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Advanced developments in nonstoichiometric tungsten oxides for electrochromic applications

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Nanoscale tungsten oxides ($\text{WO}_3/\text{W}_{18}\text{O}_{49}$) are a hot research topic because of their impeccable performance in different fields, such as multi-step responses to one stimulus or selective response to a stimulus under controlled conditions, particularly electrochromic performance and energy saving applications in dynamic controllable smart windows. To satisfy different environments, the transparency and thermal insulation of windows are particularly tailored to reduce energy consumption, thereby decreasing cooling and heating costs. In this review, the nonstoichiometric tungsten oxides (WO_{3-x}) are presented in detail, summarized, and discussed, focusing on various methods for the synthesis of $\text{W}_{18}\text{O}_{49}$ and WO_3 , such as chemical vapor deposition (CVD), physical vapor deposition (PVD), sol-gel, solvothermal, and hydrothermal methods, together with the synthetic strategies for the corresponding thin coatings. Then, the advanced strategies to improve the EC performance of nonstoichiometric $\text{W}_{18}\text{O}_{49}$ nanostructures for multifunctionality EC smart windows are discussed. Furthermore, the fundamental mechanisms in coloration/bleaching related to oxygen vacancies, hybridization, elemental doping and surface decoration are explored and discussed in detail. Finally, the key challenges, future developments, and remarks to improve the variability of EC smart windows, such as energy harvesting and energy storing multifunction, coloration enhancement, stability, durability, and reducing toxicity are also highlighted accordingly.

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1. Introduction

The growing energy and global environmental issues require advanced, renewable, and sustainable energy resources, especially energy-efficient functional materials. Smart windows are

an emerging technology that uses chromogenic materials to tailor the visible light transmission and thermal loads in buildings. Smart windows reduce energy consumption due to their astonishing energy harvesting ability and energy savings compared to the commonly used traditional glass. Therefore, according to the technological challenges, the aim of this green innovation is to build large-area, low-cost, durable, and multifunctional electrochromic (EC) smart windows.

It was first demonstrated in 1953 that WO_3 thin films can change color *via* an applied voltage after being immersed in H_2SO_4 .¹ Analogous results were achieved by Deb, at the

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American Cyanamide Corporation in the 1960s² for WO₃-based films, which were subsequently published in 1969 and 1973.^{3,4} This work was motivated by large companies to produce informative displays and liquid crystal display devices.⁵ A similar concept was extended to rear-view mirrors in automobiles and industrialized.⁵ In the 1980s, EC became widely accepted as energy-efficient technology.⁵⁻⁷ In 1984/1985, the name “smart windows” was coined, which became popular, not only to scientists but also soft media and the public. A recent development in chromogenic materials is their energy-saving multi-step application in dynamic switchable smart glasses due to their unique electrochemical properties. A smart window is a dynamically switchable device in which the optical properties of chromogenic materials change upon stimulation, such as electricity, heat, and light, and can be observed. The corresponding technologies are known as photochromic, thermochromic, gasochromic and electrochromic windows, respectively.⁸⁻¹⁰ Among them, photochromic (PC) materials are the most attractive because they change color when exposed to light and are promising in many applications. The optical transmittance in PC materials drops and returns to its original state without stimulation, such as in sunglasses, lenses, and some plastics.¹¹ In contrast to EC, PC films do not need an external power source. Recently, advancements in this area have been made using WO₃ as an active material.¹² However, utilizing WO₃ in PC smart windows is limited because the switching largely depends on the light intensity and cannot be controlled by the user.¹¹ However, this drawback can be overcome by hybrid photo-electrochromic materials that spontaneously work when exposed to sunlight

and also change color or bleached with a supplied electric potential.^{11,13,14}

Thermochromic (TC) materials can respond to changes in temperature with a clear and dark color. TC smart glasses display the automated capability of an optical transmission/reflection with variations in temperature. Operationally, it cannot be a personal preference. Nonetheless, for smart functions, no additional energy is required. Many polymers and small molecules can be activated with thermally dependent reversible reactions or temperature-driven structural rearrangements.¹⁵ The most representative material in the group of TC materials is vanadium oxide (VO₂) or VO₂-based composite materials, which allows the transmission of less heat energy at high temperatures than at low temperatures, and *vice versa*.¹⁶⁻²³ An ultra-transparent coating composed of template-free tunable VO₂ hierarchical and honeycomb-like structures showed excellent visible light transparency up to 95.4% at 700 nm with a solar modulation ability of 5.5%.²⁰ The obtained ultra-transparent honeycomb-like V₂O₅/VO₂ film exhibited an enhanced anti-reflective and transmittance performance, which was demonstrated both theoretically and experimentally. The device constructed based on the ultra-transparent honeycomb-like VO₂ coating film has potential for energy-saving applications as smart windows. Similarly, vertically aligned VO₂ hierarchical nanoplate structures grown on hollow spheroids without any template greatly enhanced the optical properties of VO₂ film(s).²² The visible light transmission and solar heat modulation were recorded to be 50% and 11.2%, respectively, with a contrast transmittance of 60 °C at 2500 nm. The film thickness was effective in varying the visible transmittance by

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lowering the solar optical modulation, *i.e.*, 87.2%, and solar optical sound of 5.3%. The 2D hierarchical nanoplate structure possibly promotes the optical properties for smart windows.

Gasochromic (GC) windows aroused wide interest at the beginning of the 21st century due to their simple, inexpensive layer configuration, and high solar transmittance. The synthesis, characterization, and electrochemical activation of WO₃ thin films used as cathodes in EC devices have also attracted interest in gasochromic applications.²⁴ WO₃-based thin films are doped or decorated with a catalyst such as Pd or Pt nanoparticles. These films are transparent in the visible spectral range; however, upon exposure to H₂ gas, the Pt and Pd nanoparticles in these systems promote the room temperature dissociation of the H₂ molecules into hydrogen atoms, which upon reaction with the tungsten oxide support, provoke the coloration (*i.e.*, blue color) of the oxide film.²⁴ The dissociated H⁺ ions and electrons spillover, and then these ion–electron pairs transfer to the WO₃ layer and cause changes in the optical transmittance based on the small polaron transition mechanism.²⁵ However, repeated coloring-bleaching cycles and long-term charge storing usually lead to structural dysfunctions and catalyst poisoning, which increase the switchable time and result in partial gasochromic irreversibility.²⁶ The coloration in a photoemission experiment with a Pt/WO₃ system was observed, where Pt nanoparticle-decorated WO₃ thin films were prepared by magnetron sputtering at an oblique angle and exposed to H₂ gas *in situ* by near-ambient photoemission spectroscopy (NAPP).²⁴ The formation of W^{*n*+} (*n* < 6) species, which are associated with oxygen network vacancies, is believed to be responsible for the blue color when exposed to hydrogen. Given that magnesium (Mg) has a high hydrogen storage capacity, which tends to be further improved by hybridizing with various metals and oxides, it has been widely used in H₂ storage and switchable mirrors.²⁷ Due to the excellent catalytic property of Nb₂O₅ for the magnesium-hydrogen reaction, fluorocarbon (FC)/Pd/Mg–Nb₂O₅ switchable films fabricated *via* magnetron sputtering and low-temperature inductively coupled plasma chemical vapor deposition technology have demonstrated.²⁸ FC/Pd/Mg–Nb₂O₅ films exhibit a high dynamic range of luminous properties, excellent optical modulation, and fast H₂ adsorption/desorption performance. Similarly, Mg–TiO₂ thin films obtained *via* magnetron co-sputtering can act as the switchable layer in smart windows. Thus, gasochromic switchable mirrors based on Pd/Mg–TiO₂ films fabricated *via* magnetron sputtering demonstrate excellent optical properties, microstructures, and strict structure–function relationships.²⁹

EC materials^{30,31} have been extensively studied to adjust the transmittance, absorbance or reflectance of visible light and solar heat for energy-saving EC smart windows, information displays, self-dimming rare mirrors for automobiles and aircraft, electronic (e-)papers, electronic (e-)skins, *etc.*^{5,6,32–39} To satisfy different environments, the transparency and thermal insulation of windows are particularly tailored to reduce energy consumption, thereby decreasing cooling and heating costs. For this purpose, different types of EC materials have been designed and implemented in EC windows, including tin-doped indium

oxide (ITO), aluminum-doped zinc oxide, and especially WO₃ nanoparticles, which are electrochromic toward NIR light.^{40–43} Hybridizing WO₃ with traditional transition metal oxides/organic electrochromics for visible light modulation makes it possible to construct dual/multifunction capacities that can independently modulate visible and NIR light. Polymer-based EC materials targeting visible light offer several potential advantages with good absorption properties in a wide range of colors, fast switching speed, and high coloration efficiency, which are usually difficult to achieve with metal oxides.^{43–45} Based on the design of the donor–acceptor (D–A) structure,^{46,47} EC polymers with a low onset potential and high reversible-switching stability including conjugated polymers,^{44,48–50} unconjugated polymers,⁵¹ and cross-conjugated polymers have been demonstrated.^{52–54} Poly(3-hexylthiophene) (P3HT) is a regioregular semiconducting polymer that shows EC modulation under a small electrical bias polarity and acts as an anodically coloring material.⁵⁵ Given that WO₃ is a well-known cathodically coloring material, a hybrid composed of P3HT and WO₃ as two active electrochromics demonstrated a dual-band EC performance.⁵⁵ In the dual EC device, P3HT was used as the anode coloring material and WO₃ as the cathode coloring material, and therefore, the transfer of charges from WO₃ to P3HT makes it colored, whereas the colors of both materials were bleached in the opposite direction.⁵⁵

To achieve multiple colors and enhance the functionality of all-in-one-type EC devices, the incorporation of several EC chromophores is inevitable.⁵⁶ They can achieve the simplest device configuration because of their facile fabrication using versatile solution processes (*e.g.*, spin coating and printing) and easily tunable device characteristics such as transmittance contrast, color, operating voltage, and cyclic stability by varying the concentration of the chromophores^{57,58} and tuning the electrochemical properties⁵⁹ or molecular structures^{60,61} of the redox species. Accordingly, 1,1'-disubstituted-4,4'-bipyridinium salts, which are known as viologens, have been extensively studied because their optical and electrochemical properties can be easily tuned by modifying their substituents. The coloration potential (*i.e.*, reduction potential) of viologens depends on the color. Therefore, uneven color contrast is obtained when a fixed voltage is applied to a device including multiple viologens.^{62,63} However, self-bleaching occurs even under open-circuit conditions, which is considered to be a weakness of all-in-one EC devices. Polymeric viologens (poly-viologens) have been demonstrated to lower the diffusivity of EC chromophores and minimize self-bleaching.⁶⁴ Compared with EC devices based on mono-viologens corresponding to the monomer of poly-viologens, EC devices containing poly-viologens exhibit the advantages of lower coloration voltage (–0.55 V) and higher coloration/bleaching cyclic stability (> 1500 cycles). In particular, as-designed poly-viologen ECDs show remarkably reduced self-bleaching, resulting in extremely low power consumption (~8.3 μW cm^{–2}) to maintain the colored state.⁶⁴ Moreover, compared with common glass, poly-viologen-based EC devices have strong NIR ray absorption capacity, which proves their feasibility as effective heat shutters at low-power consumption.⁶⁴ EC smart windows utilizing these



composite materials can simultaneously control building lighting and heating.

In the OFF state, EC devices are transparent to natural light. When a suitable voltage is applied, electrons are transferred due to the oxidation–reduction reaction, and the EC materials become colored with an optical modulation. Thus, by varying the voltage applied across electrodes modified with EC materials or their composites, they can be switched between “bright and warm”, “bright and cool”, and “dark and cool” modes. Consequently, at a lower potential, windows are in the cooling mode, blocking heat flux but allowing UV light, whereas setting smaller potential switches to dark mode limits both heat rays and UV light. These functional modes facilitate the multipurpose use of windows as follows: (i) operate under different weather conditions particularly with personal preference, (ii) provide better comfort for occupants, and (iii) features associated with energy-storage/harvesting, aseptic/sterile function, aesthetics, privacy/security, high-latitude zones, aircraft, *etc.*^{5,32,33,65,66}

Energy efficiency in buildings has become a focus, given that 40% of the global energy consumption comes from the urban environment.^{67,68} Due to the rapid increase in global warming, building cooling is becoming a key problem because every 1 °C rise in temperature leads to a 20% increase in energy consumption.⁶⁷ Building cooling systems in developed countries account for nearly 50–60% of the total building energy consumption.^{8,67} The concept of a “zero-energy building” has emerged to control the thermal load *via* the dynamic optical modulation of “smart” windows.⁶⁹ Moreover, these Wi-Fi-connected tint-controllable smart glass surfaces are equipped with environmental sensors that can read room occupancy, weather, and sunlight. The emerging smart windows installed in commercial and domestic areas can save ~170 kW h, *i.e.*, 30% of the total energy consumption annually, which is an important step towards achieving “zero energy buildings”.^{8,67} The other recent approaches based on energy-saving EC materials and technologies for novel multifunction are published elsewhere.^{67,68,70–78} Modern commercial buildings in European countries have installed functioning EC smart windows instead of active air conditioning, and approximately 4.5% energy can be saved annually in the USA and Australia.⁷⁹ Similarly, auto-dimming smart glass and automotive rear-view mirrors have been commercialized.⁸⁰

Nonstoichiometric metal oxides such as WO_{3-x} , MoO_{3-x} , TiO_x , NbO_x , NiO_x , V_xO_y , MnO_x , and Cu_xS_y have strong light absorption/reflection and thermal insulation properties. In particular, WO_{3-x} -based materials in energy-saving EC devices have application prospects because (i) they possess unusual defects and enormous oxygen vacancies, thus achieving strong adsorption affinity with electrolyte ions, (ii) oxygen vacancies provide free electrons, which increase the electric conductivity, resulting in quick electron transfer from the surface, (iii) strong local surface plasmonic resonance (LSPR) provides strong photoabsorption over a wide range of the NIR region, and (iv) a significant reduction in bandgap results in strong light absorptivity, generating electron–hole pairs.^{81–86} Because of

their unique channel/tunnel-like structures for easy electrolyte ion intercalation, defective structure, low cost, high natural abundance, excellent thermal stability, good compatibility, and high theoretical capacitance, oxygen-deficient $\text{W}_{18}\text{O}_{49}$ nanostructures are attractive for application in various chromogenic devices, particularly in EC smart windows.^{82,83,87,88}

The scope of this review is focused on the recent advancements in the synthesis of $\text{WO}_{3-x}/\text{WO}_3$ nanostructures, EC smart coatings, and future advances in smart windows. Sections 1, 2 and 3 briefly introduce the basic properties and EC mechanism of sub-stoichiometric WO_{3-x} . Section 4 provides an overview of the synthetic methods, film formation, and EC properties. Section 5 discusses the strategies for improving the electrochromism of tungsten oxide, including hybridization, elemental/plasmonic doping, and surface decoration. Section 6 describes the all-in-one solid-state next-generation multifunctional EC window principle and its configuration as a typical example for realization, followed by the conclusions, existing and prospective challenges, and finally future demands.

2. Electrochromism

EC materials coated on a transparent conducting electrode can change their optical properties reversibly and continuously by applying a small electric potential. Thus, the transport of ions can occur between the EC material film(s), ion storage/conducting film(s), and/or ion conducting electrolyte(s). The charges are balanced by intercalated/deintercalated electrons, and then transported through the transparent conductors. Smart films can maintain their physical properties with reversible optical modulation in the open circuit, and their color can be refreshed to a specific intermediate state. If energy storage devices can sense energy changes in predictable patterns, we can quickly determine that energy is running out before these devices stop working, demonstrating a wide range of potentially smart applications. Moreover, energy can be saved by using the energy stored in EC devices. If we need to color an EC device to resist sunshine/protect privacy, charges can be stored inside. If the coloration is no longer needed, the stored energy can be released through an external circuit, providing another way for energy resource utilization.

As mentioned, EC films and the complementary counter electrode may have similar optical properties. In a hybrid dual-EC device, two complementary electrodes (working and counter) coated with switchable EC materials have sufficient charge capacity for ion storage, as shown in Fig. 1. These EC devices are promising because the redox reaction that occurs in the active layer is fully compensated and can complete the color changes, resulting in an electrochemically stable battery-type cell. Therefore, EC materials can be (i) cathodic EC materials and (ii) anodic EC materials. Some EC materials that can achieve color from transparent to dark colors by ion-insertion/reduction during cathodic polarization are called cathodic EC materials. Similarly, some are anodic electrochromics, which can achieve a dark color/opaque state from a



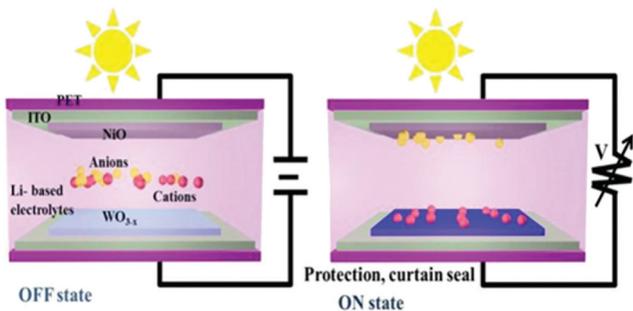


Fig. 1 Simplified EC smart window illustration in the colored and bleached state. Electron intercalation through an external circuit into the EC materials (cathode and/or anode), while ions flow through the electrolyte to compensate for the electronic charges. The panel shows sealing with a curtain for protection.

transparent state under ion-extraction/oxidation during the anodic polarization process. The key variables for evaluating the performance of EC materials are as follows: (i) optical modulation, EC materials should carry excellent optical modulation (100%), *i.e.*, fully transparent in the bleached state and fully opaque in the dark state. (ii) Switching speed, EC materials should reach 90% optical modulation between the coloration and bleaching states. (iii) Coloration efficiency is the key index to evaluate the EC performance. A high coloration efficiency ensures that materials exhibit large optical modulation with low power consumption. (iv) The cycling stability and reliability of EC materials should be maintained for more than 50 years without degradation during use.

Many inorganic materials such as TiO_2 , WO_3 , MoO_2 , VO_2 , Nb_2O_5 , Ta_2O_5 , CeO_2 , NiO , Co_3O_4 , and SnO_2 ,^{89–92} organics such as Prussian blue; and small organic molecules^{93–97} including conjugated polymers,^{96–99} transition metal complexes,^{34,40,79,100–104} viologens¹⁰⁵ and conducting polymers^{106–109} have been examined recently. The two dominant EC materials are polymers and metal oxides.¹⁰ Polymers are excellent candidates for bendable, foldable, and stretchable applications because of their flexible molecular framework interjunctions.^{10,110,111} Metal oxides possess ultra-small particle sizes with a large specific surface area and are expected to exhibit excellent EC functionality, possibly due to their short ion diffusion length, easy electrolyte accessibility, excellent coloration efficiency, cyclic stability, and durability. However, metal oxide-based films also have certain disadvantages, such as cracking and degradation, after continuous bending and stretching.

Organic EC materials exhibit controllable color, higher coloration efficiency, and polyelectrochromism similar to that of inorganic EC materials. Particularly, the well-known organic EC materials are organic small molecules, especially viologens, which are the major candidates used in EC display devices. Specifically, viologens are 1,1'-disubstituted-4,4'-dipyridinium salts, with two positive charges, which are colorless in the normal state (V^{2+}), deep-blue in the radical cation state ($\text{V}^{\cdot+}$) and reversible.³⁹ When the *N*-substituents are short alkyl chains (*e.g.*, methyl or ethyl), the colored state including the viologen cation radical shows a blue color. As the alkyl length increases

(*e.g.*, heptyl) and the electrolyte has hydrophilic character, the colored species tend to dimerize. Consequently, the colored state is magenta, but the quasi-reversible oxidation of the dimerized viologen cation radical degrades the device performance. Various aryl-substituents have been also reported to obtain other colors such as green. Based on the numerous viologens, multicolor ECDs with EC ion gels have been demonstrated.¹¹² The EC properties of conjugated conducting polymers such as polythiophenes (PT), polydimethylsiloxane (PDMS), poly(vinyl alcohol) (PVA), poly(methyl methacrylate) (PMMA), polyaniline (PANI), polyimide (PI), polypyrroles (PPy), polycarbazoles (PCz), and polyfurans (PF) vary with their molecular structure and bandgap energy. The coloration may be due to the donor moiety, which accelerates the π - π^* electronic transition. The HOMO energy level is at a higher energy state. In contrast, the acceptor moiety lowers the LUMO energy level, promoting the charge transfer between the two types of units within the backbone of the polymer in the electrolyte upon electrochemical oxidation and reduction.¹¹³ PT, PANI, and PPy have gained intense research interest due to their facile synthetic route, low redox potential and suitable bandgap. The electrochromism in conjugated polymers is tunable with changes in their bandgap, monomer type, conjugation length, side-chain/substitution groups, steric effects, and stereoregularity. Because of their unique absorptivity and redox reactions in the visible range, they change their electronic structure, optical states, and conductivity under an applied electric current. Metal-organic frameworks (MOFs) are porous materials, and their permanent porosity offers channels for the fast transport of ions, thus enhancing their EC properties.¹³ However, there are certain challenges^{9,39,110} that limit the large-scale preparation of organic EC materials as follows:

- (i) their multistep synthesis and purification processes,
- (ii) use of hazardous organic reagents,
- (iii) their photo-electrochemical instability,
- (iv) their optical contrast fade-out caused by overoxidation, and
- (v) wide range optical modulation in the visible region and lack of control over NIR light.

