculated scattering cross-section for a single MIM nanodisk. Insets shows the charge distribution of in-phase and out-of-phase modes, respectively. c(iii) and (iv) Measured colors palettes in (i) transmission and (iv) reflection mode. Reproduced from ref. 61, with permission from American Chemical Society, Copyright 2017. d(i)–(iv) d(i) Schematic diagram and d(ii) SEM images of disk-hole coupling metasurface beyond optical diffraction limit. d(iii) Full-color printing image composing of disk-hole unit cells with distinct geometric sizes. Insets show the enlarged region and corresponding SEM image. d(iv) Resolution test pattern with array of 3 × 3 structures and 2 × 2 structures. Reproduced from ref. 66, with permission from Nature Publishing Group, Copyright 2012.

Fig. 2 Structural colors generated by plasmonic resonance. (a(i)–(iii)) a(i) Photo of a plasmonic nanograting consisting of 30 nm-thick Ag film deposited on glass slide, showing the word of “OPTICS”. a(ii) Schematic diagram of the designed nanograting. a(iii) Measured transmission spectra of yellow, magenta and cyan grating with different periods. Insets show the corresponding SEM images of nanogratings. Reproduced from ref. 51, with permission from Nature Publishing Group, Copyright 2013. (b(i)–(iii)) b(i) SEM images of color filter composing Al nanoholes deposited on quartz substrate. b(ii) Measured transmission spectra of red, green and blue filters at 0° and 30° polarization angle. Insets show the corresponding top-view schematic of plasmonic nanoholes. b(iii) Optical microscope images of color filters with round holes and triangular holes on hexagonal lattice and round holes on square lattice. Reproduced from ref. 55, with permission from American Institute of Physics, Copyright 2011. (c(i)–(iv)) c(i) Schematic drawing of MIM tandem nanodisk array. Al\(_2\)O\(_3\) layer is sandwiched between identical Ag layers in the unit cell. c(ii) Calculated scattering cross-section for a single MIM nanodisk. Insets shows the charge distribution of in-phase and out-of-phase modes, respectively. c(iii) and (iv) Measured colors palettes in (i) transmission and (iv) reflection mode. Reproduced from ref. 61, with permission from American Chemical Society, Copyright 2017. (d(i)–(iv)) d(i) Schematic diagram and d(ii) SEM images of disk-hole coupling metasurface beyond optical diffraction limit. d(iii) Full-color printing image composing of disk-hole unit cells with distinct geometric sizes. Insets show the enlarged region and corresponding SEM image. d(iv) Resolution test pattern with array of 3 × 3 structures and 2 × 2 structures. Reproduced from ref. 66, with permission from Nature Publishing Group, Copyright 2012.
In addition, plasmonic colors can be achieved in a reflection mode based on specific nanostructures and mechanisms, such as MIM structures. Generally speaking, MIM nanostructures composed of a thin dielectric layer between the metal nanoantennas and the metal film can exhibit strong standing-wave resonance where the incident light is confined into the metallic gap.\textsuperscript{56–61} The weak interaction between adjacent unit cells results in encoding color into an individual nanoantenna. A single reflected peak can be formed to contribute to slightly higher saturation and a full hue of colors. Nevertheless, there is still a need to pursue higher-saturation colors performed by reflection spectra with a narrower peak. In this context, another MIM nanostructure is proposed to generate full colors in Fig. 2c.\textsuperscript{61} As shown in the schematic presented in Fig. 2c(i), a dielectric $\text{Al}_2\text{O}_3$ nanodisk is sandwiched by metallic Ag layers, which can be regarded as MIM structures. The scattering cross-section spectra of a single nanodisk has two hybridized plasmonic resonances, at high frequency $\omega_s$ and low frequency $\omega_l$. The corresponding charge distribution is shown in Fig. 2c(ii), implying the magnetic field between adjacent nanodisks is enhanced. The hybridization between Wood’s anomaly and the in-phase electric modes give rise to a narrower peak with high efficiency in the reflection mode and a lower valley in the transmission mode. As shown in Fig. 2c(iii), additive primary colors with higher saturation and brightness in the reflected field and subtractive primary colors (CMYK) in the transmissive field can be simultaneously generated by using MIM sandwich nanostructures.