Recent interest in flexible, stretchable, and foldable devices has been focused on incorporating EC metal oxides in polymers to prepare hybridized materials by carefully matching their refractive index and surface chemistry. The resulting hybrid composites exhibit improved chemical and mechanical properties, enhanced dispersion of nanoparticles, stimuli-responsiveness and provide excellent bendability, stretchability, and foldability.^{10,39,83,86,98,114} These intrinsic stretchable and transparent conductive electrochromics combine to balance the optical transparency, conductivity, and mechanical stretchability. Focusing on the explosive growth of metal oxides towards EC applications, tungsten, titanium, niobium, vanadium, molybdenum, nickel, copper, manganese, cobalt, and iridium oxides are highly addressed based on their preparation, stoichiometry, particle morphology, composition, and deposition process. Recently, $\text{WO}_3/\text{WO}_{3-x}$ has been demonstrated to be an ideal material owing to its unique EC, electrochemical, and optical properties. The unique detail of this review is the



Table 1 Advanced EC materials and their mechanisms and merits towards EC smart glass, evaluated for the desired “zero-energy buildings” “smart windows”

EC material	EC mechanism and properties	Coloration efficiency (cm ² C ⁻¹)	Transmittance contrast	Switching speed (color/bleach (s)) and cycle life (cycles)	Ref.
WO _{3-x} (amorphous)	Ion insertion, polaron absorption, charge transfer, and Vis and NIR absorption	125–80	56–70% at 550–800 nm	5 s/2.5 s, 1000 to 4000	115–118
WO _{2.72} assembled on Ag nanowires	Ion insertion, polaron absorption, charge transfer, and Vis and NIR absorption	35.7	58–86% at 550 nm	2 s/4 s, 1000	119
Mesoporous WO ₃	Ion intercalation, polaron absorption, charge storage, Vis NIR absorption	188	76% at 700 nm	0.8 s/0.4 s 1000	120
WO ₃ (crystalline)	Ion intercalation, polaron absorption, charge transfer, free carrier, and Vis and NIR absorption	42	50–70% at 550 nm	Several min, 3000	101
WO ₃ ·2H ₂ O nanoplates	Pseudocapacitive, polaron absorption, charge transfer, and NIR absorption with transmittance modulation (90.4%)	322.6	63.8% at 550 nm and 90.4% at 1600 nm	93.7% after 500 cycles	121
WO _{3-x} /POMs	Ion insertion, polaron absorption, charge transfer, and Vis, NIR selective absorption	42.31	64% at 630 nm and 88% at 915 nm	26/86, 500	122
WO _{2.72} -NbO _x on ITO-PET	Ion insertion, polaron absorption, charge transfer, capacitive, and Vis and NIR selective transmission	30	—	Mint, 2000	123
MoO ₃ -WO ₃ /Ag/MoO ₃ -WO ₃	Ion diffusion, charge transfer, and Vis and NIR selective transmission	70	72.9–79.3 at 400–800 nm	2.7/4.1, 500	124
WO ₃ /Ag/WO ₃	Ion diffusion, charge transfer, and Vis and NIR selective transmission	30.5	52.5–82.7 at 400–780 nm	15/22.4, 2500	125
NiO	Pseudocapacitive (anodic) polaron absorption, and charge transfer	30–100	68% at 650 nm	53, 10 000	117
Smart window for zero energy building	Capacitive, sensing, plasmon absorption + polaron/charge transfer, and Vis and NIR-independent absorption + reduce toxicity	Infinite	More than 75% at 550–630 nm and More than 95% at IR region	Seconds/less Greater than 50 000 Stand for over 50 years	?

prospective advancement towards next-generation smart windows using W₁₈O₄₉/WO_{2.72} among the metal oxide (Table 1).

3. Nonstoichiometric WO_{3-x}: properties and electrochromic mechanism

3.1. Properties

Tungsten oxide (WO₃/WO_{3-x}/WO₃·xH₂O) is a type of perovskite structure formed of WO₆ octahedra connected by W–O–W bonds or water molecule(s) similar to the geometry of rhenium oxide (ReO₂). Its electronic structure is composed of oxygen with a filled 2p orbital (valence band) and tungsten with an empty 5d orbital (conduction band).^{126,127} The WO₆ octahedron can be interpreted as a central tungsten metal surrounded by six equidistant oxygen atoms, sharing edges or corners, thus forming a 3D-octahedral crystal structure.¹²⁸ In the octahedral geometry, there are hexagonal tunnels and trigonal windows (Fig. 3), and the corner sites facilitate the transport of ions. One or two charge-compensated electrons are transferred to the adjacent W-atoms, reducing them to W⁵⁺ ions.¹²⁹ The charge-compensated electrons can be transferred to two neighboring W ions to form a doubly charged W⁴⁺ and transition between W⁴⁺ to W⁵⁺.^{129–131} The crystal lattice of WO_x contains considerable oxygen deficiency sites with partially reduced W⁵⁺ species. These sub-oxides with substantial oxygen vacancies become more metalized with an increase in non-stoichiometry and achieve high conductivity. The WO₆ octahedral symmetry

undergoes tilting and rotation with a change in temperature,¹³² forming typical monoclinic, hexagonal, orthorhombic and tetragonal crystal symmetries, indicating that the WO₃ crystal phases are partially reversible. At room temperature (25 °C), WO₃ crystallizes in the triclinic phase, which transforms into other phases upon annealing,¹³² such as monoclinic II (ε-WO₃, < -43 °C) → triclinic (δ-WO₃, -43 °C to 17 °C) → monoclinic I (γ-WO₃, 17 °C to 400 °C) → orthorhombic (β-WO₃, 400 °C to 740 °C) → tetragonal (α-WO₃, > 740 °C), as shown in Fig. 2. The obtained crystal structures are orthorhombic for W₃O₈ and W₃₂O₈₄, monoclinic for W₁₇O₄₇, W₁₈O₄₉, W₂₀O₅₈, and W₂₅O₇₃, and tetragonal for W₅O₁₄. The reduction reactions of monoclinic WO₃ (γ-WO₃) forms Magneli phases. Monoclinic WO₃ is the most stable phase studied as EC films in EC devices. In this reduction process, with an increase in oxygen vacancies, the position of the WO₆ octahedra in the crystal structure changes from corner-sharing to edge-sharing, which split by well-ordered crystallographic shear planes.¹³² In WO_{3-x}, the edge-sharing WO₆ octahedra with channels arrange to form hexagonal and pentagonal tunnels.

The nonstoichiometric WO_{3-x} was first demonstrated by Glemser and Sauer, where they found that the pure stoichiometric WO₃ phase structure could be converted to WO_{2.9} by introducing oxygen vacancies.¹³³ Therefore, the WO_{3-x} lattice can exhibit a large number of oxygen vacancies, resulting in a variety of non-stoichiometric compounds.^{79,134} Recent research has focused on the synthesis of sub-stoichiometric tungsten oxide (WO_{3-x}, 0 < x < 1), including W₂₀O₅₈ (x = 2.90), W₁₉O₅₅ (x = 2.89), W₅O₁₄ (x = 2.80), W₁₈O₄₉ (x = 2.72), and



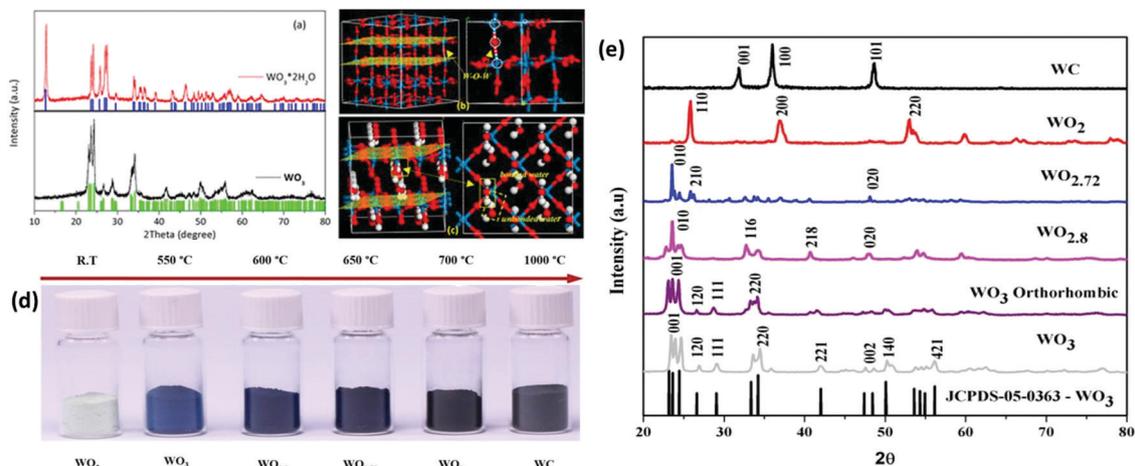


Fig. 2 (a) XRD patterns and crystal structures of (b) WO_3 and (c) $\text{WO}_3 \cdot 2\text{H}_2\text{O}$ (W, blue; O, red; and H, white).¹⁵⁷ (d) Variations in the color of tungsten oxide powders prepared via reduction under a CO atmosphere at different temperatures. (e) XRD patterns of pure tungsten trioxide before reduction and its sub-oxides after reduction under a CO atmosphere. Reproduced with permission.¹⁵⁸ Copyright©2017, MDPI.

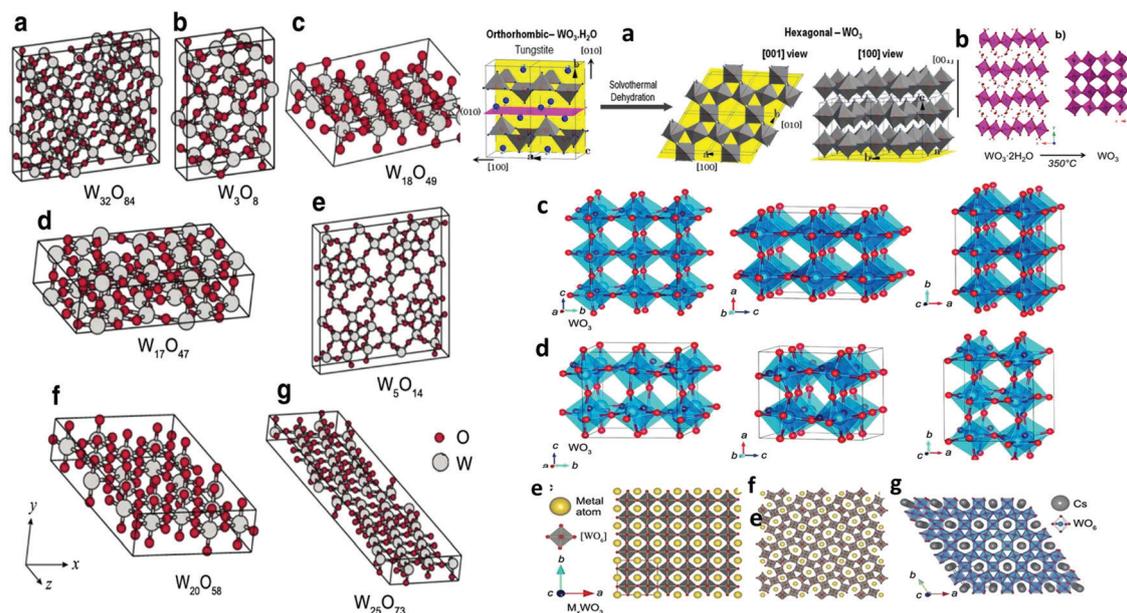


Fig. 3 Left: (a–g) Unit cells of various WO_{3-x} . The large gray and small red balls represent W and O atoms, respectively. Reprinted with permission.¹⁶⁵ Copyright©2010, AIP. Right: (a) Schematic of the transition mechanism from orthorhombic $\text{WO}_3 \cdot \text{H}_2\text{O}$ to hexagonal WO_3 , (b) crystallographic structure of monoclinic WO_3 from layered hydrated WO_3 via heat treatment process: hydrated WO_3 ($\text{WO}_3 \cdot 2\text{H}_2\text{O}$) to monoclinic (γ) WO_3 . Reproduced with permission.¹⁶¹ Copyright©2017, ACS. (c) and (d) Unit cell orientations of orthorhombic and monoclinic WO_3 along the a, b, and c planes,¹⁶⁶ and tungsten bronze ($M_x\text{WO}_3$) (e) cubic ($x = 1$), (f) tetragonal ($1 > x > 0.33$),^{167,168} and (g) hexagonal ($x < 0.33$) structures.¹⁶⁸

$\text{W}_{32}\text{O}_{84}$ ($x = 2.62$).¹³⁵ $\text{W}_{18}\text{O}_{49}$ is an n -type material semiconductor with a bandgap of 2.6–3.5 eV. The formation of oxygen vacancies in the WO_x lattice affects the Fermi level of tungsten oxide, narrows the bandgap, and increases the conductivity and free carrier electrons.^{79,133,136} These materials are usually insoluble in all solvents and can be oxidized to stoichiometric WO_3 after a long oxidation reaction. Decreasing the particle size increases the bandgap and results in a blue shift and photon absorption.^{88,136}

WO_{3-x} is an interesting LSPR candidate host due to its distinctive properties with outer d-valence electrons. The new discrete energy bands beneath the conduction band (Fig. 4b) are produced by oxygen vacancies.^{81,137,138} Due to the presence of oxygen vacancies, crystal defects, or insertion of tiny metal ions, strong LSPR with energy comparable to the bandgap energy can induce significant near-infrared absorption properties. The LSPR peak in plasmonic nanoparticles is directly proportional to the square root of the density of free electrons



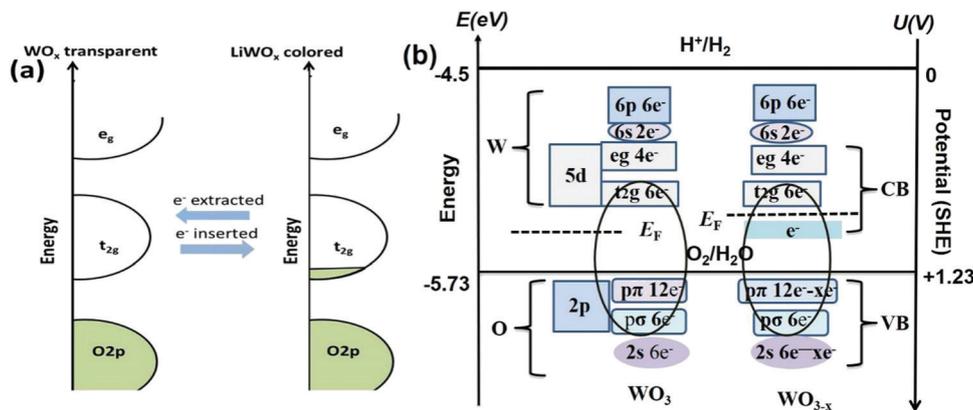


Fig. 4 Schematic illustration of tungsten oxide in the colored and bleached state. (b) Sketch of the electronic states in WO_3 and WO_{3-x} on the y-axis to the left in energy (eV) and the right in potential according to the standard hydrogen potential (SHE) under vacuum. Reproduced with permission.¹⁸⁹ Copyright©2017, Elsevier.

in the particles.¹³⁹ The LSPR depends on certain factors, such as the particle size, dimensions, structure, shape, and dielectric properties of metals.¹⁴⁰ Controlled heat treatment in a weak reducing environment is a facile but effective way to increase the number of oxygen vacancies. The dopants/desired impurities donate their electrons/holes and increase the density of free electron(s) in the conduction band. Meanwhile, the LSPR generates a rigorous local electric field, promoting the photoelectric harvesting application, which has broad application prospects for smart functions in EC windows.¹⁴¹

The monoclinic $\text{W}_{18}\text{O}_{49}$ has excellent stability and can be recovered in the pure phase with substantial oxygen vacancies. $\text{W}_{18}\text{O}_{49}$ possesses both the octahedral and pentagonal bipyramidal coordination links of the W with O atoms. Similar to h-WO_3 , ordered edge-sharing WO_6 networks produce three different tunnels/cavities on a crystal plane, such as hexagonal, trigonal, and quadrilateral, to accommodate guest ions during the electrochemical process (Fig. 3).¹⁴² Due to its astonishing defect structure and remarkable physicochemical properties, $\text{W}_{18}\text{O}_{49}$ has become a potential candidate for application in solar cells,¹⁴³ photocatalysis,^{144–147} chromogenic devices,^{119,139} photothermal agents for cancer treatment,¹⁴⁸ supercapacitors and batteries for energy storage,¹³³ and sensors toward various toxic gas molecules because of its distinctive structural faces, ultrathin nature, small particle size, high surface area, reproducible morphologies, crystal defects, and oxygen vacancies.^{149–154} Hexagonal WO_3 (h-WO_3) is a metastable phase that has received significant interest in electrochemical applications as an Li^+ and Na^+ ion host material,¹⁵⁵ and its crystal structure is presented in Fig. 3a. In other polymorphs, the octahedral units are organized into a perovskite-like structure. In hexagonal WO_3 , the WO_6 octahedra existing in the form of three- and six-membered rings in the “*ab*” plane create trigonal cavities and hexagonal windows, respectively, whereas in the “*c*” axis”, sharing of the axial oxygen results in 4-coordinated square windows.¹⁵⁶ These tunnels offer accommodation sites for a large number of guest cations during the electrochemical process. No hexagonal phase was observed during the

structural transformation; however, it can be obtained *via* the dehydration of hydrated tungsten oxides, indicating that the structural transformation can only occur in aqueous media. In addition, it can be converted into a monoclinic phase (Fig. 3b) by annealing at a temperature above 400 °C. The redox reactions of tungsten metal/tungsten oxides in aqueous electrolyte are controlled by various pH-related half-reactions, and possible explanations can be found in the review published by Shinde and Jun.¹³³

Tungsten oxide hydrates ($\text{WO}_3 \cdot x\text{H}_2\text{O}$) also exhibit properties similar to WO_{3-x} nanostructures. They can be classified into subclasses based on the number of water molecules present in their crystal lattices as follows:

- (i) WO_3 -dihydrate ($\text{WO}_3 \cdot 2\text{H}_2\text{O}$),
- (ii) WO_3 -monohydrate ($\text{WO}_3 \cdot \text{H}_2\text{O}$), and
- (iii) WO_3 -hemihydrate ($\text{WO}_3 \cdot 1/2\text{H}_2\text{O}$)/ $\text{WO}_3 \cdot 0.33\text{H}_2\text{O}$.

$\text{WO}_3 \cdot 2\text{H}_2\text{O}$ has a stratified layered structure composed of a $\text{WO}_5(\text{OH})_2$ layer sharing corners and the other water molecule is found in between the layers. Hence, it can be facially exfoliated into thin 2D nanosheets.^{81,159} $\text{WO}_3 \cdot \text{H}_2\text{O}$ is composed of highly distorted $\text{WO}_5(\text{OH})_2$ units sharing corners and is coordinated by five oxygen atoms and one H_2O molecule. As shown in Fig. 3e, it is believed that the crystal structure of $\text{WO}_3 \cdot 1/2\text{H}_2\text{O}$ resembles a cubic pyrochlore type, where the water molecules are present in its tunnels.¹⁶⁰ The inter-layer expansion can significantly reduce the ion diffusion energy barrier, ultimately improving the ion diffusion kinetics. Furthermore, the structural water molecules confined in the nano-interlayer spaces facilitate efficient proton transfer and induce a battery-to-pseudocapacitor transition in electrode materials.^{161,162} The interlayer water molecules can improve the intercalation capability of EC ions, which is significantly important in attaining superb NIR electrochromism in EC devices. Accordingly, Wang *et al.*¹⁶³ demonstrated hydrated WO_3 and studied the influence of water molecules in decreasing the energy barrier of ion diffusion and improving the ionic flux in the WO_3 crystals. They found that $\text{WO}_3 \cdot 2\text{H}_2\text{O}$ exhibited a good NIR extension with a large transmittance modulation (90.4%), high coloration



efficiency ($322.6 \text{ cm}^2 \text{ C}^{-1}$), and high cyclic stability (maintaining 93.7% after 500 cycles), with good solar heat maintenance capability. Li and coworkers¹⁶⁴ reported an EC hybrid material using $\text{WO}_3 \cdot \text{H}_2\text{O}$ nanosheets and a Persian-white film on FTO substrate as an asymmetric electrode. The film had an optical modulation of 61.7%, response speed of 1.84/1.95 s, coloration efficiency of $139.4 \text{ cm}^2 \text{ C}^{-1}$, and cyclic stability of 82%.

The optical properties of WO_{3-x} can be measured by absorbing photon energy, which excites electrons from the valence band (occupied state) to the conduction band (empty state) and generates electron-hole pairs.¹⁶⁹ The absorption coefficient (α) evaluated from the spectral transmittance, $T(\lambda)$, and reflectance, $R(\lambda)$, is given as eqn (1).

$$\alpha(\lambda) = 1/d \ln[(1 - R(\lambda))/(T(\lambda))] \quad (1)$$

Tungsten oxide is transparent in most visible wavelengths when the photon energy is lower than the bandgap energy. However, it shows a yellowish and blue color with a smaller bandgap, representing that some parts of blue light are absorbed.^{170,171} The optical absorption at a lower energy than the bandgap is because of the electronic transition between the localized states.¹⁷² However, when the photon energy is greater than the bandgap energy, the absorption can be described as follows:

$$\varepsilon \propto \infty(\varepsilon - E_g)^\eta \quad (2)$$

where “ ε ” is the photon energy and “ η ” depends on the type of optical transition, for example, if $\eta = 2$ it is an indirect and allowed transition.

Because of the reduction in particle size (quantum confinement effect), the bandgap becomes wide, resulting in light emission at shorter wavelengths.¹⁷³ Nonetheless, the structural defects and oxygen vacancies in WO_{3-x} narrow the bandgap due to the defective band below the conduction band. The high concentration of oxygen vacancies in WO_{3-x} extend the defective band into the conduction band. This forms a degenerate semiconductor, which shifts the Fermi level upward, achieving metal-like conductivity, and thus highly improves the electrochromic, electrochemical, photocatalytic, and photochromic properties.¹⁷¹ The color of WO_3 films changes from light-yellow to light-greenish and WO_{3-x} changes to dark blue, which shows a blue shift. Broad absorption bands appear in the red to NIR region and blue-green region. The absorption bands at these specific wavelengths make $\text{WO}_{2.72}$ light green, where the additional absorption bands are due to the transfer of electrons from W^{6+} to W^{5+} .¹⁷² The excess electrons can be localized, which are known as small polarons, or transferred to adjacent W ions.¹⁷⁴ The intercalated electrons may occupy the empty d-bands and develop a new electronic transition that is delocalized on crystalline WO_3 or localize on the metal sites in an amorphous form and perform polaronic absorption. Amorphous WO_{3-x} exhibits a disordered structure, indicating the presence of broken bonds with variable bond lengths, bond angles, and coordination. In WO_{3-x} , the localized states are found closer to the oxygen vacancies or tungsten atom, and

thus the electrons and/or charge compensating ions can easily jump within the localized states.^{136,169,175}

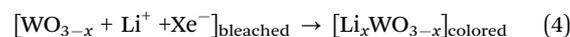
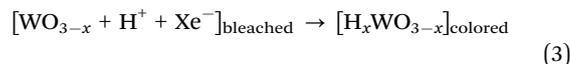
3.2. Electrochromic mechanism

The EC properties mainly depend on the particle size, dimensions, aspect ratio, crystallinity, porosity, thickness, and reversibility of films, together with their capability to accommodate guest ions (H^+ , Na^+ , and Li^+).^{13,88,176} The coloration mechanism depends on the nature and behaviors of materials. In crystalline tungsten oxide films, their coloration is attributed to the Drude-like absorption of delocalized/free electrons, similar to the behavior of heavily doped metal oxides with ionized impurities.¹⁷⁷ In amorphous tungsten oxide films, the optical absorption is caused by the electron exchange/polaron-to-polaron transition between adjacent W atoms (W^{6+} , W^{5+} , and W^{4+} states), and the inserted electrons are localized in the W^{5+} sites and polarize their surrounding lattice to form small polarons.¹⁷⁸ Moreover, compared to crystalline films, amorphous WO_3 films show a better coloration efficiency, which increases with an increase in oxygen vacancies.¹⁷⁹ However, amorphous tungsten oxide films show poor stability because of their irregular atomic distribution.⁸⁸

$\text{W}_{18}\text{O}_{49}$ is a promising EC smart material that changes color from clear to blue, dark-blue, and transparent with a faradaic charge. When protons/lithium ions intercalate and externally powered electrons are inserted into the tungsten oxide film, the transmissivity of the electrodes drops from 91.3% to 15.1% during charging.^{129,180} Two types of redox reactions occur upon voltage bias as follows:¹⁸¹ (a) a charge transfer process at the material-electrolyte interface, *i.e.*, surface-dependent process and (b) diffusion controlled redox reactions within the material lattices, *i.e.*, ion-diffusion dependent process. The intercalated cations exist in the channels formed by the WO_6 octahedra or react with the bridging oxygen of the host¹⁵⁶ by breaking the W–O–W networks and generating new W=O bonds.^{128,182} Thus, the coloration (tungsten bronzes, $\text{H}_x\text{WO}_3/\text{Li}_x\text{WO}_3$ formation) is due to the reaction of cations with W=O groups by forming colored $\text{W}^{5+}\text{-OH}$ states.¹⁸¹ It is assumed that cations can react with the bridging oxygen at a low concentration of inserted charges. However, at a high concentration, they can also react with W–O.^{128,181,183} Raman spectroscopy results show that the interaction between oxygen vacancies and Li^+ ions improves the coloration efficiency of $\text{WO}_{2.72}$ films.^{129,136,179} When the applied voltage polarity is reversed, the Li^+ ions deintercalate and the film is bleached. These redox dynamics demonstrate that tungsten oxide can be transformed from a transparent state to an absorbent state to store electric energy.