Besides color saturation and hue, resolution is another crucial property for plasmonic structural colors in practical applications. As mentioned above, the pixels of nanodisks, nanoholes and gratings are approaching wavelength scale. In order to further improve resolution, disk-hole coupling nanostructures\textsuperscript{62–66} are introduced in Fig. 2(d). Each pixel consists of four nanodisks, where Ag disks and the Au capping layers are raised above equally sized nanoholes on a backreflector (Fig. 2d(i and ii)). The Fano resonance resulting from the interference between the sharp resonance of the surface mode and the broad resonance of the nanodisk–nanohole coupling can lead to dips in the reflection spectra. Incident light flows around the nanodisks, through the nanoholes, and is absorbed by the backreflector. The interaction between Fano resonance and SP enables performing colors in a bright field. Furthermore, the resolution of pixels arranged in 250 $\times$ 250 nm units can reach 100 000 dpi. The resolution test pattern in Fig. 2d(iv) specifically shows the high resolution that the coupling nanostructures can reach. When the size of pixels is reduced to 250 nm, all colors in the chequerboard are preserved well, implying the resolution is approaching the optical diffraction limit.

Overall, high-quality structural colors both in reflection and transmission modes have been extensively generated by a variety of plasmonic metasurfaces.\textsuperscript{67–78} Compared with chemical pigments and dyes, plasmonic colors are preferable in terms of environment protection, high resolution, good durability, high flexibility and ease of fabrication. However, there are still drawbacks in plasmonic colors. The intrinsic ohmic loss resulting from free-electron scattering in metals leads to low efficiency and broad resonant peaks in the reflection spectra. This means that there is still huge potential to improve the saturation of the three primary colors.

### 2.2 Colors from all-dielectric metasurfaces

To tackle the limits of high loss in plasmonic structures, all-dielectric metasurfaces with a high refractive index and low loss have been proposed as candidates. The dielectric nanostructures that interact with electromagnetic waves by a displacement current can support not only electric-dipole resonance, but also a comparable magnetic-dipole resonance.\textsuperscript{79} Hence, all-dielectric metasurfaces, especially amorphous silicon (a-Si) nanostructures, have been extensively used to realize a range of phenomenal effects and functionalities in recent years. For instance, a magnetic reflector,\textsuperscript{80} Huygens’ effect,\textsuperscript{81} high-Q Fano resonance\textsuperscript{82} and wavefront control\textsuperscript{83} have been demonstrated in the infrared range. However, in the visible wavelength, dielectric color filters have been shown to be challenging due to the convenient dielectric materials with either high loss (such as a-Si) or relatively low refractive index (such as $\text{Si}_3\text{N}_4$ and $\text{SiO}_2$).\textsuperscript{84,85} To address this limitation, some skillfully designed all-dielectric metasurfaces with highly saturated colors have been proposed. In the following section, we will systematically discuss high-quality dielectric color filters.