The ion insertion and extraction in WO_3 films can be shown as eqn (3)–(5):

Cathodic reaction,



Anodic reaction





Protic electrolytes are used for H^+ intercalation. Similarly, Li^+ electrolytes are widely used in $\text{WO}_{2.72}$ EC devices (Fig. 1).¹⁵⁶ The colored state should be H_xWO_3 or Li_xWO_3 , whereas some reactions reach $X < 0.5$ states, resulting in complete reversibility. The hybridization in the octahedral geometry can be shown between the oxygen 2p orbital (valence band) separated from the tungsten 5d orbitals (conduction band) with a bandgap (Fig. 4a and b). The d-orbital of tungsten in WO_3 splits into two sub-orbitals, conventionally represented by “eg” & “t_{2g}”,¹⁸⁴ as given in Fig. 4a and b, respectively. In stoichiometric WO_3 , the “O2p” orbital is filled and the W5d orbital is empty (Fig. 4a), where the bandgap is wide enough, which results in glowing transmission in the thin films. The intercalation of external ions and the charge compensating electrons partially fill the empty d-orbitals, resulting in optical absorption.¹⁸⁵

The removal of the oxygen atom (oxygen vacancy) from the WO_3 structure produces two additional electrons that can be paired in a single spin state (closed-shell) (Fig. 4b) or an unpaired triplet spin state. Furthermore, single/two electrons can be transferred to nearby W atoms, resulting in W^{5+} and $\text{W}^{5+}-\text{W}^{5+}$ centers, respectively. The formation of an oxygen vacancy changes the W–W bond distance across the O vacancy sites. Thus, the W–O bond length and bond angles in the crystal change as the bonded orbitals split, leading to the hybridization of the W5d and O2p orbitals. The inserted electrons and ions diffuse and localize on the tungsten atoms, reducing some of the W^{6+} to W^{5+} states. Because of the absorption of photons, the inserted electrons gain sufficient energy, which can be transferred to the adjacent sites¹⁸⁴ or oxygen vacancies, indicating that oxygen vacancies have a significant impact on both the photochromic and EC properties.^{130,131} The W^{4+} state is considered energetically unfavorable, although XPS results prove the existence of W^{4+} ions.^{136,186} However, XPS is surface sensitive and is not consistent about the bulk state of different tungsten oxide films.¹⁸⁶ The most favorable condition is the creation of a double charge vacancy and two W^{5+} ions. The transfer of charges between two sites is represented by “i” & “j”, as given in eqn (6).



It is believed that the intercalated electrons diffuse in the localized state near the conduction band and the surrounding atoms move, and thus a potential well is generated. The obvious interaction of the electrons and phonons results in the formation of polarons. An electron interacting with a cloud of phonons is called polarons.¹⁸⁷ The polaron-induced optical absorption in $\text{W}_{18}\text{O}_{49}$ produces a broadband absorption spectrum, making the film dark blue.¹⁸⁴ Similarly, in the polaron-induced coloration,¹⁸⁸ the electrons produced due to the oxygen vacancies are trapped in the 5d orbitals of $\text{W}_{18}\text{O}_{49}$ (or defect sites) and then polarize the surrounding lattice to produce small polarons. The resulting polarons then simultaneously yield polaron-to-polaron transitions between two adjacent non-equivalent tungsten sites. In contrast, localized

surface plasmons are the mutual oscillations of free electrons/conductive electrons in materials with large carriers.¹³⁵ Therefore, NIR absorption is widely interpreted as the result of the small polarons hopping and/or local surface plasmon resonance (LSPR).⁸⁸

4. Synthesis and film fabrication strategies of $\text{WO}_{3-x}/\text{WO}_3$

Substantial efforts have been dedicated to synthesizing WO_3 nanostructures with different morphologies, sizes, and dimensions (0D, 1D, 2D, and 3D), as displayed in Fig. 5. To date, tungsten oxide particles have been synthesized by varying the synthetic conditions and technologies, which greatly affect their size, morphology, and functional applications. According to the relevant reaction media, the techniques can be grouped into two main categories, *i.e.*, gas-phase technologies and wet-chemical synthesis methods. Gas-phase technology is classified as plasma sputtering,¹⁹⁰ chemical vapor deposition,¹⁹¹ and physical vapor deposition. Wet-chemical synthesis includes Langmuir–Blodgett deposition,¹⁹² electrophoretic deposition,¹⁹³ electrospinning,¹⁹⁴ and hydrothermal¹⁹⁵ and solvothermal synthesis. Similarly, spray-coating, inkjet printing, spin-coating, roll coating, screen printing, and layer-by-layer self-assembly have also been employed.¹⁹⁶ However, the material synthesis and film fabrication processes are still challenging to adopt on a large scale. These technologies will be reviewed in this section.

4.1. Gas-phase deposition methods

4.1.1. Chemical vapor deposition (CVD). CVD is a common industrial technology used for depositing extremely high-quality thin films on a variety of substrates. The commonly used CVD methods include atmospheric pressure chemical vapor deposition (APCVD) and aerosol-assisted chemical vapor deposition (AACVD). APCVD has certain merits, such as large area scalability, uniformity in thickness, and possibly low cost. For example, the APCVD process is used to manufacture large-area energy efficient FTO/ITO coatings for windows over three meters.¹⁹⁹ The working principle of both the CVD and PVD methods can be found in the review published by Jiang *et al.*²³ Blackman and Parkin²⁰⁰ used the APCVD method to deposit WCl_6 dissolved in different solvents (ethanoic anhydride,

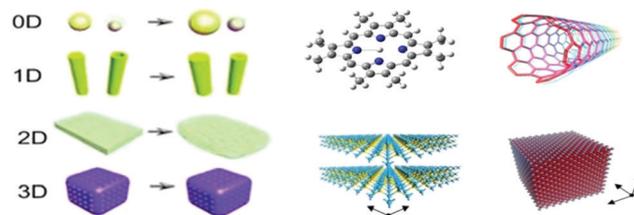


Fig. 5 Schematic illustration of the design of $\text{WO}_{3-x}/\text{WO}_3$ nanostructures with different shapes/dimensions.¹⁹⁷ 0D nanoparticles, 1D nanowires and nanotubes, 2D thin layers, and 3D porous nanostructure. Reproduced with permission.¹⁹⁸ Copyright ©2019, ACS.



ethanoic acid, ethyl ethanoate, methanol, ethanol, 2-propanol, 2-methyl-2-propanol, and water) on a glass substrate. The thin films crystallized with ethanoic anhydride, water, and ethanoic acid were fully oxidized to a monoclinic phase of WO_3 at 625 °C. The deposited films revealed different surface morphologies, *i.e.*, cubic to platelets, under similar conditions, which displayed significantly different performances in different applications, such as photocatalysis, gas sensors, and photochromism.

Brescacin *et al.*²⁰¹ developed phosphorus (P)-doped amorphous WO_3 films *via* CVD processes using volatile, low-melting P-doped tungsten carbonyls. The P-doping preserved the amorphous nature of the material and deposited film, even upon calcination at high temperatures. The tiny amount of dopant released by the precursor during its decomposition was sufficient to inhibit the crystallization of tungsten oxide on the matrix. The films deposited on common glass displayed remarkable EC performances with the maximum coloration efficiency of $66 \text{ cm}^2 \text{ C}^{-1}$. Ou *et al.*²⁰² synthesized tungsten(vi) oxo-complexes of the type $\text{WO}(\text{OR})_3\text{L}$ having partially fluorinated alkoxide and/or chelating ligands (Fig. 6). The thermal decomposition behavior and mass spectrometry (MS) of selected examples were studied. With $\text{WO}(\text{OC}(\text{CF}_3)_2\text{CH}_3)_3$ as a single-source precursor, tungsten oxide (WO_x) was deposited *via* CVD and AACVD and compared. The obtained deposited coatings exhibited different crystallinity, structures, and

morphology (nanowires, nanoplates, and nanocubes). These variations in morphology demonstrated that the solvent used during AACVD controls the growth of the film. The obtained results suggested that the materials deposited by both deposition methods allow not only the procedure to be scaled up, but also have the potential to generate the formation of a film on a wide area with EC properties.

Studies emphasized that MoO_3 and WO_3 have similar EC properties, and thus the APCVD method can be used solely to obtain a film composed of either Mo-doped WO_3 or mixed Mo–W oxides *via* the pyrolytic decomposition of metal carbonyls.²⁰⁵ The visible light transmission of the film deposited on ordinary glass was about 80%, while that on conductive glass (*e.g.*, $\text{SnO}_2:\text{Sb}/\text{glass}$) was about 60%. It is also found that after annealing, the optical transmission of the single oxide films improved, while that of the mixed oxide films slightly decreased. The mixed oxide showed deep-blue coloring only at the cathode, minimizing the visible-light transmittance. Similarly, excellent reversibility with an optical modulation of about 30% was obtained. Numerous reports based on mixed Mo–W oxides have been published.^{206,207}

4.1.2. Physical vapor deposition (PVD). PVD is another gas-based coating process, which involves metal transfer at the atomic level under vacuum conditions. This method may be related to CVD, except that in PVD, the depositing material is

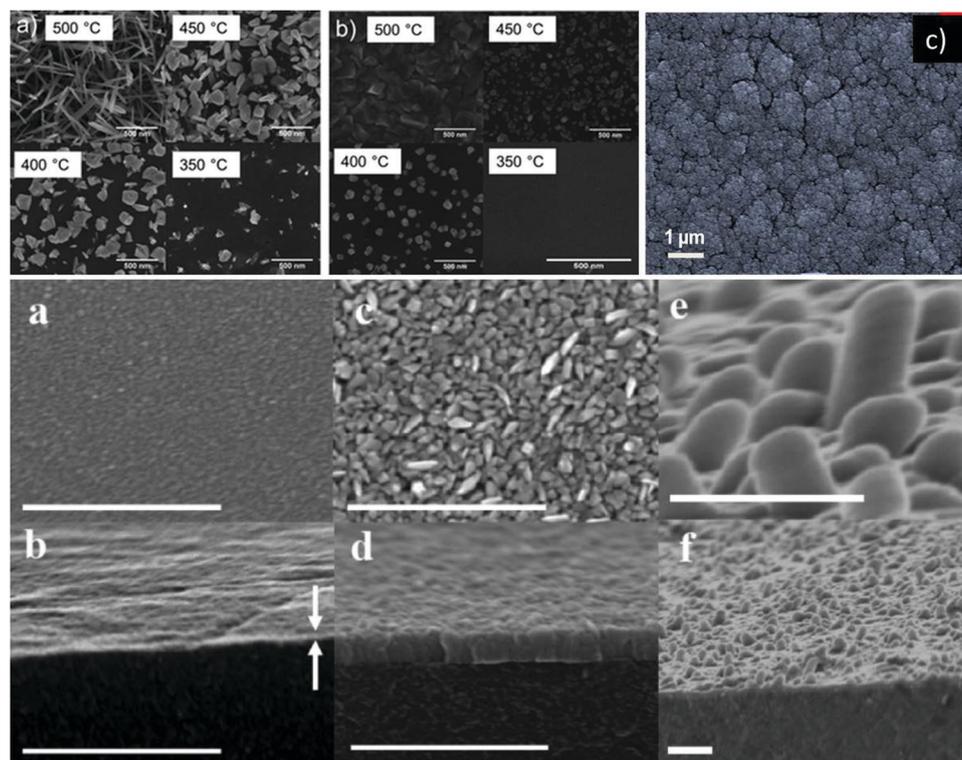


Fig. 6 (top) Plane-view SEM images of deposits grown on Si(100) at 500 °C, 450 °C, 400 °C, and 350 °C from the precursor *via* (a) AACVD and (b) CVD. Reproduced with permission.²⁰² Copyright©2019, ACS. (c) High-magnification SEM images of γ - WO_3 film deposited *via* the HW-CVD (hot wire) method. Reproduced with permission.²⁰³ Copyright©2017, Wiley-VCH. (bottom) Deposition at different times. Top view (a) over 5 min, (c) over 60 min, and (e) over 60 min with 48 h of additional substrate heating during and after deposition. (b, d and f) View of fractured cross section of the corresponding films. All scale bars represent 500 nm. Reproduced with permission.²⁰⁴ Copyright©2010, Wiley-VCH.



initially in the solid state, whereas in CVD, the gaseous form of the precursor material is introduced into the reaction chamber.

PVD is comprised of four steps as follows:

- (i) evaporation of materials deposited by high energy sources (*e.g.*, electron beam/ions that evaporate atoms from the surface);
- (ii) transport of the vapor to the substrate to be coated;
- (iii) the reaction between the metal atoms and the appropriate reactive gas (such as oxygen, nitrogen, and methane) during the transport stage; and
- (iv) deposition of the coatings on the surface of the substrate.

The PVD method has several advantages including (i) PVD thin coatings demonstrate better properties compared to the substrate materials; (ii) all types of inorganic materials and some types of organic materials can be used; and (iii) the procedure is more environmentally benign than several other techniques, *e.g.*, simple electroplating. Meanwhile, the PVD method also has certain disadvantages, including (i) problems with coating complex shapes; (ii) high cost and low output; and (iii) complexity of the process.

Zhang *et al.*²⁰⁸ demonstrated the growth of a WO₃-WS₂ bilayer hybrid with WO₃ monolayer species attached on the surface of large-size WS₂ monolayers, as shown in Fig. 7a-d. The optical analysis showed that the photoluminescence quantum yield (PLQY) of the as-prepared WO₃-WS₂ hybrid was excellent and could reach 11.6% higher PL than that of the bare WS₂ monolayers, which is about 13-times greater than that *via* the mechanical exfoliation of WS₂ (Fig. 7e-g). The improved PL enhancement mechanism can be investigated by carrying out time-dependent optical measurements. The generation of WO₃-WS₂ hybrid structures with ultrahigh PLQY can provide an efficient method for developing highly effective 2D integrated photonic applications. Hadifakoor *et al.*²⁰⁹ fabricated a

DMD skeleton of WO₃/Ag/WO₃ thin films with various thicknesses through the PVD approach *via* electron-beam evaporation under the vacuum at pressure of 10⁻⁵ torr and calcination temperature of 100–400 °C. The structural and optical properties of the nano-multilayer composites structures were studied in heat mirror applications. Nanosized Ag layers with thicknesses of 10 nm, 12 nm, and 14 nm were deposited on monolayer WO₃ films. Atomic force microscopy (AFM) showed the morphology of the individual layers, indicating that a smoother layer can be achieved after annealing at 300 °C. Ellipsometry analysis was performed to examine the bilayers, thickness of the Ag film layer, and inter-connectivity between the WO₃-Ag-WO₃ layers. There was almost no interference among the WO₃-WO₃ layers in the samples with Ag thicknesses of 12 nm and 14 nm. The efficiency of the heat mirrors was evaluated according to their optical characteristics and best performance. Recently, Hoseinzadeh *et al.*²⁰⁹ studied a WO₃ and Ag composite deposited *via* the layer-by-layer (LbL) PVD method as an EC electrode on an FTO glass substrate. Nano-sized silver noble metal can behave as an electron-trap center, which enables the departure of charges. The electrochemical and optical properties were examined through CV and UV-visible absorption experiments, which showed that heating the deposited film at 200 °C can provide much better conductivity (90 mA) and transmittance change ($\Delta T = 90\%$ at a continuous switching step), endowing WO₃-Ag-WO₃-Ag with excellent EC properties. Esmail *et al.*²¹⁰ demonstrated amorphous WO₃ smooth films thinner than 200 nm, which exhibited an excellent coloration efficiency with a switching speed in the order of seconds.

4.2. Wet-chemical synthesis methods

The gas-phase deposition of coatings is a highly pure and efficient route for the fabrication of WO₃ films. However, the required instrumentation is very complex and expensive. Conversely, solution-based particle synthesis and film formation are cost effective, controllable, and high yielding, which provide an alternative reliable route for the production of smart coatings, reflective displays, and sensors.²¹¹

4.2.1. Sol-gel method. The “sol-gel” terminology describes a wide-range process, which is the production of a solid phase through the colloidal dispersion/gelation of a colloidal suspension (the sol). Further reactions lead to bonds between the sol particles, creating an infinite network of particles (gels). The “dry gel” (xerogel) state results after drying the gel, and subsequent heating can remove the unreacted organic impurities, stabilize the gel, densify, and crystallize the material.^{80,212} Thus, a metal solution can be converted to an oxide chain-like complex compound such as a metal alkoxide *via* a simple polymerization reaction and/or using a template(s). The compounds used for the oxide polymerization include common organic solvents, organic acids, and templates, and the polymerization occurs through hydrolysis/condensation. The sol-gel method was used only to prepare metals and semi-metal oxides, and then extended to organic-inorganic hybrids and occasionally for the production of pure organic

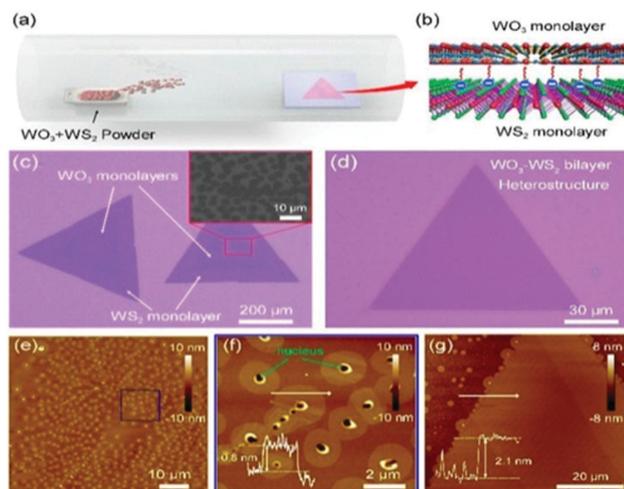


Fig. 7 (a and b) Schematic of the synthetic process and 3D representation of the WO₃-WS₂ heterostructure. (c and d) Typical optical image of (c) partially covered and (d) completely covered WO₃-WS₂ heterostructures. The inset in (c) shows the SEM image of the area within the red box. (e-g) AFM images of (e and f) partially covered and (g) completely covered heterostructures. The insets show the corresponding height profile. Reproduced with permission.²⁰⁸ Copyright©2019, ACS.



networks.⁸⁰ WO₃, V₂O₅, MnO₂, and many other transition-metal oxides display numerous oxidation states. The composition and structure of their precursors are susceptible to the pH and redox states, and hence they can control the sol and gelation.⁸⁰ The sol-gel deposition techniques include dip-coating,^{213–216} spin-coating,²¹⁷ inkjet-printing,^{218,219} Langmuir-Blodgett deposition,^{220–222} and spray-pyrolysis.^{76,223} This technology requires cheaper equipment for large-area surface coatings compared to gas-phase technologies.

The sol-gel method is attractive due to the following reasons:

(i) it provides an easy route for the preparation of metal oxide microstructures and mixed oxides with a controlled composition and homogeneity,

(ii) processing produces an amorphous hydrogel, which is better than crystalline films due to its excellent elasticity,

(iii) can reform volume changes caused by redox reactions, and improve conductivity,

(iv) films can well adhere to ITO/FTO films and be processed at ambient pressure, and

(v) deals with a new deposition method of WO₃-based coating on a large scale.