For a-Si color filter, there are invariably two aspects limiting the saturation and range of structural colors, which are high loss and substrate effect.\textsuperscript{82} That is, compared with the presence of the substrate, nanostructures can exhibit narrower spectra transitions from zero backward to suppressed forward scattering in free space. Therefore, an Si color filter can perform improved saturation of structural colors by suppressing the substrate effect or reducing material loss. On one hand, as shown in Fig. 3a, a 70 nm-thick $\text{Si}_3\text{N}_4$ membrane as an antireflective layer was coated over the silicon substrate.\textsuperscript{86} The antireflective layer can create a substrateless environment that effectively suspends the Si nanodisks in free space. A direct correlation between the observed reflection and the particles’ backscattering associated with Kerker’s conditions\textsuperscript{87,88} is established based on the $\text{Si}_3\text{N}_4$ index matching layer. The multipolar decomposition in Fig. 3a(iii) indicates that the first Kerker’s condition can be fully met in the dip at longer wavelengths, where reflection can be totally suppressed. The second Kerker’s condition at shorter wavelengths is approximately satisfied since the phase difference is approximately $\pi$ when the magnetic and electric modes hold the same amplitude. As a result, highly saturated colors generated from this metasurface are able to occupy about 120% standard RGB (sRGB) space by mimicking the Si nanostructure in free space in the reflection mode. The color printing in Fig. 3a(iv) intuitively exhibits full-hue and high-saturation colors. On the other hand, reducing loss of a-Si in the visible range is another way to improve the quality of the colors. In this context, hydrogenated amorphous silicon (a-Si:H), which is obtained via thermal annealing a-Si, has a lower extinction coefficient under the premise of maintaining a high refractive index.\textsuperscript{89,90} Compared with polycrystalline silicon (p-Si) with comparable loss, a-Si:H nanostructures have the advantages of low cost,
efficient growth on substrates at relatively low temperatures and compatibility with the complementary metal oxide semiconductor (CMOS) process.\textsuperscript{90,91} Recently, three primary subtractive colors (CMY) with nearly 100\% efficiency have been generated by all-dielectric metasurfaces incorporating a-Si:H nanodisks in transmission mode.\textsuperscript{91}

Besides a-Si and p-Si, monocrystalline silicon (c-Si) has lower loss in the visible wavelength. When the wavelength is fixed at 410 nm, the extinction coefficients for a-Si, p-Si and c-Si are 2.02, 0.752 and 0.269, respectively, implying c-Si is a kind of ideal material for color filters.\textsuperscript{92} Nonetheless, the c-Si nanostructure is difficult to fabricate due to the incompatibility with the CMOS process.\textsuperscript{84,93} Recently, this challenge has been addressed by utilizing electron beam lithography (EBL) to fabricate the c-Si sample. As shown in the fabricated process of c-Si nanoblocks presented in Fig. 3b(ii), 150 nm-thick chemical resist is spin coated on a c-Si layer on a silica substrate. A 30 nm-thick Cr hard mask is deposited by a resist lift-off process after the designed patterns are drawn by EBL. Finally, the Cr mask is removed by immersing in a Cr etching solution.\textsuperscript{94} By using the fabricated c-Si nanoblocks, various brilliant structural colors are realized via tuning of the geometric parameters. Moreover, due to the high refractive index of c-Si, the pixels composed of a single nanoblock can present high-quality three primary colors in Fig. 3b(iii), indicating that this all-dielectric metasurface processes high resolution of about 85 000 dpi.

Although a variety of Si metasurfaces have realized highly saturated colors, the invariably existing loss in Si materials still hampers higher saturation and higher efficiency. As a result, TiO\textsubscript{2} as an alternative dielectric material with negligible loss in the visible range is a superior substitute to improve the purity of structural colors. To date, TiO\textsubscript{2} nanostructures with a large aspect ratio have been successfully fabricated by atomic layer deposition or electron beam evaporation.\textsuperscript{95–97} Due to the fact that the refractive index of TiO\textsubscript{2} is lower than in Si materials, a periodic arranging array of TiO\textsubscript{2} unit cells can be an effective way to enhance Mie resonances. The coupling between photonic crystal radiation and electric dipole resonances will significantly increase efficiency and reduce the bandwidth of the reflected peak;

Fig. 3 Structural colors generated from all-dielectric metasurface. (a(i)–(iv)) a(i) Schematic and a(ii) SEM image of all-dielectric nanostructures consisting of amorphous Si nanodisks on 70 nm Si\textsubscript{3}N\textsubscript{4} layer and SiO\textsubscript{2} substrate to mimicking free space. a(iii) Reflection spectra and multipolar decomposition of scattering cross-section when the diameter and gap size is 130 nm and 70 nm, respectively. a(iv) Color printing pictures before and after annealing. Reproduced from ref. 86, with permission from American Chemical Society. Copyright 2017. (b(i)–(iii)) b(i) Structural geometry of monocrystalline silicon nanostructures over silica substrate. b(ii) Fabricated process flow chart of the all-dielectric metasurface. b(iii) SEM image and measured image of letters “RGB” with three primary colors, respectively. Reproduced from ref. 94, with permission from American Chemical Society. Copyright 2017. (c(i)–(iv)) Schematic diagram of TiO\textsubscript{2} nanopillars arranged in distinct varying periods P\textsubscript{x} and P\textsubscript{y} under c(i) x and c(ii) y polarization states, respectively. Reflection spectra under c(iii) x and c(iv) y polarization states with varying size of P\textsubscript{y} when P\textsubscript{x} is fixed. Reproduced from ref. 98, with permission from Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim, Copyright 2018. (d(i)–(v)) d(i) Configuration and d(ii) SEM image of multi-dielectric nanostructures resorting to stacked SiO\textsubscript{2}, TiO\textsubscript{2} and Si\textsubscript{3}N\textsubscript{4} nanodisks in reflection mode. d(iii) Reflection spectra and d(iv) multipolar decomposition of scattering cross-section of designed structure. The insets present magnetic field profile. d(v) Measured colors generated from all-dielectric nanostructures with changing gap and period. Reproduced from ref. 105, with permission from American Chemical Society. Copyright 2019.
in other words, saturation of visible colors can be improved dramatically. For example, Fig. 3c shows a highly saturated color filter in reflection mode with nearly 100% efficiency based on TiO2 nanopillars.95 Fano resonance resulting from interference between the radiation waves of Mie resonance and directly reflected waves enables the generation of narrow-band spectra. In addition, the designed strategy of dependently varying periods along the x and y directions is utilized to simultaneously tune saturation and hue. By changing the polarization angle from 0° to 90°, three primary colors (RGB) with high saturation can be flexibly transformed, indicating that this color filter is anticipated to be applied in high-density optical data storage, hyperspectral microscopy and security encryption, for example.

Except for the all-dielectric nanostructures mentioned above, Si3N4 guided-mode resonant grating99,100 perovskite nanostructures101,102 and SiGe resonators103,104 have also been studied for structural colors in recent years. In particular, a kind of multi-dielectric nanostructure consisting of stacked SiO2, TiO2 and Si3N4 layers is proposed to realize ultra-highly saturated colors, as shown in Fig. 3d.105 The index matching of SiO2 capping layer between TiO2 layer and air, and Si3N4 bottom layer between silica substrate and TiO2 layer can significantly suppress higher order modes at shorter wavelengths, and further improve the monochromaticity of the reflection spectra (Fig. 3d iii and iv). More than 83% energy of reflected light is confined in reflected colors. Based on this novel design, 171% sRGB space and 127% Adobe RGB space are taken up in the 1931 CIE chromaticity diagram, which is the largest color space generated from nanostructures as far as we know.

In summary, structural colors generated from all-dielectric metasurfaces offer high saturation and high efficiency, which can be extensively used in high-end displays and imaging applications.96,97,98,105 However, additive colors in the transmission mode are still rarely used within all-dielectric nanostructures. The urgent need now is to realize high-quality additive colors in transmission spectra.

### 3. Dynamically tunable structural colors

Having dynamically tunable colors in situ is essential for advanced displays with novel functionalities, such as color printing,
animations and security encryption. However, the tuning of most structural colors generated from plasmonic and dielectric metasurfaces is dependent on changing physical parameters, which is difficult in fabricated processes.\textsuperscript{106,107} To overcome this limitation, several methods have been used to realize dynamic tuning.\textsuperscript{108–110} For instance, changing the relative orientations of resonators to tune different resonant modes, or using phase-change materials, voltage-control coupling and thermal stimulus to tune metasurfaces.\textsuperscript{111–113} Here, four tunable methods of mechanical stretching, chemical reaction, electrical control and photon doping are reviewed.