In 1984, Chemseddine *et al.*²²⁴ first reported the results of a sol-gel EC WO₃ film, which was then subjected to different sol-gel processes for the production of pristine WO₃ and mixed oxides. The coloration efficiency values compared to the reported sol-gel films were 70 cm² C⁻¹ (at 685 nm) and 223–167 cm² C⁻¹ (at 800 nm)²²⁵ compared to 115 cm² C⁻¹ (at 633 nm) for the amorphous films²²⁶ produced *via* thermal evaporation and 42 cm² C⁻¹ (at 650 nm) reported for sputtered polycrystalline films.^{227–229} The advantage of the sol-gel technique is its capability to generate various mixed oxides from a variety of hybrid mixtures with controlled compositions.^{230–240} Nguyen *et al.*²⁴¹ used the sol-gel spin coating process and further annealing to form amorphous EC WO_x thin films. The precursor solutions were prepared from ammonium metatungstate hydrate ((NH₄)₆[H₂W₁₂O₄₀]·4H₂O) and dimethylformamide (DMF) to produce a smooth and transparent amorphous WO_x film *via* spin-coating followed by calcination in air at 250 °C for 2 h. The obtained EC film exhibited excellent

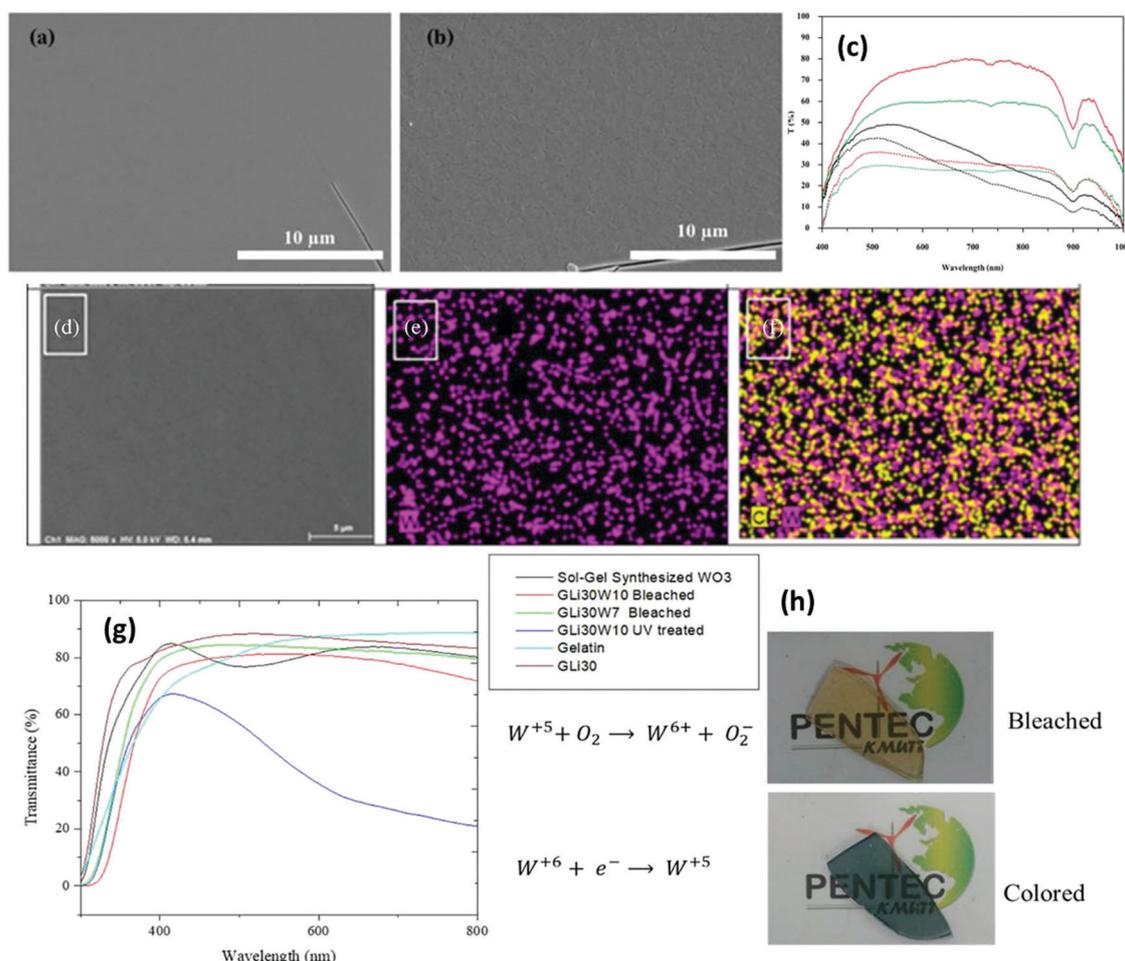


Fig. 8 Surface morphology of the 5% MoO₃–95% WO₃ film annealed at (a) 300 °C and (b) 400 °C. (c) Transmittance spectra. Reproduced with permission.²⁴³ Copyright@2017. Elsevier. (d) Film containing 30 wt% LiClO₄ and 10 wt% WO₃ (sample denoted as GLi₃₀W₁₀), (e) tungsten map and (f) chlorine and tungsten distribution mapping. (g) Transmittance spectra of free-standing films and (g) optical images of colored and bleached states. Reproduced with permission.²⁴² Copyright@2020, Wiley-VCH.



stability over 4000 switching cycles with a coloration efficiency of $125 \text{ cm}^2 \text{ C}^{-1}$ and color/bleach times of 5/2.5 s. Amorphous WO_x film-based devices are impeccable for next-generation smart windows and other electronic display devices. Recently, Azarian *et al.*²⁴² explored amorphous electro-photochromic films formed *via* the *in situ* sol-gel synthesis of 10 wt% WO_3 and 30 wt% LiClO_4 (denoted as $\text{GLi}_{30}\text{W}_{10}$) in a gelatin/lithium cosolvent mixture (Fig. 8d-f). The transparent single-layer film was flexible and adhesive, active under UV radiation with a photo-response time of 30 s, and gradually reversed once the UV source was blocked. Besides, in the presence of a conductive ion storage anode, the film on ITO glass exhibited an excellent EC response. The EC device configured with ITO/ $\text{GLi}_{30}\text{W}_{10}$ /NiO/ITO achieved a coloration efficiency of $51.54 \text{ cm}^2 \text{ C}^{-1}$ and fast color-bleach response times (2 s/1 s). Jittiarporn *et al.*²⁴³ prepared hybrid MoO_3 - WO_3 thin films on ITO glass *via* a sol-gel dip-coating process (Fig. 8). The formation of the film could be influenced by the calcination temperature, the concentration of MoO_3 (0–10%), and the copolymer template of Pluronic P-123 triblock (0–20% w/v). The lowest amount of 5% MoO_3 -95% WO_3 films without the use of the template and 200–300 °C annealing resulted in a high CE, excellent cycling stability, and low response time. The surface roughness and low crystallinity of the films were thought to be attributed to their enhanced EC features. Amorphous mesoporous WO_3 with a pore size in the range of 20–30 nm was fabricated *via* evaporation induced self-assembly and used as an electrochromic supercapacitor (ECS).¹²⁰ The resultant device based on ECS exhibited excellent optical modulation (76% at 700 nm), ultrafast switching speeds (0.8 s for coloration and 0.4 s for bleaching), and a high areal capacitance (2.57 mF cm^{-2}), even at a high current density (1.0 mA cm^{-2}).

Sun *et al.*²⁴⁴ prepared mixed molybdenum-tungsten oxides with different stoichiometries ($\text{Mo}_x\text{W}_{1-x}\text{O}_3$, $0 < x < 1$) *via* sol-gel deposition from peroxo-polymolybdotungstate solutions onto FTO glass substrates. The $\text{Mo}_x\text{W}_{1-x}\text{O}_3$ films were annealed at 450 °C. The transmittance modulation of the mixed oxide films was improved compared to the pristine WO_3 film and no broadening of spectral modulation was observed. Kharade *et al.*²³¹ also employed a similar method to prepare MoO_3 mixed WO_3 thin films *via* hybrid physicochemical synthesis involving the microwave-assisted sol-gel preparation of WO_3 and vacuum evaporation of MoO_3 on an FTO glass substrate. Similarly, a mesoporous WO_3 - TiO_2 composite film was fabricated *via* a sol-gel-based two-stage dip-coating technique on ITO glass, and subsequently annealed at 450 °C, 500 °C, and 600 °C.²⁴⁵ The composite film calcined at 600 °C contained orthorhombic WO_3 and anatase TiO_2 . Mesoporosity was achieved in the film by employing silicates as templates. Mixed hexagonal and monoclinic phases were obtained upon calcination at 450 °C and 500 °C. The results show that nanocrystal anatase TiO_2 can stabilize the orthorhombic WO_3 phase by calcination at 600 °C. The mesoporous dye-sensitized WO_3 - TiO_2 composite film calcined at 600 °C showed an optical modulation of 51% in the NIR region. In contrast, the devices examined for the composite layers annealed at 450 °C and

500 °C displayed a moderate optical modulation of 24.9% and 38%, respectively.

4.2.2. Hydrothermal and solvothermal methods. The solvothermal and hydrothermal methods are commonly used for the synthesis of nanostructured materials. These methods allow the generation of various nanoscale morphologies and the possibility of structural control, which may not be afforded by high-temperature wet-chemical reactions. Hydrothermal synthesis involves the use of water-soluble precursor compounds at room temperature, which is cost-effective, and also depends on the relevant reaction variables, *e.g.*, concentration, pH, temperature, time, and reducing agents, that control the structural evolution, morphology, and size in the required dimensions (Table 2).

Due to its distinctive features, the hydrothermal method is considered the most suitable route for the synthesis of tungsten-based nanostructures. The solvothermal route involves nonaqueous media, which provide unique physicochemical characteristics to the material design for various high-tech devices. Because of the wide applications of WO_3 and WO_{3-x} nanostructures in energy-saving high-tech devices, many research groups have focused on these materials in the last few years. For example, Guo *et al.*²⁷⁷ prepared $\text{W}_{18}\text{O}_{49}$ nanorods by pyrolyzing the $(\text{NH}_4)_x\text{WO}_{3+x/2}$ precursor, which was synthesized in the presence of an inorganic surfactant ($(\text{NH}_4)_2\text{SO}_4$) in a reducing environment. Pan *et al.*²⁶⁰ used glycerol and ammonium sulfate ($(\text{NH}_4)_2\text{SO}_4$) as a surfactant and capping agent, respectively, for the synthesis of coral-like nanostructures *via* the hydrothermal method. The obtained product showed excellent EC properties, including a fast-switching time (6/5 s for bleaching/coloration), a good CE ($109.6 \text{ cm}^2 \text{ C}^{-1}$), and a large optical modulation ($\sim 78.1\%$) at 630 nm by applying a potential of $\pm 1 \text{ V}$. Similarly, Kadam *et al.*²⁷⁸ used propylene glycol (PG) as a surfactant to increase the size of the nanocrystals on the surface of ITO glass, which was then used for EC application. Chu *et al.*²⁷⁹ used a seed layer and a surfactant-free hydrothermal method to synthesize nanoplates on FTO glass. The obtained highly oriented 2D nanostructure exhibited potential for practical applications in smart windows due to its fast-switching speeds of 12 s and 3 s for coloration and bleaching, respectively. Mathuri *et al.*²⁸⁰ also used an FTO glass substrate and synthesized WO_3 nanoplates; however, they used different hydrothermal reaction times (8–16 h), and temperatures (120–180 °C), which resulted in hexagonal and monoclinic WO_3 nanoplates. The WO_3 -deposited films at 120 °C and 180 °C for different reaction times of 8 h, 12 h, and 16 h possessed different surface morphologies, such as buds-like, bricks-like, sheets, and platelets-like structures. The optical bandgap of the deposited WO_3 films varied in range of 2.5–2.9 eV and 2.6–2.75 eV, respectively.

Liang *et al.*²⁸¹ produced a WO_3 film using FTO-coated glass *via* the hydrothermal method. Their results showed that different WO_3 nanoparticles such as nanosheets, nanoflakes, nanocuboids, and 3D nanowire flowers could be synthesized by adjusting the precursor solution composition. The solvent plays



Table 2 List of the preparation methods and EC properties of 0D, 1D, 2D, and 3D WO_{2.72}/WO₃-based nanoparticles

Technology	Methods	Sub-method	Materials	Morphologies	Coloration efficiency, color/bleach response and cyclic stability (cycles)	Ref.	
Mechanochemical process (Ball milling)			Na _x WO ₃ (bronze)	Nanoparticles		246	
Gas-phase technologies	CVD/PVD	Magnetron sputtering	WO _x		46.9 cm ² C ⁻¹ , 7.8/117.8 s	247	
		R.F. sputtering	WO ₃		46.45 cm ² C ⁻¹ , 8.33/4.16 s	248	
		R.F. sputtering	WO _{3+x}	Granular	~213 cm ² C ⁻¹ , 5000	249	
		Thermal vapor deposition	WO _{3-x}	Nanorods		250	
		HIPIMS	WO ₃ /Ag/WO ₃		30.5 cm ² C ⁻¹ , 15/22.4 s, 2500	125	
Wet-chemical technologies	Hydrothermal/solvothermal	APCVD	WO _{3-x}	Nanowires, nanoplates		200	
			WO _{2.72}	Sub-micro fibers		251	
			WO _{2.72}	Nanodots	154 cm ² C ⁻¹ , 1/1 s	252–255	
			WO _{2.72}	Nanocrystals/particles	124.5 cm ² C ⁻¹ , 1.9/2.8 s, 1000	139 and 256	
			WO _{2.72}	Nanowires	(—)	135	
			WO _{2.72}	Nanowires assembled	35.7 cm ² C ⁻¹ , 2/4 s, 1000	119 and 257	
			WO ₃	Nanorods/Nanowires arrays	102.8 cm ² C ⁻¹ , 25.2/18 s, 7.6/4.2 s, 3000	229 and 258	
			WO ₃	Nanoparticles	151.9 cm ² C ⁻¹ , 1.3/1.7 s	259	
			WO ₃	Nanofibers	109.6 cm ² C ⁻¹ , 2.7/7.9 s, 4000	260	
			WO ₃	Particles	178.7 cm ² C ⁻¹ , 1.9/2.5 s, 5000	261	
			WO ₃	Nanoflakes/sheets	55.6 cm ² C ⁻¹ , 5.2/2.2, 3000	262	
			WO _{2.72}	Nanowires/Nanotrees	75.35 cm ² C ⁻¹ , 2.64/7.28 s	263	
			WO _{2.72}	Nanoneedles	150 cm ² C ⁻¹ , 500	264	
			WO _{2.72}	Nanobelts	(—)	265	
			WO ₃	Nanobricks	39.24 cm ² C ⁻¹ , 9.7/6.9 s	266	
			WO ₃ /NiO	Nanoporous structures	85.9 cm ² C ⁻¹ , 21.76/4.54 s	267	
			WO ₃ ·H ₂ O, WO ₃ ·2H ₂ O	Nanoplates	384.6 cm ² C ⁻¹ , 322.6 cm ² C ⁻¹ , 500	163	
			WO _{2.72} /POMs	Nanowires	42.31 cm ² C ⁻¹ , 52/26 s, 500	122	
			WO ₃ -TiO ₂	Hierarchical nanostructure	128.3 cm ² C ⁻¹ , 6 s, 1000	268	
		Sol-gel			WO _x -NbO _x	Nanowires	
	WO _{2.72}			Mesoporous	80.5 cm ² C ⁻¹ , 5.3/3 s	247	
	WO _{2.72}			Mesoporous	46.9 cm ² C ⁻¹ , 7.8/117 s, 8	247	
	WO _x				125 cm ² C ⁻¹ , 5/2.5 s, 4000	241	
	WO ₃			Mesoporous	35.8 cm ² C ⁻¹ , 16/30 s	269	
	Ag-WO ₃			Nanowires composite	45.3 cm ² C ⁻¹ , 12/2 s	270	
	WO ₃ /TiO ₂ /WO _x				480 cm ² C ⁻¹ , 3.3/2.8 s	271	
	WO ₃ -PEDOT:PSS			composite	68.8 cm ² C ⁻¹ , 2.4/0.8 s, 10 000	272	
	Langmuir-Blodgett			WO ₃	Nanobundles	41.8 cm ² C ⁻¹	273
	Electrostatic force-assisted printing			WO ₃ ink	WO ₃ powder	51.5 cm ² C ⁻¹ , 65.5% at 700 nm, 25 s/24 s	274
	Spray pyrolysis						
	Pulsed electrochemical deposition			WO ₃	Poros films	118.3 cm ² C ⁻¹ , 6/2.7 s	275
	Electrochemical deposition			WO ₃	Nanorods/nanoparticles	~90 cm ² C ⁻¹ , 3/2 s	185
	Electrochemical deposition			WO ₃	Nanogranular	41.16 cm ² C ⁻¹ , 7.5/3.9 s	276

a significant role in controlling the shape and size of nano-WO₃. The transmission spectra indicated that the WO₃ film showed 50% and 70% transmittance in the visible spectrum both in the colored and bleached state, respectively, which with

the applied potential of ±1.0 V, was maintained for 1000 continuous coloration/bleaching cycles. Bhosale *et al.*²⁶¹ used a wet-etching process and fabricated a WO₃ EC film on ITO glass *via* the hydrothermal method. The etched WO₃ exhibited



roughness, porosity, and an open-tunnel structure. The resultant films showed excellent coloration efficiency ($\sim 178.7 \text{ cm}^2 \text{ C}^{-1}$), a fast response time (1.9/2.5 s), transmittance modulation ($\sim 49\%$ at 630 nm), and stability (5000 cycles).

Lu *et al.*²⁸² prepared uniform WO_3 nanorods using NaCl as a capping agent. The solution pH and amount of NaCl played key roles in the morphology and microstructure evolution. The prepared nanorods were aligned on ITO glass *via* the drop-casting method. Sun *et al.*²⁸³ synthesized $\text{W}_{18}\text{O}_{49}$ nanowires *via* the solvothermal method using WCl_6 and cyclohexanol precursors. The sample calcined up to 400°C was stable and no phase change was observed; however, upon increasing the calcination temperature to $>400^\circ\text{C}$, a phase conversion to monoclinic WO_3 occurred. Lu *et al.*²⁸⁴ grew a vertical nanowire array on FTO glass for the fabrication of an EC working electrode. They used WCl_6 , CH_3OH , and polyethylene glycol (PEG) as a capping agent. The WO_3 nanowires coated FTO glass showed maximum coloration/bleaching of 41.63% at 632.8 nm and the switching times of 9.8 s and 2 s, respectively. Li and co-workers²⁶³ synthesized hierarchical nanotree-like WO_3 architectures directly *via* seed-mediated growth on FTO, as shown in Fig. 9a–f. They used H_2WO_4 , $\text{SC}(\text{NH}_2)_2$, $\text{C}_4\text{H}_4\text{O}_4$, aqueous HCl, and FTO glass and hydrothermal treatment to form a WO_3 seed layer. The same precursors were used and the seed layer on FTO was put in an autoclave to grow the WO_3 nanotree architecture. They reported that the nanotree trunks showed

(002) orientation growth, whereas the branches exhibited (100) and (200) oriented growth directions. The optical modulation was 74.7% at 630 nm at a potential 0.2 V (Fig. 9g). The coloration efficiency (Fig. 9h) was $75.35 \text{ cm}^2 \text{ C}^{-1}$ with fast switching responses of 2.64/7.28 s. The electrochemical and EC properties of the WO_3 nanotrees were attributed to their large surface area, showing potential for application in EC smart windows.

Li *et al.*²⁶⁴ synthesized needle-like $\text{W}_{18}\text{O}_{49}$ to form a thin film on an ITO substrate. The needle-like $\text{W}_{18}\text{O}_{49}$ nanocrystals exhibited a good performance and control over NIR light. Their powder-based optical properties also showed that the needle-like $\text{W}_{18}\text{O}_{49}$ nanocrystals have excellent photoabsorption in the NIR region, corresponding to 780–2500 nm. The LbL-coated film exhibited high color contrast, quick switching kinetics, improved coloration efficiencies ($150 \text{ cm}^2 \text{ C}^{-1}$ at 650 nm and $255 \text{ cm}^2 \text{ C}^{-1}$ at 1300 nm), good long-time stability (reversibility of 98% after 500 cycles), and wide EC spectral coverage for both the Vis and IR regions. Huang *et al.*²⁸⁵ also obtained WO_3 hexagonal nanorods, although they used tungstate powder, commercial tungsten oxide powder, H_2O_2 , oxalic acid, Na_2SO_4 for the synthesis of nanorods and nanofibers *via* the hydrothermal method. Guo *et al.*²⁸⁶ obtained nanowires; however, they also used graphene sheets and ethanol as the solvent. Park *et al.*²⁸⁷ synthesized urchin-like $\text{W}_{18}\text{O}_{49}$ particles *via* the simple sonication of a WCl_6 and ethanol mixture. Gu *et al.*²⁸⁸

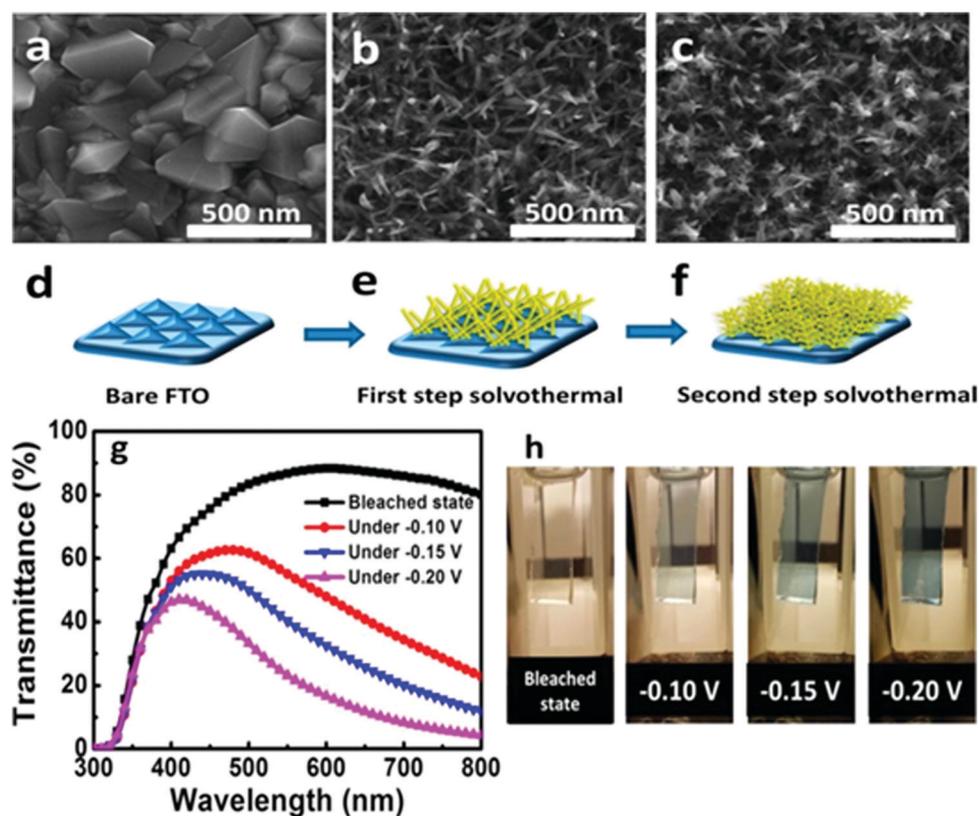


Fig. 9 (a–c) SEM images of WO_3 nanowires and nanotree grown on FTO and (d–f) corresponding schematic illustrations, respectively. (g) UV-vis spectra of WO_3 nanotree at various potential and photographs of its color change. Reproduced with permission.²⁶³ Copyright©2018, ACS.



synthesized hierarchical structures composed of 1D nanowires/nanorods *via* a simple hydrothermal method with various surfactants, *e.g.*, K_2SO_4 , Rb_2SO_4 , Na_2SO_4 and $(NH_4)_2SO_4$. The urchin-like and ribbon-like structures were produced with K_2SO_4 and Rb_2SO_4 based on a sulfate-induced oriented growth mechanism. Guo *et al.*²⁸⁹ synthesized $W_{18}O_{49}$ nanostructures with different morphologies, *i.e.*, nanowires, nanotubes, and nanoparticles, *via* the solvothermal method (Fig. 10a–i) by changing the alcohol and tungsten source (WCl_6 and $W(C_2OH)_6$). They reported that using primary alcohol resulted in $W_{18}O_{49}$ nanostructures, whereas pentanol and secondary alcohols, *e.g.*, 2-propanol and 2-butanol, gave a mixture of WO_3 – $W_{18}O_{49}$. The dispersed $W_{18}O_{49}$ particle suspension was paint coated on quartz glass using an applicator (Fig. 10j).