Mechanical stretching is the most common way to realize dynamic tuning of near-field interaction via varying period of lattice. In stretched technologies, polydimethylsiloxane (PDMS), as an elastomeric matrix, holds the advantages of elasticity, inertia, non-toxicity and non-flammability.\textsuperscript{109,114} In recent years, mechanically stretched devices have been widely used in interactive LEDs,\textsuperscript{115} solar cells\textsuperscript{116} and tunable metasurfaces.\textsuperscript{117} In particular, the full-color tunable filter by means of mechanical stretching is proposed across the entire visible range, as presented in Fig. 4a.\textsuperscript{118} Al nanostructures with high bulk plasma frequency from the ultraviolet to infrared range are patterned over the PDMS substrate via the EBL process. The scattering spectra depending on the excitation and restricted collection angles is highly sensitive to the period of lattice. As a result, when the nanostructure is designed to perform green in the relaxed state, the scattering spectra will red shift and blue shift in the state of stretching along the x and y axes, respectively (Fig. 4a(i)). The mechanically stretched filter is an ideal device to integrate into a micro-electromechanical system (MEMS), such as ultracompact MEMS-based spectrometers.

Chemical transition is another way to realize dynamic tuning of structural colors; namely, tuning resonance via changing components of nanostructures, where Mg material with superior plasmonic properties is a perfect choice in the visible wavelength.\textsuperscript{119,120} In the field of solid-state hydrogen storage, Mg is one of the most promising candidates, because up to 7.6 wt% of hydrogen gas can be absorbed.\textsuperscript{119} Therefore, metallic-state Mg and dielectric-state MgH\textsubscript{2} are able to undergo chemical transition via the processes of hydrogenation and dehydrogenation, respectively. Based on this unique property of catalytic Mg, Fig. 4b shows a dynamic color filter comprising an Mg layer sandwiched between Ti/Pd capping layers and a Ti adhesion layer.\textsuperscript{121} Upon hydrogen exposure, Mg nanoparticles will go through a series of complicated chemical processes, and the fraction of metal Mg will gradually decrease, rendering a dynamic change from plasmonic colors to dielectric colors. Due to the fact that the Mie resonances of MgH\textsubscript{2} nanoparticles are weak in the visible range, the colors can be erased after hydrogenation. More importantly, the erased process is reversible through dehydrogenation in the presence of oxygen (Fig. 4b(iii)). The flexible colors are promising to be used in dynamic color displays with functionalities of tunable colors, encryption and animation.

Besides physical and chemical tuning, electronic control is also a good way to tune colors due to ultra-thin, low-power-consuming and flexible merits.\textsuperscript{122,123} In electronic technology, the phase transition from the nematic to the isotropic state of liquid crystals (LC) can be used to tune resonance under an applied voltage.\textsuperscript{106,124,125} However, the high cost, lack of bistability and complicated fabrication of LC nanostructures still remain a challenge. Hence, several electrochromic materials, such as conducting polymers and transition metal oxides, have been utilized to dynamically tune colors.\textsuperscript{126–128} For instance, Fig. 4c illustrates full-color electrochromic switching by use of polyaniline (PANI) and poly(2,2-dimethyl-3,4-propylenedioxythiophene) (PolyProDOT-Me\textsubscript{2}), which can both be conformally and controllable electrodeposited on Al nanoslits.\textsuperscript{129} To further improve the purity of colors, an Si\textsubscript{3}N\textsubscript{4} waveguide as a buffer layer is set underneath the Al nanoslits to narrow the linewidth of the spectra (Fig. 4c(i)). By applying different voltages, oxidized and reductive PolyProDOT-Me\textsubscript{2} with changing refractive indices will be switched, as shown in Fig. 4c(i), which means switching colors can be realized at ON and OFF states by using different voltages. The electrochromic switching with fast speed and high-contrast colors has promising applications, ranging from catalysis to photovoltaics.