We also attempted to use similar routes to synthesize 1D stoichiometric and nonstoichiometric tungsten oxide with different morphologies (nanotubes (NT), nanowires (NW), nanowire bundles (NWB), nanorods (NR), nanoplates (NP), and urchin-like nanospheres (Fig. 11a–i)). The different morphologies were tuned by changing a variety of pertinent variables such as organic/inorganic surfactants (K_2SO_4 , Na_2SO_4 , Li_2SO_4 , $(NH_4)_2SO_4$, KCl, KNO_3 , citric acid, oxalic acid, PVP, and CTAB), temperature (120–180 °C), reaction time (1–24 h), tungsten salt, ($Na_2WO_4 \cdot 2H_2O$ and WCl_6) and solvent (water, ethanol and ethylene glycol). Controlled experiments resulted in highly crystalline 1D WO_3 nanostructures. The Brunauer–Emmett–Teller (BET) surface area was derived for the h- WO_3 nanotubes ($71.95 \text{ m}^2 \text{ g}^{-1}$), WO_{3-x} nanowires ($51.83 \text{ m}^2 \text{ g}^{-1}$), h- WO_3

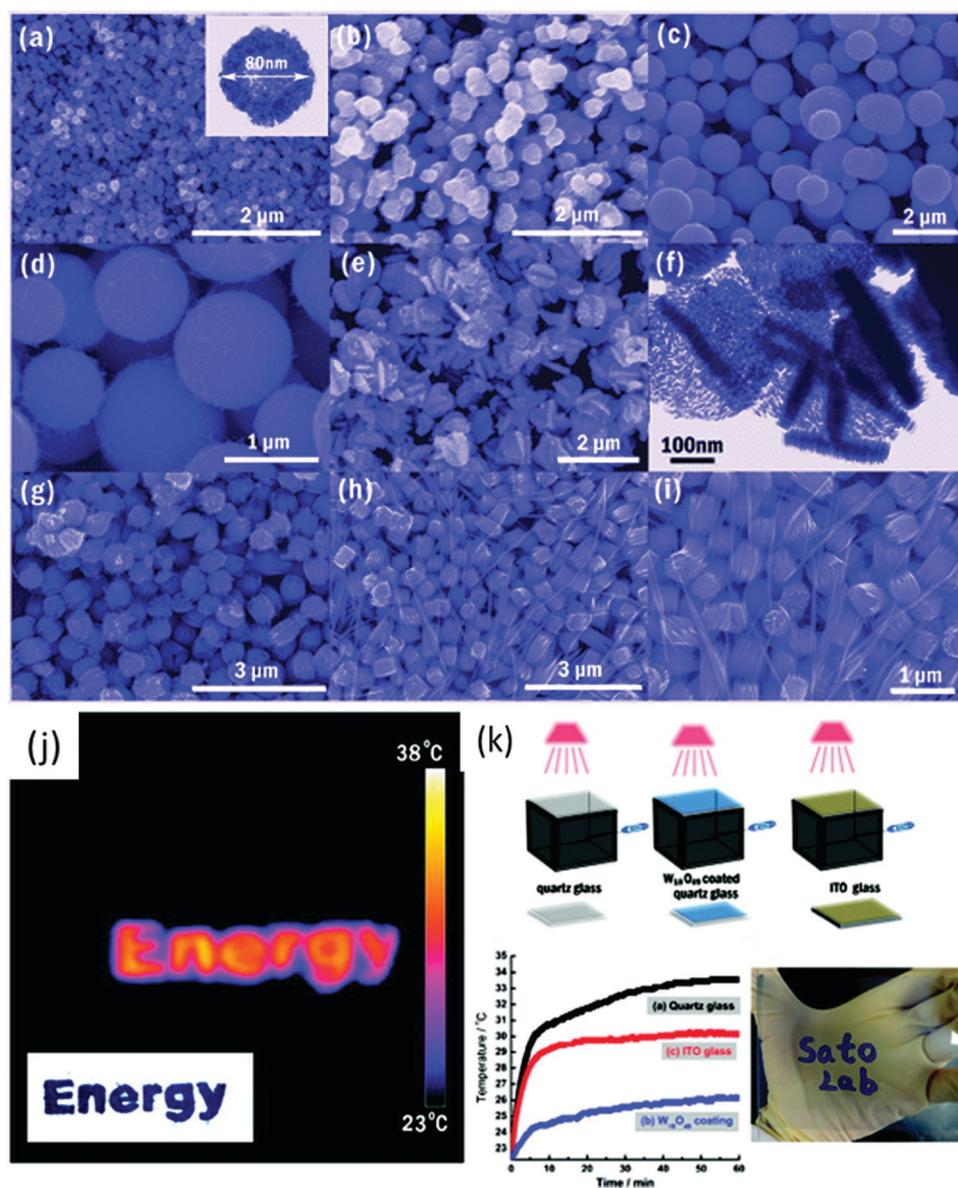


Fig. 10 (a–i) SEM and TEM images of the $W_{18}O_{49}$ nanomaterials. (j) Schematic illustration of the simulated experiment. Sealed boxes with a facet covered by quartz glass, $W_{18}O_{49}$ nanorods coated on quartz glass and ITO glass were irradiated by a 50 W halogen lamp. The temperature changes were determined using an electronic thermometer. (k) Time-dependent temperature in the box. Reproduced with permission.²⁸⁹ Copyright © 2012, ACS.



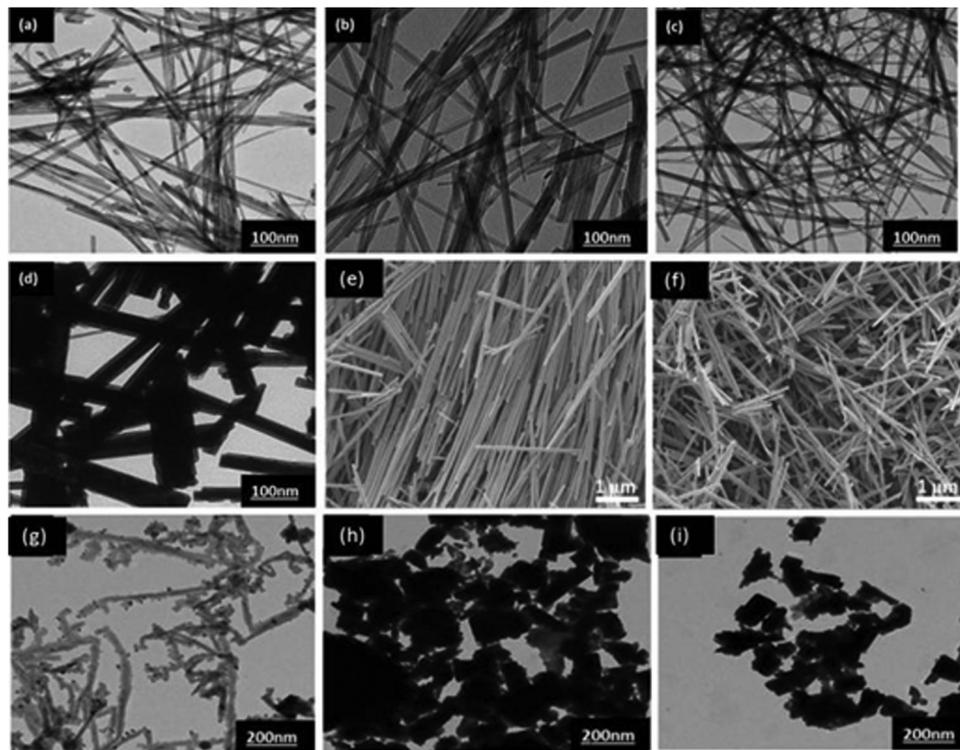


Fig. 11 TEM and SEM images of tungsten oxide nanoparticles: (a) WO_{3-x} -NW, (b) WO_3 -NT, (c) h- WO_3 -NW, (d–f) h- WO_3 -NR, and (g) $(\text{NH}_4)_2\text{SO}_4$, and (h) and (i) CA and OA without K_2SO_4 , respectively. Reproduced with permission.¹⁵² Copyright@2020, Elsevier.

nanowires ($42.70 \text{ m}^2 \text{ g}^{-1}$), nanorods ($24.60 \text{ m}^2 \text{ g}^{-1}$), and nanowire bundle ordered structures ($18.4 \text{ m}^2 \text{ g}^{-1}$). The effects of pertinent variables such as surfactants were explored by carrying out a series of experiments. Under hydrothermal conditions, the ions in the solutions and organic acids (citric acid and oxalic acid) adsorbed on the WO_3 crystal facets and produced highly oriented 1D WO_3 nanostructures. Meanwhile, the K^+ , Na^+ , Li^+ and SO_4^{2-} ions adsorbed on the WO_3 crystal planes of h- WO_3 .¹⁴³ The SO_4^{2-} anions preferentially adsorbed on the surface parallel to the c -axis at the $\langle 001 \rangle$ plane of WO_3 , resulting in the formation of 1D

single-crystal nanostructures.¹⁹⁵ CA and OA allowed the controlled growth and conversion of the crystal phases. It was concluded that an appropriate number of ions and CA and OA played a crucial role in controlling the nanocrystal phases and crystallinity. Moreover, ultrathin aggregated nanotubes and nanowires were produced, which then simultaneously assembled into bundles due to the self-assembly and oriented attachment mechanism.²⁸⁸

Zheng and co-workers²⁹⁰ reported various morphologies of WO_3 , *i.e.*, nanorods, nanofibers, and nanoflakes, by changing the surfactant and reaction conditions, as shown in Fig. 12a–d.

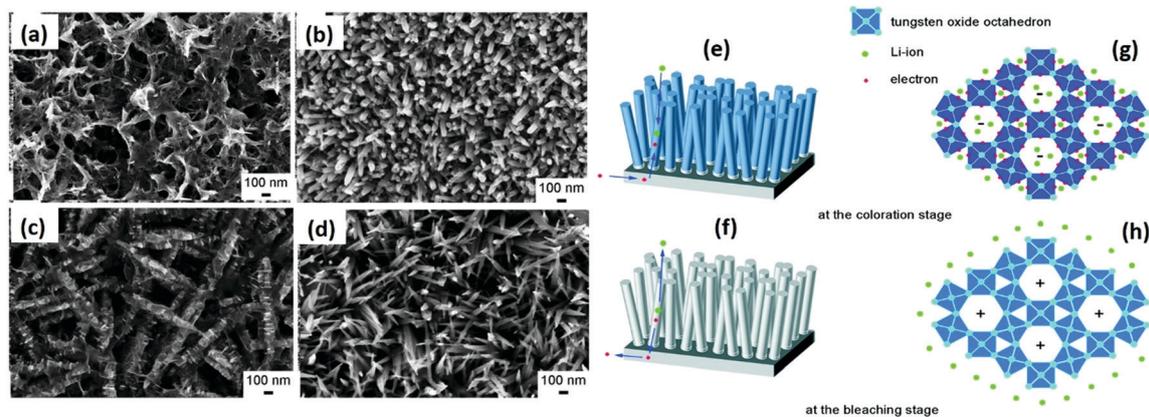


Fig. 12 SEM images of (a) porous WO_3 nanofibers, (b) WNRs-1, (c) WO_3 nanoflake arrays, (d) WNRs-2. (e) EC mechanism of the WNRs the interface reaction process of Li-ions and electrons between WO_3 and the electrolyte and the diffusion process of Li-ions in the WO_3 crystal lattice. Reproduced with permission.²⁹⁰ Copyright@2015, RSC.



The impact of crystallinity and morphologies on the EC properties was also investigated. Accordingly, nanorods were grown *via* seed-mediated growth on an FTO substrate at pH 2 *via* the hydrothermal process without any hard organic templates, followed by annealing (Fig. 12b). The nanofibers grew on FTO at pH 2.4 with the assistance of KCl as a capping agent, keeping the temperature 160 °C for 4 h (Fig. 12c). For the nanoflakes, the pH was 1.5 and NaCl is used as the capping agent at 180 °C for 40 min (Fig. 12d). The WO₃ nanorods grown on FTO showed a high optical modulation of 64%, coloration efficiency of 61 cm² C⁻¹, and switch speeds of 5 s and 6 s as shown in Fig. 12e–h. It was reported that the highly ordered WO₃ nanorods grown on an FTO seed layer showed the best EC performance among the morphologies. High-temperature annealing also affected the EC properties, given that the annealed WO₃ nanorod sample possessed a low optical modulation due to its long hexagonal tunnel, but short response time due to its high crystallinity.

5. Strategies to improve EC properties

5.1. WO_{2.72} hybridized/mixed composites

It is believed that composite materials can aggregate and boost the performance of two/more ingredients. The mixing of two active materials with matching behavior can achieve a dominant interfacial area to a large extent. In nanoscale composites, the structural organization is advantageous because the physicochemical features arise from the confined interfaces. However, in arbitrary mixing, even the active components of material blends may be affected/detrimental due to the fact

that their passageways are tortuous, and there may be some inactive sites. These composites may be more beneficial for the deliberate design of architectures from their mesoscale structure.^{291,292} The metal oxide nanocomposites with template mesoscale architectures have fast switching kinetics and long cycle life. In this case, it has been reported that two degenerate metal oxide nanocrystals may efficiently modulate the NIR spectral transmittance *via* electrochemical charging, and then discharging of free electrons, which are responsible for LSPR trigger absorption.^{40,293,294} Hence, it can be concluded that NIR spectral light is much more intense at shorter wavelengths (700–1300 nm), which can be ideal for composite nanocrystals with LSPR absorption, and thus beneficial for dynamic solar light.

Polyoxometalate nanoclusters (POMs) denote a familiar class of metal oxide cluster compounds with framework topologies, which possess electronic and optical properties.^{295,296} POMs as an electron reservoir undergo reversible multi-electron transfer reactions and significantly impart EC properties. Gu *et al.*¹²² studied the EC performance of crown-type polyoxometalates, K₂₈Li₅H₇P₈W₄₈O₁₈₄·92H₂O (P₈W₄₈), and W₁₈O₄₉ nanowires coated on ITO glass *via* the LbL process, as shown in Fig. 13. The P₈W₄₈ anion is a visible light absorption material owing to its wide range of stability. In the reduced state, it may exhibit good absorptivity of visible light, while the W₁₈O₄₉ NWs are a candidate with excellent absorptivity of NIR light. The effects of the W₁₈O₄₉ NWs and P₈W₄₈ on near-infrared and visible-light selective modulation were studied *via* the self-assembly technique. Meanwhile, the W₁₈O₄₉ NWs or P₈W₄₈ anion are negatively charged species, which hybridized with the positively charged poly(ethylenimine) (PEI)

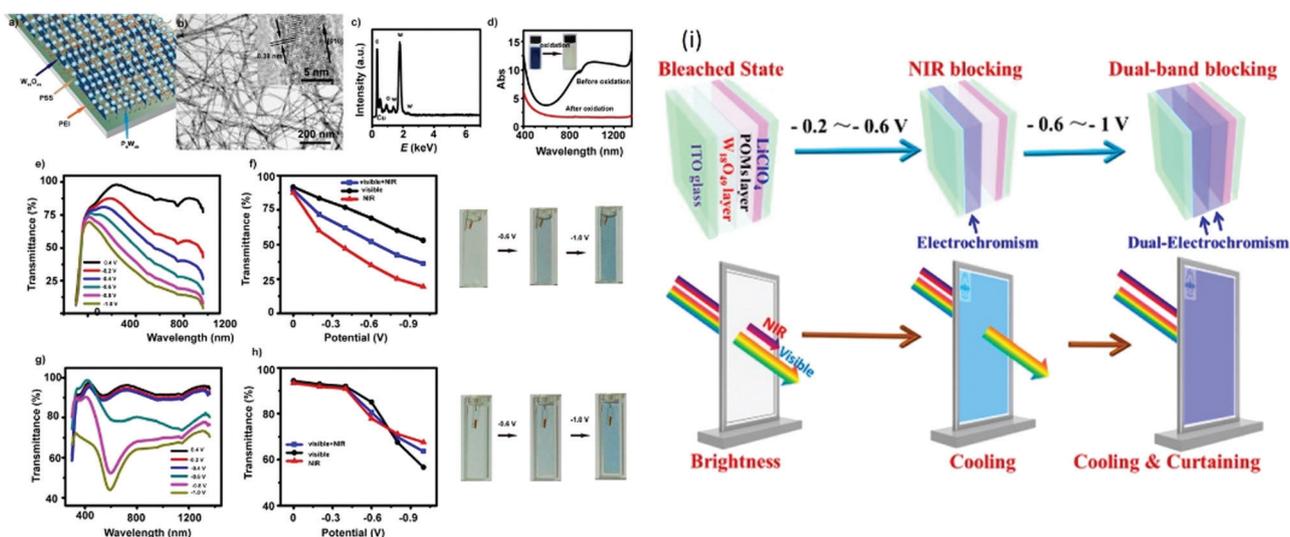


Fig. 13 Transmittance spectra of [PSS(PEI/PSS)₃(PEI/W₁₈O₄₉)₃₀/(PEI/P₈W₄₈)₂₀] film at different potentials, held for 4 min. (b) Relative transmittance change for [PSS(PEI/PSS)₃(PEI/W₁₈O₄₉)₃₀/(PEI/P₈W₄₈)₂₀] film under different applied potentials; blue curve: NIR + Vis region, red curve: NIR region, and black curve: visible region, and digital photographs of the film at 0.4 V, -0.6 V, and -1.0 V, respectively. (c and d) *In situ* optical response of [PSS(PEI/PSS)₃(PEI/W₁₈O₄₉)₃₀/(PEI/P₈W₄₈)₂₀] multilayer when switching between -0.60 V (100 s) and -1.0 V (100 s) to 0.4 V (200 s), where each step is measured at 1060 nm (c) and 500 nm (d). (e and f) Optical density variation as a function of charge density at 1060 nm (red curve) and 500 nm (black curve). The applied potential is -0.6 V (e) and -1.0 V (f) for 100 s. Corresponding schematic (i) shows the dual-band optical modulation. Reproduced with permission.¹²² Copyright©2018, ACS.



polymer to create electrostatic interactions for alternating layer deposition of a multilayered film (abbreviated as $(\text{PEI}/\text{W}_{18}\text{O}_{49})_n$ or $(\text{PEI}/\text{P}_8\text{W}_{48})_n$). The ITO glass was first activated *via* the LbL deposition of a polyelectrolyte multilayer of poly(sodium 4-styrene-sulfonate) (PSS) and PEI to increase the surface charges and surface affinity before the deposition of the $(\text{PEI}/\text{W}_{18}\text{O}_{49})_n$ or $(\text{PEI}/\text{P}_8\text{W}_{48})_n$ composite multilayer thin film. Due to the excellent stability of the films prepared using P_8W_{48} and $\text{W}_{18}\text{O}_{49}$ and their precise structural control, the nanocomposites film achieved the high coloration efficiency of $42.31 \text{ cm}^2 \text{ C}^{-1}$ at 500 nm and $287.49 \text{ cm}^2 \text{ C}^{-1}$ at 1060 nm (Fig. 13e–h). The switching speeds for the transparent, NIR-blocking and broadband-blocking were 86 s, 52 s, and 26 s, respectively. The electrochemical cycle stability of the composite film after 500 cycles at wavelengths of 1060 nm and 500 nm was only 3.20% and 2.40%, respectively. The obtained results provide an effective way for the formation of composite-based functional smart glass.

Liu *et al.*²⁵⁷ and Wang *et al.*¹¹⁹ demonstrated that ITO-based coating films are hard, which can be damaged and lose conductivity after a certain degree of bending of the device structures. They used Ag nanowires as a conductive substrate support and deposited $\text{W}_{18}\text{O}_{49}$ nanowires for EC device applications. The Ag nanowires and $\text{W}_{18}\text{O}_{49}$ nanowires were orderly assembled *via* the Langmuir–Blodgett (LB) method²⁵⁷ to develop several layers with elevated conductivity and transmittance. The Ag– $\text{W}_{18}\text{O}_{49}$ nanowire co-assembled thin film obtained *via* the LB method was continuously transferred to a PET substrate as an EC smart cell electrode, as demonstrated in Fig. 14. An increase in the number of layers of $\text{W}_{18}\text{O}_{49}$ nanowires significantly increased the coloration efficiency, cyclic stability, and transmissivity, as shown in Fig. 14b, c, f and g. The Ag– $\text{W}_{18}\text{O}_{49}$ co-assembled nanowire film could bend to a very small radius of 1.2 cm without any mechanical loss, which showed an excellent EC performance and mechanical stability (1000 cycles), as shown in Fig. 14d and e.

An Ag– $\text{W}_{18}\text{O}_{49}$ co-assembled film was used to develop a multilayered nanowire network on an ITO-PET film over a large-area conductive substrate with tunable conductivity ($7\text{--}40 \Omega \text{ sq}^{-1}$) and transmissivity (58–86% at 550 nm), as shown in Fig. 15a–e.¹¹⁹ The Ag– $\text{W}_{18}\text{O}_{49}$ – LiClO_4 /PMMA/PC-ITO-PET hybrid all-in-one solid-state portable and flexible EC device is shown in the colored and bleached states, which could be bent to a radius of about 1.2 cm with 90% resistance retention ($\Delta R/R \approx 8.3\%$) after more than 1000 cycles. Heo *et al.*¹²³ produced a low-temperature mesoporous $\text{WO}_{2.72}$ – NbO_x composite thin film on an ITO-PET supporting substrate, as shown in Fig. 16. Consequently, they realized a random/disordered arrangement of nanocrystal structures in organic media, which giving the opportunity to develop mesoporosity and overcoming the compact structure formulations (Fig. 16E and F). Ligand-stabilized $\text{WO}_{2.72}$ nanorods were synthesized using $\text{W}(\text{CO})_6$ and Me_3NO dissolved in oleylamine in a three-neck flask in an N_2 glove box and annealing directly at $300 \text{ }^\circ\text{C}$ for 90 min. After the wet-chemical reaction, the colloidal solution was washed using isopropyl alcohol (IPA) and redispersed in toluene, which were repeated several times to produce clean high aspect ratio $\text{WO}_{2.72}$ nanorods. They also synthesized low aspect ratio $\text{WO}_{2.72}$ nanorods by reducing the Me_3NO and oleylamine concentration, as shown in Fig. 16A. The obtained $\text{WO}_{2.72}$ may contain highly encapsulated ligands, which were stripped using Meerwein's salts (Et_3OBF_4). Et_3OBF_4 was dissolved in acetonitrile (AN) and mixed with $\text{WO}_{2.72}$ nanorods in toluene at a 1:1 ratio with constant stirring, centrifuged and then redispersed for repeated washing. Both the ligand-stripped and ligand-capped $\text{WO}_{2.72}$ nanorods were spin-coated on clean ITO-PET substrates using different dispersion media to form thin films of about 300 nm. The results showed that ligand-stripped $\text{WO}_{2.72}$ nanorods produced a thin mesoporous film due to the opposite electrostatic links, which prevented the spontaneous packing of the nanorods (Fig. 16E and F). After the fabrication of the $\text{WO}_{2.72}$ nanorod thin film, a guest component

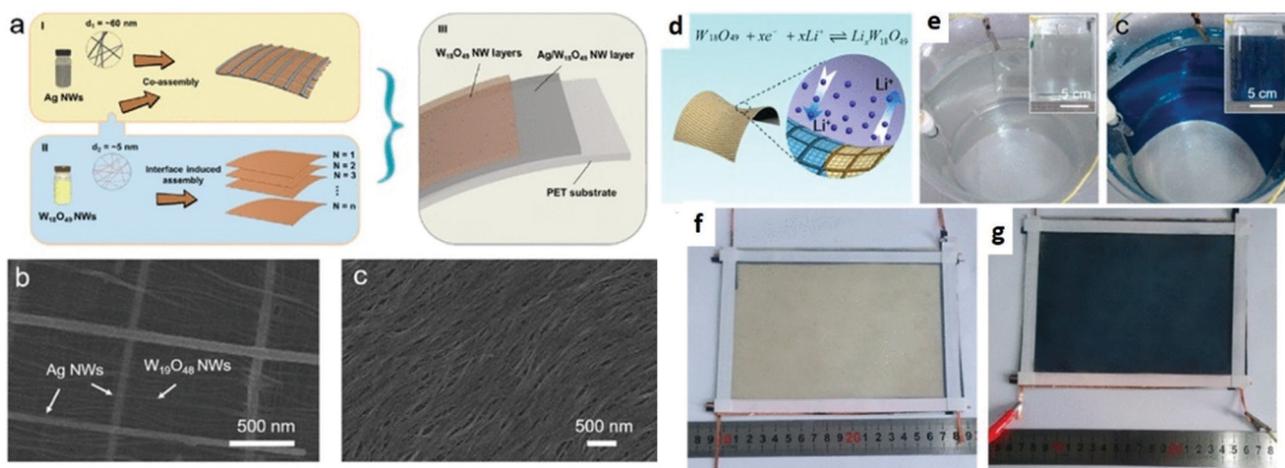


Fig. 14 (a) Schematic of the fabrication of a flexible device. (b and c) Images of Ag– $\text{W}_{18}\text{O}_{49}$ nanowires. (d–f) Coloration and bleaching of Ag– $\text{W}_{18}\text{O}_{49}$ nanowires film before and after an applied potential. Reproduced with permission.¹¹⁹ Copyright©2017, ACS.



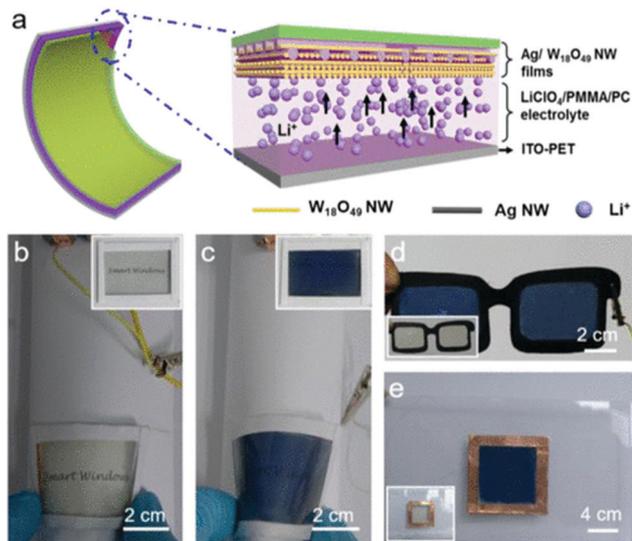


Fig. 15 (a) Schematic diagram of a solid EC device. Photographs showing the (b) bleached and (c) colored state of the EC devices in the bending state. The insets are photos without bending. (d) Photographs of the EC glasses model and (e) EC window model. Reproduced with permission.¹¹⁹ Copyright©2017, ACS.

solution composed of NbO₂POM clusters and [N(CH₃)₄]₆Nb₁₀O₂₈·6H₂O was drop-casted on the fabricated film, and then immersed in a formic acid and absolute ethanol mixed solution and heated at 150 °C. The WO_x-NbO_x composite film fabricated on ITO-PET has the advantages of selective control of the Vis-NIR light transmission, fast switching, and long cycle life (Fig. 16G and H).

Kim *et al.*⁴¹ designed WO_{3-x} nanocrystals and an amorphous niobium oxide nanocomposite mesoporous film. The obtained WO_{3-x}-NbO_x composite exhibited high charge-discharge kinetics and stability for about 2000 cycles. However, wet-synthesis of WO_{3-x} nanocrystals *via* the direct heating of a

mixture of tungsten salts without applied pressure requires harsh conditions, high temperature, and the product yield is small. Vuong *et al.*²⁹⁷ also employed similar procedures and reported the fabrication of a porous core-shell TiO₂-WO₃ nanowire structure film *via* sputter deposition using carbon nanotubes as a substrate and frequent thermal treatment. The particle size, morphology, electrochemical, and EC performance were studied and then compared with that of thin TiO₂-WO₃ composite films and single WO₃ and TiO₂ nanowires. The porous composite nanowires exhibited high proton intercalation capability with good reversible electrochemical cycles through intercalation and de-intercalation of the charge-compensating ions. Dulgerbaki and co-workers²⁹⁸ fabricated WO₃/PEDOT hybrid films and Li co-workers²⁹⁹ fabricated W_{0.71}Mo_{0.29}O₃/PEDOT:PSS hybrid film-modified electrodes, which exhibited highly improved EC performances, including fast switching kinetics, high coloration contrast, and tremendous coloration efficiency of >100 cm² C⁻¹ compared with the pure WO₃ films.

5.2. WO_{3-x}/metal/WO₃/other EC bilayer architecture films

To strengthen the transparency and EC functionalities, soft coatings as dielectric/metal/dielectric (DMD) blends have been appealing in recent years as alternatives to ITO-coated substrates.³⁰⁰ Various dielectric materials, for example WO₃, TiO₂, MoO₃, and ZnS, have been reported for the construction of DMD blends recently. The DMD assembly has also been extensively used in electrodes in organic optoelectronic devices, such as light-emitting diodes and organic solar cells.³⁰¹⁻³⁰³ In DMD coatings, the WO₃ film may perform a dual function, where it is anti-reflective for light in the transparent state and becomes absorbent in the colored state.^{293,304-307} Furthermore, DMD bilayer or trilayer films possess optimum resistivity (>15 Ω), high transparency (<80%) in the visible region, low thickness and superior flexibility.^{124,125,305-307} For dual functionality, the matrix design by simulation of D/M/WO₃ as glass/ZnS

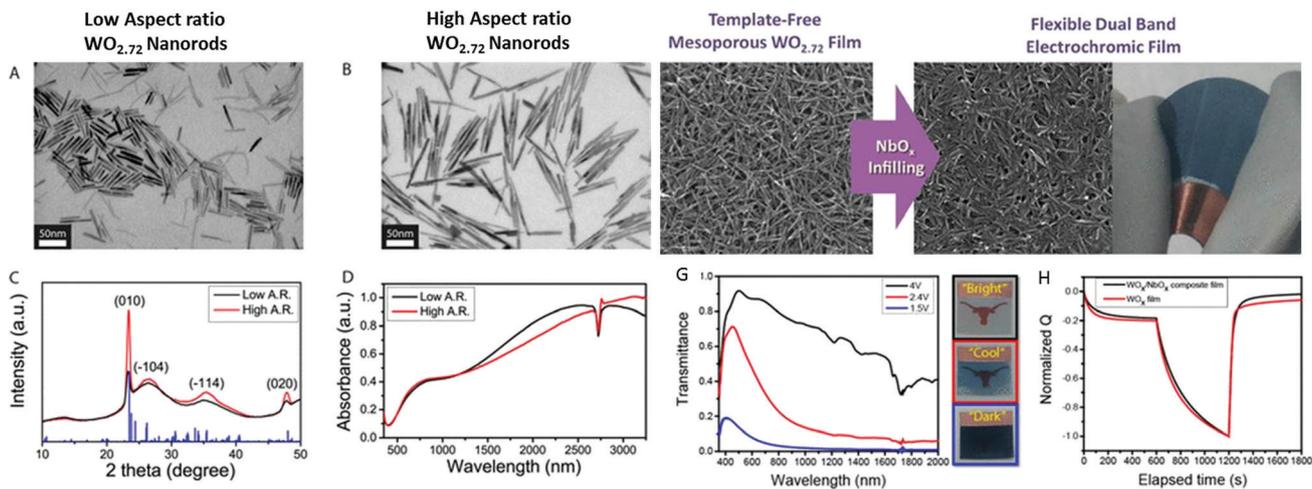


Fig. 16 (A and B) Low and high aspect ratio WO_{2.72} nanorods (top) and (C and D) their XRD and UV-vis absorbance spectra, respectively. Low and high aspect ratio (E) WO_{2.72} mesoporous film and (F) Nb-POM in-filled WO_{2.72} film (top). (G) Transmittance of composite film at various applied potentials and their corresponding photographs. (H) Charging and discharging profile at various potentials. Reproduced with permission.¹²³ Copyright©2017, ACS.



(34 nm)/Ag (15 nm)/WO₃ (326 nm) was formulated and verified via experiments by Leftheriotis' team.³⁰⁴ The color coating simulation studies showed the asymmetry of the radiation propagation path, which is absorptive in one direction and highly reflective in the opposite direction. They also substituted the layer of ZnS with an optimum thickness of WO₃ and the Ag layer with Au, and the addition of a 4th protective layer of Al₂O₃ demonstrated feasible replacements toward the optimum design. The film was produced via electron-beam gun deposition with a transparency of 57% and sheet conductivity of 9 Ω. The obtained results showed that the outer WO₃ coating layer should be thick enough to protect the Ag nanolayer from the liquid electrolyte. The optical contrast was 90 cm² C⁻¹ and a coloration efficiency of 30 cm² C⁻¹ was recorded for 500 coloration/bleach cycles. The influence of liquid electrolyte on the EC coatings was observed with an increase in resistance and stability due to degradation, where the Ag nanolayer degraded because of the interactions of the electrolyte through the pores of the coated ZnS layers.³⁰⁸ Hence, a layer thickness of WO₃ of up to 300 nm was adequate for the protection of the Ag nanolayer over the ZnS/Ag/ZnS stack.

Koubli *et al.*³⁰⁶ employed electron beam deposition for the formation of a WO₃/Ag/WO₃ coating film. The fabricated film exhibited excellent deep coloration. However, with the intercalation of electrical charges, the film was unstable and damaged with the reversal of electrical potential. It became dysfunctional in the bleach state due to the reaction of Ag with Li⁺ ions, and the resistance of the Ag layer increased. Hence, for safe mode operation, the charge density was lowered to 12.6 mC cm⁻² from 15.7 mC cm⁻². Li *et al.*³⁰⁵ demonstrated that a WO₃/Ag/WO₃ film could sustain up to 3000 cycles without EC performance degradation, as shown in Fig. 17. The poor adhesion and rapid degradation regarding color/bleaching of the bifunctional WO₃/Ag/WO₃ film were improved by sputtering

deposition using metal targets.¹²⁵ They successfully added an ultrathin tungsten sacrificial layer followed by an external WO₃ layer. Metal-to-metal bonds are generally stronger than metal-to-oxygen bonds. Therefore, the Ag-to-W layer film was adjusted to the desired thickness and the resulting film in the 2000th bleaching cycle showed a stable EC performance and avoided oxidation of the silver metal nanolayer in the oxygen atmosphere. Based on this strategy, the oxidation of silver metal (Ag⁰) can be eliminated and the conductivity of the film can be lost. Moreover, according to the conventional DMD three-layer structure, the introduction of a sacrificial W layer presents complexity to the deposition process in the form of a fourth structural layer in the device configuration.

Najafi-Ashtiani *et al.*³⁰⁹ used high-power impulse magnetron sputtering (HiPIMS) instead of DC/RF magnetron sputtering technology to produce a DMD film without a fourth sacrificial metal layer, as illustrated in Fig. 18. The HiPIMS EC deposition method has a wide range of use in industry and production. Converting a traditional sputtering system to HiPIMS operation only requires the application of a suitable power supply to derive the target in the HiPIMS mode.³¹⁰ The high instantaneous power supply produces high-density plasma of the sputtered target material with a high ionization degree.³¹⁰ The limited supply of oxygen is promising for the controllable growth of WO₃ nanoparticles and external layer deposition to reduce the oxidation of the Ag layer.³⁰⁹ The thickness of WO₃ (30 nm)/Ag(10 nm)/WO₃ (50 nm) was optimized with the sheet resistance of ~23 Ω and transmittance of 80% (Fig. 18a). The fabricated coating exhibited an EC performance with good transparency, excellent cycling stability for 2500 cycles, fast switching kinetics, and Vis-NIR light modulation (Fig. 18b).

The EC performance of DMD films is not satisfactory in terms of switching time, switching speed, optical modulation, and durability. To this end, the pioneering work of

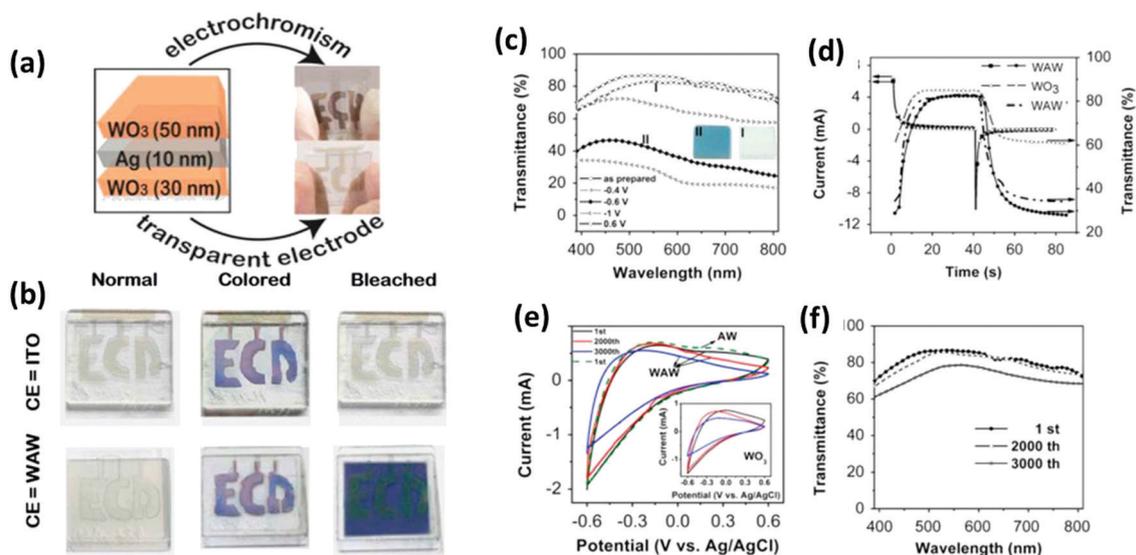


Fig. 17 (a and b) Different layer thickness of WAW (30/10/50 nm) used as an electrode and their corresponding color and bleached states. (c) Transmittance variations with a change in voltage (0.6 to -0.6 V). (d) Current and optical contrast with step potential. (e) CV curves and (f) transmittance variations with different cycles. Reproduced with permission.³⁰⁵ Copyright©2015, Elsevier.



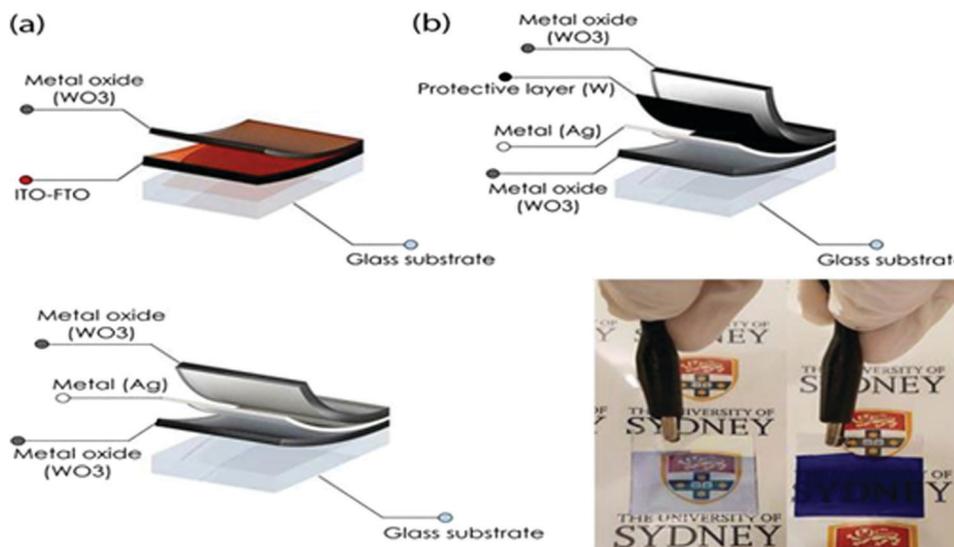


Fig. 18 HiPIMS-sputtered EC $\text{WO}_3/\text{Ag}/\text{WO}_3$ (DMD) three-layer structure. (a) WO_3 deposited on ITO-FTO-coated glass. (b) Transparent, conductive DMD structure deposited on glass without ITO/FTO with a tungsten (W)-protective layer on the Ag layer. (c) DMD structure (ITO-FTO free) fabricated via HiPIMS without a protective layer. Optical images of bleached and colored states of the DMD (30 nm/10 nm/50 nm) coating. Reproduced with permission.³⁰⁹ Copyright@2019, ACS.

Dong *et al.*¹²⁴ proved the application of mixed oxides of W–Mo-encapsulated Ag such as an $\text{MoO}_3\text{--}\text{WO}_3/\text{Ag}/\text{MoO}_3\text{--}\text{WO}_3$ trilayer film for ITO-free EC devices, as demonstrated in Fig. 19. The film thickness was optimized at 40 nm, resulting in the sheet resistance of 9Ω with a fast-switching speed and high coloration efficiency ($70 \text{ cm}^2 \text{ C}^{-1}$).

5.3. Doped $\text{WO}_{2.72}$ nanostructures

Using dopants or intended impurities to switch the behavior of nanomaterials is at the heart of various key technologies. For semiconductors, the doping process is crucial, otherwise they would be electrically insulating. Certain applications such as wavelength-tunable lasers,³¹¹ bioimaging,³¹² photocatalysis, and solar cells³¹³ may eventually rely on the doping of nanocrystals to tune their electrical properties. Doping is an effective method to tailor the bandgap of semiconductor materials by introducing a transition state (defect band) between their valence bands and the conducting bands. Moreover, dopants can strongly alter the electronic, optical, and magnetic properties of bulk semiconductors.³¹⁴ The EC efficiency of $\text{WO}_{3-x}/\text{WO}_3$ has been tested by doping³¹⁵ with Mo, Nb, Ti, Ni, Al, Ag, Pd, Cu, Au, *etc.* The use of impurity atoms to tune the conductivity of WO_{3-x} can be better understood by considering (i) substitutional doping and (ii) interstitial doping.

5.3.1. Substitutional doping. Substitutional doping plays a key role in tuning the electrical conductivity in semiconductor materials, where the dopant atom replaces an atom in a nanocrystal, thereby integrating discrete components into circuits and devices.³¹⁶ Semiconducting nanocrystals possess low free carrier concentrations, and thus do not exhibit LSPR phenomena. Accordingly, substitutional dopants with one additional valence electron can be thermally ionized and donate their extra electron to WO_{3-x} (n-type doping). Similarly,

dopants with one fewer/deficient valence electrons can offer an extra hole (p-type doping). These electrons/holes acting as charge carriers can support LSPR in the NIR region, thereby increasing the electrical conductivity. For conducting thin-film technology, the capability to introduce these carriers is necessary, as published for tin-doped indium oxide (ITO),³¹⁷ aluminum-doped zinc oxide (AZO),³¹⁸ indium-doped cadmium oxide (ICO),³¹⁹ and niobium-doped anatase titania.³²⁰ Hasani *et al.*³²¹ doped CdSe quantum dots with WO_3 nanocrystals *via* wet-synthesis followed by annealing. The optimized CdSe QD-doped WO_3 showed an excellent EC performance with a coloration efficiency of $112.3 \text{ cm}^2 \text{ C}^{-1}$. Ni and Ti doping has been reported recently to introduce oxygen vacancies, crystal defects, and nanostructural changes in WO_3 for improved EC properties.^{322,323}

Mo, Cu, and Nb are considered effective sensitizers, increasing the oxygen vacancies and crystal defects. Thus, the absorption peak position of the films shifts to a low wavelength range (blue shift). Moreover, they can also accelerate the reversible switching kinetics of the film.^{12,324} Al ion doping also adds advantages based on the shape, size, and structural variations.³²⁵ Li and Ru doping successfully enhanced the electrochemical performance, cycle stability, and diffusion coefficient of WO_3 -based thin films.^{77,326} As the size of nanocrystals decreases, their energy increases, and thus the carriers in the nanocrystals must occupy any confined electronic states.³²⁷ For robust/stable conductivity, adjacent nanocrystals should be uniform in size to have resonant electron states, and eventually, be narrowly spaced to confirm their quantum-mechanical overlap. Yao *et al.*³²⁸ synthesized Cr-doped WO_3 polyhedra by controlling the parameters of RF thermal plasma. The results demonstrated that the 2.5 at% Cr-doped-layer could reduce the crystal growth rate of WO_3 along the $\langle 001 \rangle$, $\langle 010 \rangle$, and $\langle 100 \rangle$ planes.