Finally, photon doping can be used to realize dynamically tunable \textit{in situ} structural colors. In addition to the extrinsic structural colors generated from light–matter interaction, the intrinsic emission colors caused by photoluminescence (PL) of the active materials is introduced in the photon-doping method.\textsuperscript{130,131} As shown in the schematic presented in Fig. 4d, methylammonium lead halide perovskite (MAPbBr\textsubscript{3}, where MA = CH\textsubscript{3}NH\textsubscript{3}+) gratings are patterned on a glass substrate via a top-down process.\textsuperscript{132} MAPbBr\textsubscript{3} material as a semiconductor possesses intrinsic emissions generated from the direct bandgap. The intensity of the intrinsic emission colors can be determined by the excitation; that is, the density of the pumping laser. Hence, mixing of the extrinsic structural colors and intrinsic emission colors leads to a new color. Based on Grassmann’s law, changing the intensity of intrinsic colors by use of a pumping laser enables the \textit{in situ} new color to be dynamically tuned. The dynamic tuning is reversible with decreasing pumping intensity. More significantly, the color-transition time is mainly determined by buildup, and ring down time of PL is of the order of nanoseconds, which is orders of magnitude faster than previous studies. The enlarged measured printing in Fig. 4d(iii) shows flexibly dynamic tuning from red to green by changing the intensity of pumping light.

4. Applications

As shown in the previous sections, static and dynamic structural colors have been widely realized in metasurfaces. Because of the ultra-thin, ultracompact, highly durable and environmentally benign properties of the nanostructures, the full-color filters with high saturation, high efficiency and high resolution are promising alternatives to replace conventional pigments in many practical applications. In the following section, displays, encryption and sensing applications based on structural colors are introduced.

To date, high-end and advanced displays have attracted the most attention in the study of structural colors. In Fig. 5a, a
dynamic plasmonic nanostructure based on Au/Ag nanodomes is integrated into a designed matrix display covering the entire visible range. The method of electrodeposition can be utilized to dynamically alter the thickness of Ag shells on Au nanodomes by applying a voltage, and rendering a continuous color change based on a reversible redox reaction of Ag. During the process of redox, the colors will gradually change with the time of the oxidation/reduction reaction, which is consistent with the cyclic voltammogram measurement. In the design of a biomimetic chameleon (Fig. 5b(ii)), the chameleon covered with plasmonic nanodomes is equipped with color sensors to sense the environment's color. And then the color information is automatically analyzed and delivered into the nanodomes so as to change the color of the chameleon accordingly. This implies that optical wearable devices are possible. In Fig. 5a(iii), rapid display with the responsive speed of a unit cell and static display, where the unit cells are operated by a line-by-line matrix-addressing technique, are respectively demonstrated, showing the potential of high-end displays and imaging applications.

Fig. 5 shows a dynamic cryptographic nanostructure based on asymmetric a-Si:H nanoantennas on a silica substrate. Kerker's conditions can be fully met, and lead to a narrowed resonant peak and suppression of unwanted light; that is, improved saturation of structural colors. More importantly, the polarization-dependent property resulting from the asymmetric structures gives rise to the possibility of storing different optical information in distinct polarization states. As presented in Fig. 5b(i), a fixed nanostructure shows different quickmark codes in 0° and 90° polarization angles. Hence, the nanostructure can be a possible design strategy to realize cryptographic nanoprints via controlling polarization angles, such as the test of cryptograph shown in Fig. 5b(ii). Four kinds of pixel with 0° and 90° rotated angles for ‘‘R, O, A’’ and 45° and 135° rotated angles for ‘‘H, L, B’’ are used to encrypt information in designated polarization angles. These cryptographic nanoprints can be compatible with a smartphone, security tag, payment, identification and anti-counterfeiting.