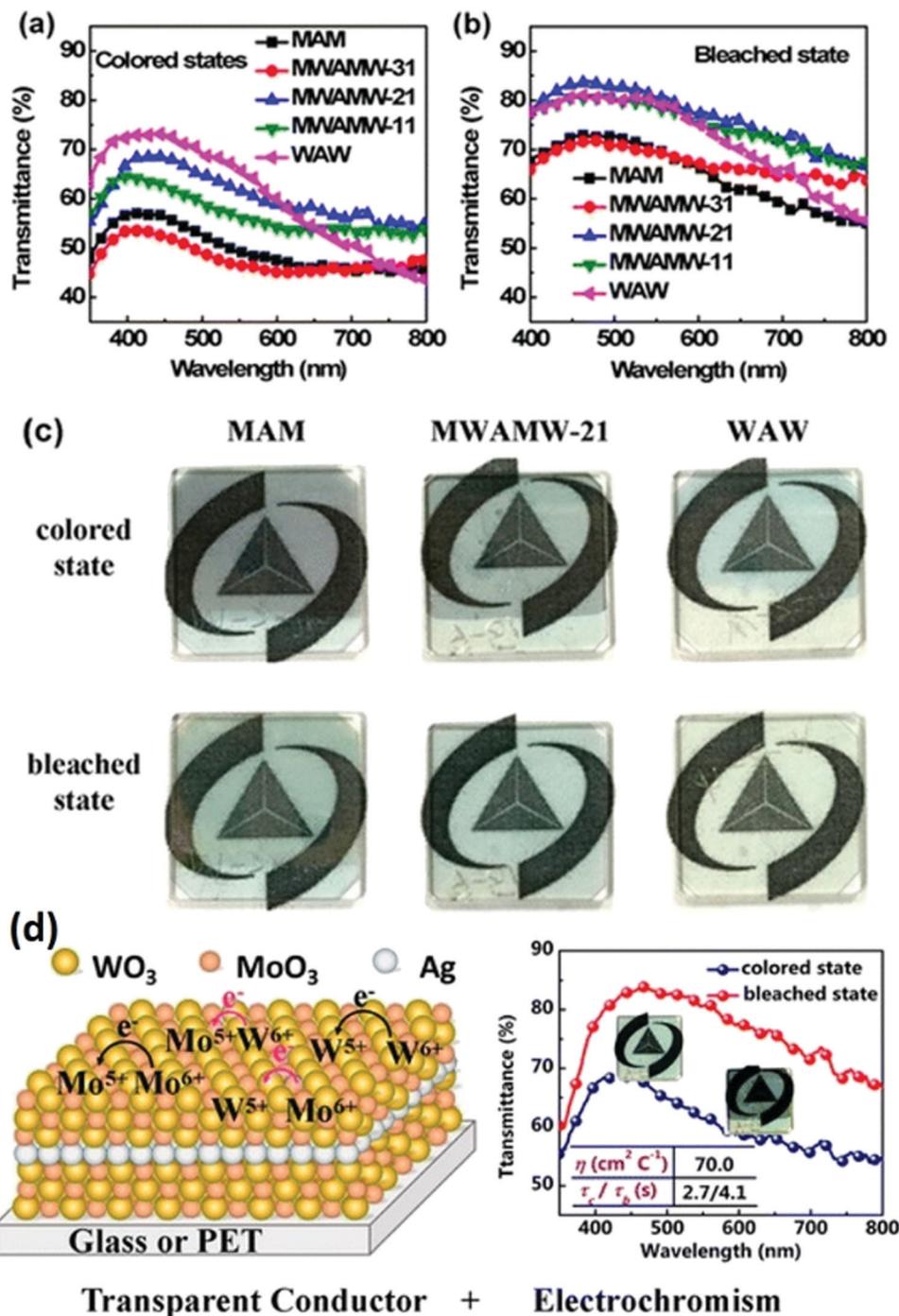


Fig. 19 Optical transmittance of MAM, MWAMW, and WAW films in (a) colored and (b) bleached states at potential steps from -0.50 to 0.55 V (vs Ag/AgCl) with a 20 s interval. (c) Pictures of MAM, MWAMW and WAW films in the colored and bleached states. (d) Schematic of MWAMW composite deposited on PET or glass and corresponding color and bleached state optical transmittance. Reproduced with permission.¹²⁴ Copyright@2016, ACS.

Higher carrier concentrations are required for superior conductivity. However, nanocrystals with additional free electrons and holes can also be used as strong reducing/oxidizing agents. Therefore, the electrochemical reactions on the surface can consume the charge carriers,³²⁹ and the carriers will not participate in electrochemical reactions. Thus, avoiding these reactions of films fabricated using doped nanocrystals is a big

challenge! The n-type and p-type doping of CdSe quantum dots using In, Sn, Al, or Ag as dopants have been a research focus, as illustrated in Fig. 20a and b.^{330,331} The critical temperature affecting the doping processes can be summarized as follows.³³² (i) surface adsorption, (ii) lattice incorporation, (iii) lattice diffusion, and (iv) lattice ejection (Fig. 20a and b). It is noteworthy that each nanocrystal can be switched, but it may



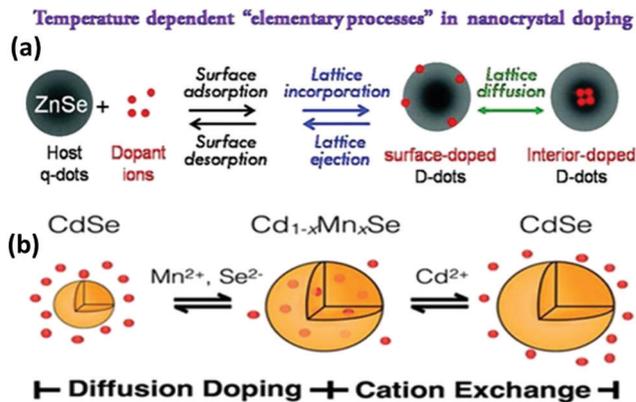


Fig. 20 (a) Temperature-dependent surface adsorption, lattice incorporation, lattice diffusion, and lattice ejection doping in nanocrystals.³³² (b) Mn^{2+} diffusion doping of CdSe nanocrystals and Mn^{2+} expulsion from $\text{Cd}_{1-x}\text{Mn}_x\text{Se}$ nanocrystals via cation exchange with Cd^{2+} .³³² Reproduced with permission. Copyright © JACS.

be impossible to dope to achieve the desired doping characteristics.

5.3.2. Interstitial doping. Interstitial doping is a principal means for new atoms to be incorporated within the crystal structure because of the atomic vacancies/defects within the lattices. Interstitially doped tungsten oxides, which are known as tungsten bronze, have the formula M_xWO_3 , where M is Li^+ , Na^+ , Cs^+ , or Rb^+ .³³³ Tungsten bronze materials retain high visible light transmittance, while exhibiting high NIR light absorption, making them ideal candidates for spectrally selective optical devices. A high doping level/degenerate doping in semiconductor metal oxides can effectively modulate the NIR wavelength transmittance through electrochemical charging and discharging of the free electrons responsible for LSPR absorption.^{41,294} It has been found that solar NIR light is the most intense at shorter wavelengths (700–1300 nm), and thus nanocrystals with LSPR absorption in this spectral range will be ideal for dynamic solar light control.⁴¹ Due to their outer-d valence electrons, WO_{3-x} nanoparticles can be ideal candidates with intense visible and NIR light absorption, where their metallic characters exhibit tunable LSPR peaks.¹³⁹ Therefore, interstitially doped/oxygen vacancy-doped nanocrystals can achieve metallic characters with LSPR modulation in the NIR region, besides the substitutionally doped transparent conductive oxides, e.g., ITO/AZO,^{41,139,294,334} used for tuning the LSPR absorption in the IR range (1400–3000 nm). When dopants are incorporated in these nanocrystals, they can emerge as new advances for certain well-known applications and effectively predicted, as found in the literature.^{294,331,334–338}

To improve the energy saving proficiencies of EC smart windows, it is necessary to maximize the plasma resonance displacement and modulations of NIR rays. For example, in ITO, a high doping concentration shifts its LSPR peak to the high frequency spectrum. In contrast, a decrease in the nanocrystal size to $\sim 2\text{--}5$ nm in accordance with an increase in optical modulation remains enigmatic. Smaller nanocrystals have a higher surface charge density, which facilitates high

capacitive behavior and allows more electrons to be injected. Therefore, smaller size and highly doped nanocrystals will exhibit improved EC performances in the desired multifunctional EC smart windows. Because of the impurity atom doping and higher formation energy of defective nanocrystals, nanocrystals tend to expel dopants during annealing to minimize their energy and undergo self-purification/healing (Fig. 20).³³⁹ It was found that surface adsorption and diffusion³³⁹ can be influenced by three main factors: (i) surface morphology, (ii) nanocrystal shape, and (iii) surfactants in the growth solution. Based on the kinetic control, dopants can be incorporated in the host nanocrystals if they can bind and reside on the host crystal surface. Diffusion of impurities into the host nanocrystals may occur on the surface of nanocrystals. It was also found that nanocrystals contain active surface sites and the binding energy of the impurities adsorbed on the nanocrystals determine their residence time.³³⁹ The diffusion doping strategy is based on incorporating relatively incompatible anions/cations in semiconductors, which is driven by thermodynamic and high-temperature cation randomization. For, example $\text{Cd}_{1-x}\text{Mn}_x\text{Se}$ has been studied (Fig. 20b).³⁴⁰ The slow diffusion kinetics of Mn^{2+} ions and high-temperature treatment possibly accelerate their diffusion, although it is associated with self-purification due to the diffusion of Mn^{2+} ions out from the nanocrystals.³³²

It is worth noting that only effective doping that successfully incorporates dopants in the host crystal lattice rather than adsorbed on its surface can alter the characteristics properties of nanocrystals; however, plasmonic nanocrystals are quite interesting in this regard.^{314,340} Au and Ag are believed to have strong surface-enhanced Raman spectroscopy (SERS) performance. They have no signals upon Raman scattering but can produce strong LSPR under irradiation of light. Because Au couples with Ag substrates, and then non-noble metals, metal oxides, graphene, silicon-metal and metal-tellurides are also considered to become a new type of SERS materials due to the charge transfer mechanism. Liu and coworkers³⁴¹ reported that TiO_2 and WO_{3-x} do not have free electrons because of their wide bandgaps, indicating that these semiconductors will not show the LSPR effect. However, their conductivity can be greatly enhanced due to the introduction of dopants into their large vacancy/defect sites, given that the dopant materials have the LSPR effect due to the free electrons found in their d-orbital, as shown in Fig. 4b and 21. However, the SERS induced due to the LSPR in TiO_{2-x} and WO_{3-x} cannot be generated due to the presence of oxygen vacancies and crystal defects. This can be attributed to the difference in excitation wavelength between NIR and SERS, which are found in the visible region. However, the conductivity (charge transfers) greatly increases due to the vacancy-doped features between the substrates and the adsorbed molecules. Therefore, if the LSPRs of vacancy-rich materials can be adjusted from the NIR region to the visible region (near the SERS excitation wavelength), then due to the charge transfer process, their SERS activity can be further improved. The high density and oriented $\text{W}_{18}\text{O}_{49}$ ultrafine nanowire bundles with an abundance of oxygen vacancies



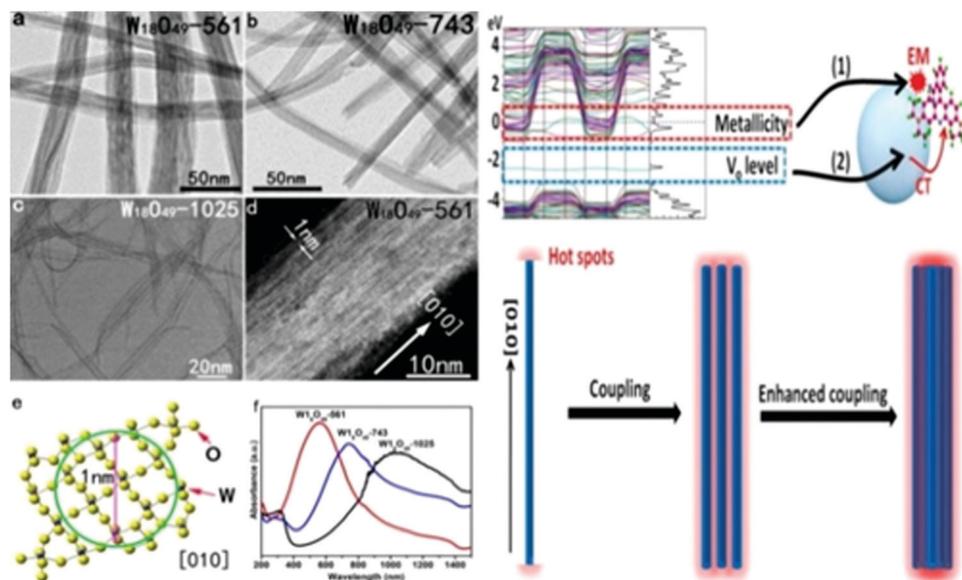


Fig. 21 (a–c) TEM images of $W_{18}O_{49}-561$, $W_{18}O_{49}-743$, and $W_{18}O_{49}-1025$, distinguished due to various in their LSPR peaks. (d) HRTEM image of $W_{18}O_{49}-561$. (e) Cross-section of a 1.0 nm nanowire inside of one $W_{18}O_{49}$ unit cell oriented along the [010] direction. (f) UV-visible-NIR absorption spectra of the $W_{18}O_{49}-561$, $W_{18}O_{49}-743$, and $W_{18}O_{49}-1025$. (left) Top: Band structure of $W_{18}O_{49}$ and the two possible SERS enhancement ways in $W_{18}O_{49}$: (1) EM and (2) CT. Bottom: LSPR coupling between nanowires. Adopted from ref.341 Copyright©2018, ACS.

and defects have strong LSPR due to the charge transfer and electromagnetic enhancement factors. With an increase in thickness due to the aggregation of the nanowires, the absorption peaks gradually change from the NIR to visible region (Fig. 21). However, the XRD peaks remain almost the same, which was concluded to be due to electromagnetic oscillations found in the highly thin $W_{18}O_{49}$ nanowire bundles. These pioneering results may reveal SERS properties.³⁴¹ The SERS enhancement mechanism can be attributed to the synergistic effect of the LSPR-SERS coupling that occurred among the ultrathin oriented nanowires, defects, and oxygen vacancy, which significantly improves the charge transfer process.³⁴¹

5.4. Plasmonic surface-decorated $WO_{2.72}$ composites

Plasmonic nanocrystal-modified interfaces significantly affect the optical absorption properties of nanomaterials. Noble metals nanocatalysts, such as Au, Pd, Pt, Ag, Rh, and Cu, are promising for improving the optical properties *via* LSPR effective modulations and Schottky junction formation. This is because the difference in the work functions of noble metals and the semiconductor metal oxide ($WO_{2.72}$) induce electronic sensitization, affecting their electrical properties. The rectifying Schottky barrier at the interface enhances the electrical transport properties. Thus, a higher work function and Schottky junction formation can significantly facilitate the separation of photogenerated electrons and holes. Ordered-shaped plasmonic nanocrystals such as nanorods (1D), sheets (2D), cubes, and triangular prisms (3D) possess sharp edges and corners, and hence exhibit excellent LSPR and considerable electromagnetic field enhancement.

Plasmonic-mediated nanocrystals can be classified into four categories as follows: (i) isotropic/bulk, (ii) assembled, (iii)

responsive, and (iv) interfacial plasmonic nanocrystals. In bulk/isotropic plasmonic nanocrystals, the nanocrystal composite is monodispersed throughout the matrix compared to the bulk in a medium. The dielectric matrix may cause an insulating barrier between the plasmonic nanocrystal candidates or may also contribute to their optoelectronic properties. In assembled nanocrystals, the plasmonic component species are arranged into groups, clusters or certain anisotropic areas in the dielectric matrix. This self-assembly causes the nanoparticles to acquire strong plasmonic coupling due to the formation of nanojunctions and their collective optical properties. New research is devoted to reconfiguring the capability of plasmonic nanocrystals to modulate external stimuli that trigger a high optical response. Thus, the responsive plasmonic nanocrystals can acquire tunable and unique optical functionalities.¹⁰ It is noteworthy that the gaps between the nanocrystals often have an impact on their optical properties. The interfacial plasmonic nanocrystals connect the interfaces of the metal–dielectric (M–D), dielectric–dielectric (D–D) or dielectric–metal–dielectric (D–M–D). These composite architectures are frequently 2D/3D structures. Electron injection in 2D/3D plasmonic nanocrystal composites results in a shift towards the high-frequency region and optical modulation in the NIR region of the solar spectrum. Plasmonic composites are specially engineered to acquire optoelectronic properties different from their bulk components. The blend symmetries of plasmonic-incorporated particles can be understood based on Fig. 22, and detailed explanations can be found in the chemical review published by Su-Wen Hsu *et al.*³⁴²

The Vis-band optical modulation of WO_{3-x} can be shifted towards the NIR wavelength due to the coupling of traditional metals/cation intercalation, resulting in NIR-LSPR extinction.



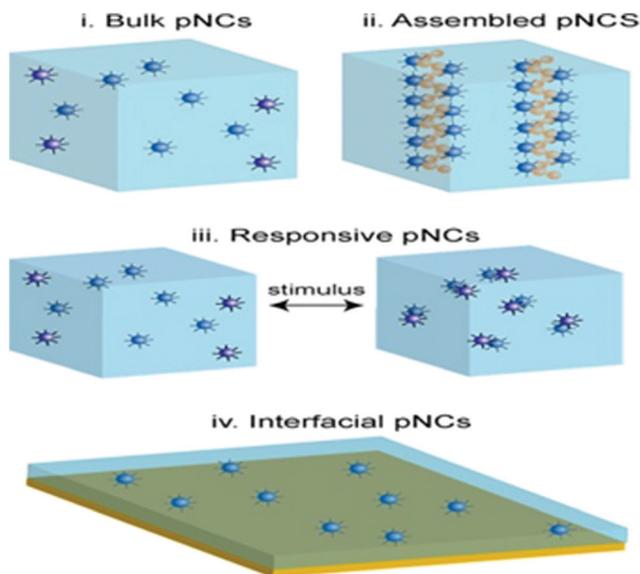


Fig. 22 Schematic of the various classes of pNCs. Reproduced from ref. 342 Copyright©2018, ACS.

Thus, the surface decorating/synthetic doping of noble metals in WO_{3-x} nanocrystals should be capable of independent modulation of spectral bands in both the Vis and NIR regions within a single-component functionalized film. Thus functionalized WO_{3-x} can be capable of dual-mode optical modulation as a nonradiative material to achieve multifunctional smart windows for controlling visible light transmittance, photo-thermal conversion of NIR solar loads, and also eliminating microorganisms living on windows.^{67,293,343,344} The growth of plasmonic nanocrystals on metal oxide layers can support the tunable LSPR, which will interact with the solar spectrum to shift the plasmon resonance for color changes.⁴² Moreover, Li-ion intercalated WO_{3-x} can be used for energy storage applications, such as supercapacitors and batteries.^{133,345} Improving the slow diffusion kinetics of Li ions and the conductivity of bulk WO_{3-x} can improve the charge capacity and charging capability of WO_{3-x} -based electrodes. Assuming that surface-modified WO_{3-x} can be charged, its charge capacity is almost equivalent to that of lithium insertion reactions in ultra-small particles of about 10 nm.^{42,346} Besides decreasing the size of WO_{3-x} , doping can also improve the electrochemical properties of WO_{3-x} electrodes (see Section 5.3) to boost their inherent low conductivity. Furthermore, surface-decorated nanocrystals with a diameter of <10 nm can support LSPR, which can be tuned through the Vis to NIR region by altering the size, shape, composition, and Fermi energy level of the nanocrystals, as reported in the literature.^{67,347,348} Au, Ag, and Pd nanoparticles may exhibit a strong, robust, stable, and tunable LSPR absorption band at both visible and NIR absorption wavelengths.^{69,349} Au and Pd nanoparticles with a spherical shape and size of ~4–6 nm were deposited on the surface active sites of WO_3 nanowire bundles in HAuCl_4 and PdCl_2 solutions with additional oxidation^{67,149,350} (Fig. 23). In this scenario, the absorbed NIR light could subsequently be

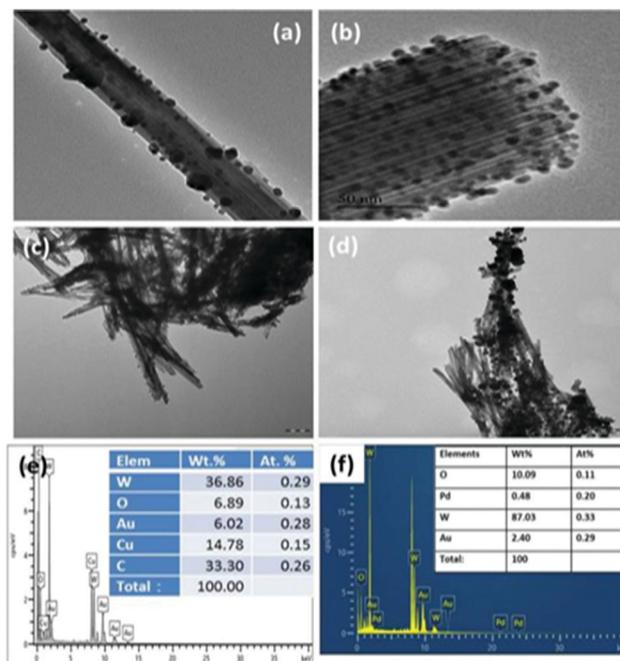


Fig. 23 (a–d) TEM and HRTEM images displaying WO_{3-x} nanorods: nanowire assembly decorated with Au (a and b), Pd (c), and (d) bimetal Au, Pd NP-decorated WO_3 . (e and f) Corresponding EDS elemental mapping. Reproduced with permission.¹⁴⁹ Copyright©Elsevier.

converted to heat energy due to the electron–phonon/phonon–phonon interactions. Au NP-decorated WO_3 may enhance the catalytic and photocatalytic activity and exploit the LSPR in Au NPs. The extensive work functions of $\Phi(\text{Au}) = 5.1$ eV, $\Phi(\text{Pd}) = 5.5$ eV, and $\Phi(\text{WO}_3) = 4.8$ eV may induce electronic sensitization and trap photo-excited electrons. The LSPR in gold-decorated WO_{3-x} has been applied in photothermal ablation therapy,³⁵¹ biological sensors,^{352,353} gas sensors,^{354–356} and optical detection.³⁵⁷

Xu *et al.*²⁹³ fabricated an EC-photothermal dual-band composite film of plasmonic Au anchored with 3D honeycomb-like WO_3 , as shown in Fig. 24a–d. The obtained composite film was tested for the optical transmission and photothermal conversion of visible and NIR light and reducing the attack of microorganisms (Fig. 24e). The Au LSPR and broadband non-radiative plasmon decay are assumed to be tunable as an electric field on the surface of the WO_3 substrate. The photothermal conversion was attained in a colored state, which is ascribed to the coupling of traditional visible-band optical switching with the NIR-LSPR extinction, as given in Fig. 24f and g. The as-prepared EC and photothermal films were also examined and proven to be effective in reducing the risk of microorganisms such as *E. coli* bacteria. Therefore, photothermal EC film-coated glass as an advanced multi-functional smart sterile window has broad application prospects.