Moreover, colorimetric sensing based on plasmonic structural colors has been introduced to detect analytes by directly observing changes of visible colors. However, the broad resonant spectra caused by localized surface plasmon resonance (LSPR) generally leads to reduced colorimetric resolution due to the strong radiative damping. For example, the highest LSPR sensitivity of an Au sensor is focused on the near-infrared range. To overcome the...
limitation of LSPR sensing, a skillfully designed strategy is used to improve colorimetric capacity. For instance, 2D Al nanoclusters have been designed to provide Fano resonance with energies in the region from blue to green, which is the most sensitive part for human eyes.\textsuperscript{138} The detected colors can gradually change from blue to orange when the analytes are air ($n = 1.0$), PDMS ($n = 1.4$), poly(methyl methacrylate) (PMMA) ($n = 1.5$) and diphenyl ether ($n = 1.6$), respectively. Furthermore, in the field of biosensing, quantitative colorimetric sensing of ricin is also reached in Fig. 5c based on nanopin-cavity resonator,\textsuperscript{139} by which bright-field structural colors are fully realized (Fig. 5c(ii)). The location of the resonant peak in the visible range can be flexibly tuned by changing the medium indices. As presented in Fig. 5c(iii), an antibody-functionalized biosensor with high sensitivity and fast response is fabricated to detect ricin. The antibody protein can be immobilized by use of the amide linkage formed by the reaction of the terminal succinimide group with the amine of protein. Based on the biosensor, the density of ricin can be observed by a dramatic color change at less than 10 min.

5. Summary and prospects
We have reviewed the development of structural colors based on plasmonic and dielectric metasurfaces to date. The plasmonic additive and subtractive colors caused by SPP resonance have been widely realized via nanogratings, nanoholes, disk-hole coupling and MIM nanostructures in both transmission and reflection modes. Although the inherent loss of metal materials hinders higher saturation and efficiency, remarkably high resolution approaching 100 000 dpi is possible by use of disk-hole coupling. While for dielectric colors, Mie resonances generated from high-index dielectric nanostructures lead to high-quality reflection spectra with narrow linewidth and high efficiency; namely highly saturated colors. Significantly, a multi-dielectric metasurface based on multi-polar modulation has recently obtained a huge color space occupied 171% sRGB space. However, there is still the challenge of realizing additive colors in the transmission mode within all-dielectric metasurfaces. Based on these color filters, a number of methods of dynamically tunable colors, including mechanical stretching, chemical reaction, electrical control and photon doping, are systematically discussed in this paper. The ultra-thin thickness of the color filters has the potential to integrate on a chip, indicating that structural colors generated from metasurfaces are promising in terms of a new paradigm for practical applications; for example, biosensing, colorimetric sensing, high-end displays, encryption, anti-counterfeiting, high-density optical data storage, holograms and so on. In summary, there is great significance and enormous potential application in the study of structural colors.

However, looking ahead, there are still obstacles that need to be addressed in the development of practical color filters. First of all, high-quality additive colors with both high saturation and high resolution are rare in the transmission mode, which is of significant importance for holograms and advanced imaging applications. In addition, color filters need to be combined with practical applications as much as possible. Mass production is still a challenge since Ag and Au are too expensive within plasmonic nanostructures, and TiO\textsubscript{2} and c-Si are not compatible with CMOS process within dielectric nanostructures. Although Al or a-Si seem to be good choices for practical production, high loss invariably hampers the generation of highly saturated colors. That means new designed strategies and fabricated technologies are imminently required in the subsequent research. For instance, hybridization between dielectric and plasmonic materials can be a promising platform for additive structural colors in the transmission mode by meticulous design. On the other hand, possible approaches in terms of nanoimprint lithography, self-assembly technique and injection molding need to be developed in practice to realize mass production. Overall, there is still huge space to be explored in relation to structural colors before commercialization. We believe colors generated from metasurfaces can be extensively used to take the place of chemical pigments, and dramatically improve the quality of our daily life in the future.

Conflicts of interest
There are no conflicts to declare.

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Notes and references


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