Moreover, Ag nanoparticles were distributed isotropically within the bulk polytetrafluoroethylene (PTFE) matrix through direct current (DC) or co-sputtering of Ag and PTFE to leverage the LSPR extension of the Ag nanoparticles in the visible



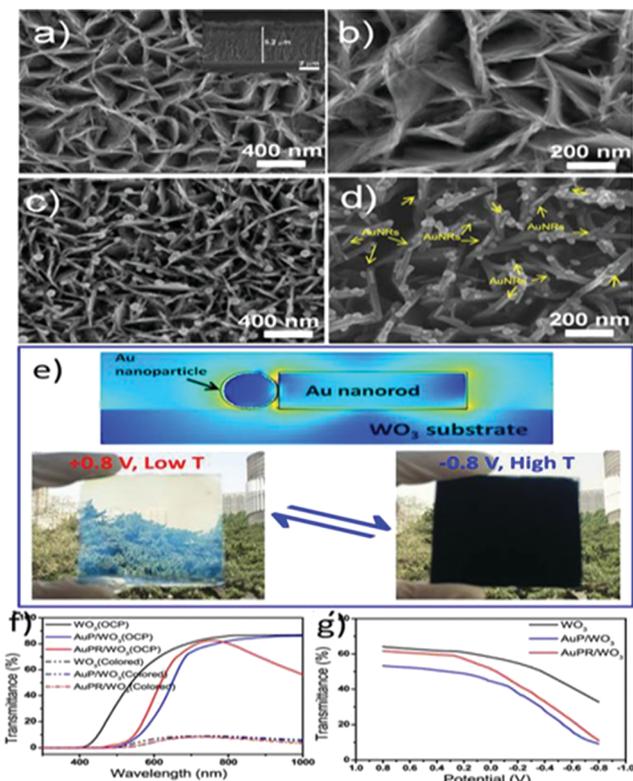


Fig. 24 SEM images (a) top view and (b) magnified view of a blank WO_3 film; (c) top view of Au nanoparticle-decorated WO_3 film; and (d) top view of Au nanoparticle- and nanorod-decorated WO_3 film. (e) Schematic of Au nanoparticle-attached nanorods, and digital photographs of the AuPR/ WO_3 film on FTO in the bleached and colored state. (f) Transmission spectra of the WO_3 , AuP/ WO_3 , and AuPR/ WO_3 film-coated FTO slides at OCP and colored states. (g) Relationship of transmittance at 915 nm with an applied electrochemical potential at a scan rate of 10 mV s^{-1} . Reproduced with permission from ref. 293 Copyright@2018, ACS.

range.^{342,358} The deposition process is important for developing flexible substrates that utilize matrix/nano-inks and target nanoparticles with a very small size. The plasmonic nanoparticle-based composite-generated film on the flexible substrates not only selectively controls the solar spectrum as an EC device but also has retrofit installation features.³⁵⁹ During *in situ* synthesis, the small size and morphology, and aspect ratio of plasmonic nanoparticles can cause impediments. Accordingly, plasmonic nanoparticle-polymer blends can successfully overcome this disadvantage *via* the incorporation of a polymer matrix in the already produced metal nanoparticles. Therefore, each material component of the plasmonic nanocrystals/composite can be synthesized independently and combined with the prospective polymer blend to generate a multifunctional nanocomposite.^{360–362}

Nanoscale WO_{3-x} particle decoration is advantageous for the fabrication of plasmonic nanocrystal-composites, which prevents undesirable aggregation or may be beneficial for phase separation. Similar to NiO, WO_3 , TiO_2 , and VO_2 , the color and optical transmittance can be tailored by external stimuli, including an electrical field, light, and temperature.¹⁰

In these scenarios, the nanocrystal components are grafted with polymer templates, which are miscible in a polymer matrix, *i.e.*, dispersion of fullerenes in polystyrene.³⁶³ Similarly, they can also be grafted through the large gyration radius of linear polymers, *i.e.*, polyethylene nanoparticles in polystyrene,³⁶³ and/or tethered with soft/hard templates having a high surface area, which endows the highest possible stability to the nanocrystal composite-polymer templates.³⁶⁴ Au nanorods were grafted with polystyrene polymer, and then deposited as a thin film.³⁶⁵ The thin film showed an optical response at 780 nm, which is reliable with the excitation of the longitudinal dipolar LSPR modes. The LSPR resonance function showed a blue shift in the optical absorption, resulting from the long linear-polymer templates and the side-by-side assembly of the aligned nanorods. In addition, the plasmonic EC film-based electrodes with nanocrystals in a disordered packing¹²³ generated open network channels, which increased the surface area and could be easily accessed by the electrolyte. In this area of research, the Milliron team demonstrated the fabrication of porous films and the effect of the nanopore sizes obtained by using amphiphilic block copolymer micelles to bind ligand-stripped ITO nanocrystal architectures into an ordered/more alignment conductive mesoporous network.³⁶⁶ The mesoporosity was controlled by altering the molecular weights of the hydrophilic/hydrophobic block polymers, with the pore size in the range of 29–37 nm and thicknesses of 9–13 nm.³⁶⁶ The ordered mesoporous structures resulted from the interaction between the ligand-stripped nanostructure surface and hydrophilic block-copolymers.³⁶⁷ Increasing porosity increases the speed of ion transport and switching through the surface of the nanocrystals.^{347,367}

In addition, the controllable size, quality, and assembly structure of plasmonic nanocomposites can be integrated into traditional multi-functional smart windows, where each component can be activated independently for specific functions. The independent modulation of visible and NIR light will stimulate the window to function in hot/cold climates to respond perfectly to changes in weather conditions. Recent research relies on plasmonic nanocomposite architectures for visible and NIR-selective transmission, enabling dual-band functionality to meet the challenges of traditional smart windows.^{122,123} Dahlman *et al.*⁴² fabricated a single-component colloidal aliovalent niobium-doped anatase TiO_2 nanocrystal film for dual-band optical modulation. The Nb- TiO_2 -deposited film showed LSPR in the NIR region and could modulate light with capacitive charging. A variation in the composition, size, shape, and doping in the TiO_2 nanocrystals greatly affected the efficiency of the EC film. The anatase TiO_2 has low electrical conductivity and slow ion diffusion, which was improved by composite formation and impurity doping.

Moreover, multifunctional integrated devices have received tremendous research interest due to their low cost, lightweight, flexibility, and for monitoring health and the environment. Recent research achievements such as gas sensors, strain sensors, and photodetectors have been reported in various self-powered multifunctional integrated systems, thus reducing



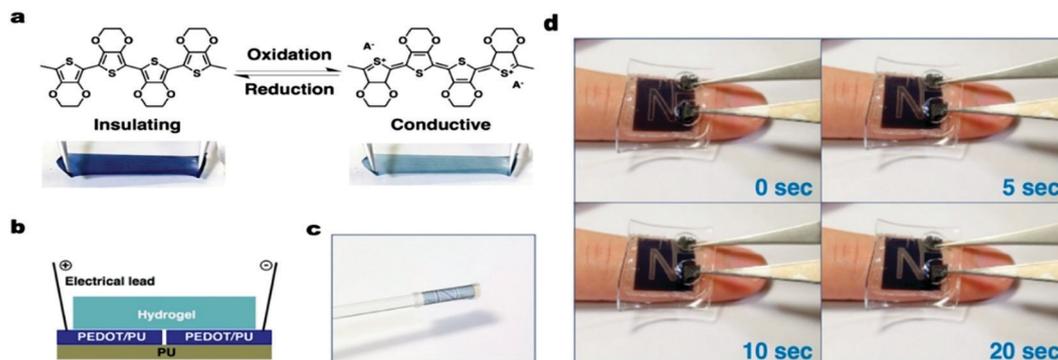


Fig. 25 Stretchable EC film of PEDOT/PU. (a) Molecular structures of PEDOT and photographs of the stretched EC film of a PEDOT/PU composite in the reduced and oxidized states. (b) Schematic illustration of an EC film/hydrogel hybrid. (c) EC film/hydrogel hybrid wrapped around a glass rod and human finger. (d) Color change in the EC film/hydrogel hybrid device placed on a human finger. Reproduced with permission.³⁷⁵ Copyright©2017, ACS.

energy cost by cooling, heating, and lighting.^{68,293,368} For example, flexible supercapacitors have been reported together with photodetectors based on CdSe nanowires,³⁶⁹ Co₂O₃ nanowires on Ni fibers with graphene,³⁷⁰ TiO₂ nanoparticles,³⁷¹ and TiO₂/MXene heterostructures³⁷² deposited on different substrates. The resultant flexible all-solid-state integrated systems exhibit energy-storage performances together with an excellent response to white light. Yun *et al.*³⁷³ fabricated a stretchable SiC gas sensor for NO₂ on an Ecoflex substrate, which could detect gas with a sensitivity of about 50 min after being charged. Moreover, flexible, self-powered devices have been reported, which are sensitive to human touch or motion.³⁷⁴ Kai *et al.*³⁷⁵ synthesized a stretchable conductive composite by mixing poly(3,4-ethylene dioxythiophene):*p*-toluene sulfonic acid (PEDOT:PTS) mixed with polyurethane (PU). The film fabricated by spin coating showed dynamic electrochromism in a hydrogel without supplying an external voltage, as given in Fig. 25. The stretchable PEDOT/PU film is suitable for a wearable display device; hence, it can be ultimate proof for the multifunctional features of wearable-to-self-powered with sensory smart window applications.

6. State-of-the-art EC devices: principal factors and composition

6.1. Principle factors

In an exemplary EC device (Fig. 1), cathode EC materials exhibit their inherent optical properties without an applied electrical potential, and the electrolyte ions (cations and anions) are uniformly dispersed in the device. When an electric potential is applied, the EC material captures the injected electrons, and thus the material gets reduced and then ionized, with a change in optical property to the “switching” mode. Meanwhile, the cations from the electrolyte move toward the opposite electrode and gather in the area of negatively ionized EC materials to balance the charge near the cathode. Also, the anions from the electrolyte enter the counter electrode to balance the charge and are stabilized by oxidization. Thus, fully reversible redox reactions occur with an appropriate reverse voltage. Therefore,

the performance of EC devices depends not only on the electrochemical reaction of heterogeneous electron transfer between the electrode and the EC film and/or ion storage film, but also on the homogeneous ion(s) transfer through the EC layer(s), the ion-transport layer, and the ion-storage layer. Thus, the EC performance can be optimized as follows: (i) by increasing the surface activity between electrodes and the EC film and/or ion storage film; (ii) increasing the electronic conductivity of electrodes and/or the ionic conductivity of electrolytes; and (iii) advancement to EC materials.

The EC material films are the key components defining the EC device performance, where the reversible optical modulation occurs under a suitable voltage. The dynamic windows consisting of WO_{3-x} can be electrochemically/electronically color/bleached with the insertion of a tiny electric potential (1–3 V) to dynamically control the inlet/exit of thermal loads and daylight *via* smart windows of buildings, vehicles, aircraft, *etc.*

EC devices can be (i) single EC and (ii) dual/multiple EC.

(i) Single EC devices are developed from one EC material coated as a thin layer. This type comprises materials with alternative one coloured state and one bleached state, which is useful in absorption/transmission-type devices. These devices may have fewer redox states compared to that composed of more than one EC active species.

(ii) Dual/multiple EC devices consist of two/more active EC materials triggered for multifunctional features in a single platform (Fig. 1). However, two/more active materials on the same platform can cause system complexity. Nevertheless, they have significant advantages in the possibility of multiple color changes at small applied potentials and multifunction. Dual-function devices designed with cathode and anode EC materials in one platform (Fig. 1) are considered to be more efficient, powerful, and therefore can save a tremendous amount of energy.

Based on their operation, EC devices can be transmissive type, which operates on the principle of optical transmission, and reflective type, regulating the optical reflectance. Further progress remains to be achieved in developing more efficient multifunctional EC devices with different capabilities in a



single platform. The key parameters used for evaluating the EC device performance are as follows:

- 6.1.1 Coloration efficiency
- 6.1.2 Switching time
- 6.1.3 Cycling life
- 6.1.4 Device stability and durability

6.1.1. Coloration efficiency. The change in optical modulation (optical density) occurring per unit injected charge is known as the coloration efficiency (CE), which has an impact on selecting the EC material and device performance. It can be expressed as follows:

$$CE = \Delta OD/Q = \log[T_{\text{bleached}}(\lambda)/T_{\text{colored}}(\lambda)]/Q \quad (7)$$

where T_{bleached} and T_{colored} are the transmittance in the bleached and colored states, respectively. The coupling of electrochromism and energy storage involves a high CE ($\text{cm}^2 \text{C}^{-1}$) and fast switching speed with the supply of low power for high optical modulation and charge density. Generally, an EC device having a high coloration efficiency will be more durable and switch faster.⁷⁹

6.1.2. Switching time. Switching time is the response time required to achieve switching efficiency between the transparent and different colored states. The switching speed depends on the (i) conductivity of the electrodes, (ii) ionic conductivity of the electrolyte, (iii) diffusion of ions in the thin films, (iv) magnitude of the applied potential, (v) film thickness, (vi) microstructure size, and (vii) surface morphology.

6.1.3. Cycling life. The cycle life refers to repeatable cycles between the colored state and bleached state under an applied voltage. Due to repeatable cycles, device failure may occur, such as solid-phase changes, and/or side reactions may occur, and thus the lifetime of thin layers decreases with changes in the electrode composition.

6.1.4. Device stability and durability. The stability towards redox reactions generally limits the utility of EC materials. For practical impact, EC smart windows should sustain steady switching after thousands (10^4 – 10^6) of repeatable cycles, which is a key obstacle for the perspective development of next-generation EC devices. In addition to long-term switching stability, device durability is also a key parameter. Moreover, the device should remain stable in an environment that is very cold (below -20°C) or very hot (above 80°C). Moreover, EC devices are generally sealed to avoid the adverse consequences of humidity, oxygen, air, and water in the external environment. Finally, the given appropriate voltage schemes and combination of electroactive materials candidates inside the EC devices are beneficial for promising EC properties.

6.2. Device configuration

Optical transparency and mechanical flexibility have become new developments in the next generation of flat panel electronics, enabling new technologies, including EC windows, (e)skin, wearable optoelectronics, and rolled-up displays.^{376,377} A traditional EC device is composed of a 4–5 layer configuration, and nanomaterials are widely used in each layer to achieve better performances and reliability. The recent

advances on EC materials (inorganic/organic) have been reviewed by Yang *et al.*,⁸⁴ Rai *et al.*,⁸⁷ Kim and Yang,¹⁰ and Eh *et al.*,¹⁹⁶ and the state-of-the-art approach of EC materials towards multifunctional EC smart-windows by Cai *et al.*⁸³ and recent progress in smart windows based on electro-, thermo-, mechano-, and photochromics is summarized by Ke *et al.*⁸ The more recent development in EC materials triggered in each layer to build the next-generation smart windows for “zero-energy buildings” is summarized in this review, as illustrated in Fig. 26.

- 6.2.1 Transparent conductive substrates
- 6.2.2 Ion conducting electrolyte
- 6.2.3 Ion storage/conductive oxide counter electrodes (anode)
- 6.2.4 EC material ($\text{WO}_{2.72}$) cathode

6.2.1. Transparent conductive substrates. For EC devices, the ideal electrode materials not only need to possess tremendously high conductivity, but also excellent transparency and high chemical/electrochemical corrosion resistance capabilities. The CVD, PVD, PLD, spray coating, spin coating, inkjet printing solution casting, and dip coating methods have been considered to be the best techniques for forming transparent thin films.³⁷⁸ The transparent conducting electrode materials used in EC devices will be summarized as:

(i) *Doped metal oxides.* For EC glasses, thin transparent conducting films consisting of doped metal oxides are generally employed as conductive electrode supporting substrates. Common examples include $\text{In}_2\text{O}_3:\text{Sn}$ (ITO), $\text{SnO}_2:\text{F}$ (FTO), $\text{SnO}_2:\text{Sb}$ (ATO), $\text{In}_2\text{O}_3:\text{Zn}$ (IZO), $\text{ZnO}:\text{Al}$ (AZO), $\text{ZnO}:\text{Si}$ (SZO), $\text{ZnO}:\text{Ga}$ (GZO), $\text{ZnO}:\text{B}$ (BZO), $\text{ZnO}:\text{F}$ (FZO), $\text{ZnO}:\text{Nb}$, $\text{ZnO}:\text{Sc}$, and $\text{SnO}_2:\text{Ta}$.^{5,379} All these co-doped oxides have wide bandgaps and are transparent in both the visible and NIR light spectrum. They can develop several bilayer architectures with a combined

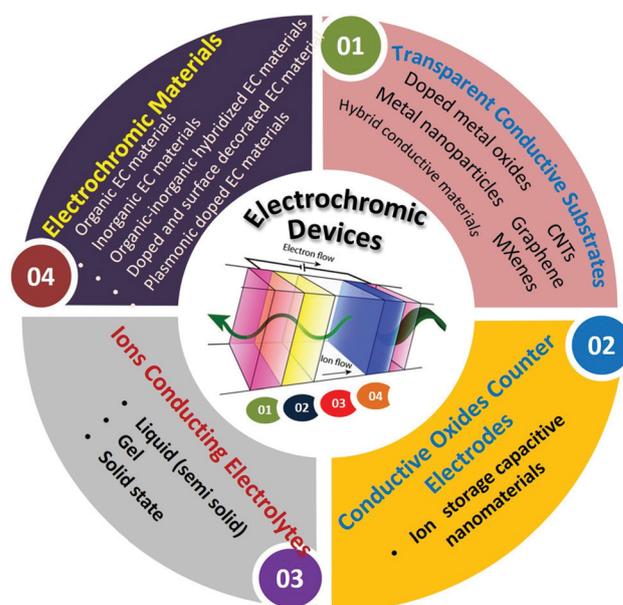


Fig. 26 Schematic illustration of the composition of EC devices.



resistivity of about $\sim 1 \times 10^{-4}$ – $10^{-3} \Omega \text{ cm}^{-1}$, expected thickness of up to $\sim 300 \text{ nm}$, and can be reduced to a certain adequate value with excellent durability. With the arrival of flat-panel display devices, ITO became the most commonly used transparent conductive oxide material for transparent electrodes;^{378,379} however, it is still not available on a large scale for EC smart windows. All these oxides are transparent in the solar spectrum.

The large-scale integration of traditional wearable electrochromic devices requires the development of flexible, stretchable, and deformable substrates.^{196,372,375,377,380} Generally, polymer, CNTs, graphene, MXenes, metal film, plastic, carbon fibre, silicone, hydrogels, and paper are used as supporting substrates due to their outstanding mechanical flexibility and physiochemical stability (resistivity, heat, good acid–base corrosion resistance, *etc.*).³⁸⁰ ITO is cost-expensive and brittle. Thus, conductive coatings on the surface of flexible EC substrates may be destroyed and cracked due to excessive stretching and bending processes at a lower strain of 2–3%. Concerns have also been raised over the limited indium source, stringent deposition conditions, and consequently high costs.³⁸¹ It is unstable in acidic/basic environments, which leads to a shortened device lifetime. Moreover, ITO films have a high refraction index, and thus can cause unwanted reflections.³⁸¹

Polyethylene terephthalate (PET) supporting substrate is a commonly used bendable electrode material due to its easy manufacture, low cost, excellent flexibility, super transparency (>85%), and constancy in weak acids–bases.^{382,383} Thermal evaporation and photolithography can be applied for deposition on PET substrates; however, its brittleness and processing at high-temperature (150 °C) limit its use in flexible EC devices. Therefore, it is difficult to distribute the functional electrode materials and sensing materials on PET substrates through direct chemical preparation. ITO- and FTO-coated glass substrates have been replaced by PET/polymers (polyimide), and because of their excellent conductivity, these materials improve the cycle stability and colour efficiency, and hence are suitable for use in EC devices (Fig. 1).³⁷ Polyimide (PI) sustains a high temperature (400 °C < 3%) without affecting wearability and transparency.³⁸⁰ Gas-phase and wet-chemical methods can be used to deposit materials on PI substrates, which demonstrates that the active materials have strong adhesion with the substrates, thereby improving their electrochemical properties. Moreover, ultra-thin silicon films (5–10 nm) and mica plates are also a good choice for EC material growth and device manufacturing due to their high thermal stability (>600 °C). Graphite, carbon paper and MXenes have attracted great attention because of their foldable, bendable, and rollable properties. Flexible substrate-based electrodes in EC devices have retroflected installation ability on existing windows, which significantly reduces costs compared to new installations. However, for large-area practical applications, the development of flexible EC devices at present is in its infancy. For example, in advanced flexible EC devices, the effect of UV light on polyester/PET foils can cause degradation and will affect the lifetime of EC windows. To avoid the brittleness of FTO and ITO

glass, some flexible transparent substrates such as graphene,^{384,385} carbon nanotubes (CNTs),³⁸⁶ metal nanowires,^{387,388} conducting polymers,^{302,303} and MXenes³⁸⁹ have been investigated as a substitute. However, the electrodes made up of these prevailing materials may suffer from limited transmittance in the bleached state.

(ii) *Metal nanostructures.* Metal grids and 1D noble metals nanoparticles such as Ag, Au, Cu, and Al have been tested as coatings for wearable substrates, which exhibit decent flexibility and excellent conductivity (Fig. 27).³⁹⁰ The metal-based coated layer should retain a low refractive index in the visible region and d-band contribution to the plasma wavelength oscillations in the UV range ($\sim 320 \text{ nm}$). The d–d electronic transitions provide minimum reflectance of a shorter wavelength in the visible region closer to the plasma wavelength.^{300,391}

Gold exhibits great potential because of its environmental stability and LSPR peaks that spectrally match with the solar emissions.^{293,393} Ag nanowire-based films are rapidly emerging due to their excellent conductivity, transparency, softness, and toughness, making them an auspicious candidate for the electrodes of next-generation wearable optoelectronics.⁸⁴ The refractive index of silver is 0.06 and its extinction coefficient of 3.59 at $\lambda = 550 \text{ nm}$ is the most ideal choice among the noble and non-noble metals in recent research. It has limited absorption in the visible region with a neutral color and has excellent optoelectronic properties.^{300,394,395} Cu multi-layers films also increased in popularity and became the preferred choice for thin-coating technology because of their tunable performance and reduction in overall material size and cost.³⁰⁰ Metals nanowire-based electrodes have good connectivity, flexibility, and anti-deformation ability. However, they have certain limitations, where the nanowires have weak wire-to-wire interaction and poor adhesion with the substrates, making it easy for them to detach from the surface, thus shortening the coating lifetime. It is worth noting that pure metal nanowires are easily oxidized in air and have poor stability due to long-term electrochemical cycles. Therefore, new materials with ultrasmall size and excellent morphologies should be exploited to construct flexible substrates with lightweight, high electrochemical stability, high transparency, high interface adhesion of conductive layers, and high thermal management properties.

To improve the conductivity stability of silver-based transparent conductive electrodes, Cai's group^{83,114} and Li's group²⁹⁹ proposed replacing ITO with silver grids as flexible transparent conductive substrates, as shown in Fig. 28. A high conducting silver grid/PEDOT:PSS hybridized composite film with outstanding stability was developed. An ultrathin layer of WO_3 nanoparticle coating was formed on an Ag-grid/PEDOT:PSS hybridized film, exhibiting the performance of flexible/foldable electrochromics. The hybrid-blend structure showed a large optical modulation of 81.9% at 633 nm, fast switching kinetics, and a high coloration efficiency of $124.5 \text{ cm}^2 \text{ C}^{-1}$. Moreover, the hybrid film showed outstanding electrochemical cycling stability (sustaining 79.1% of its initial



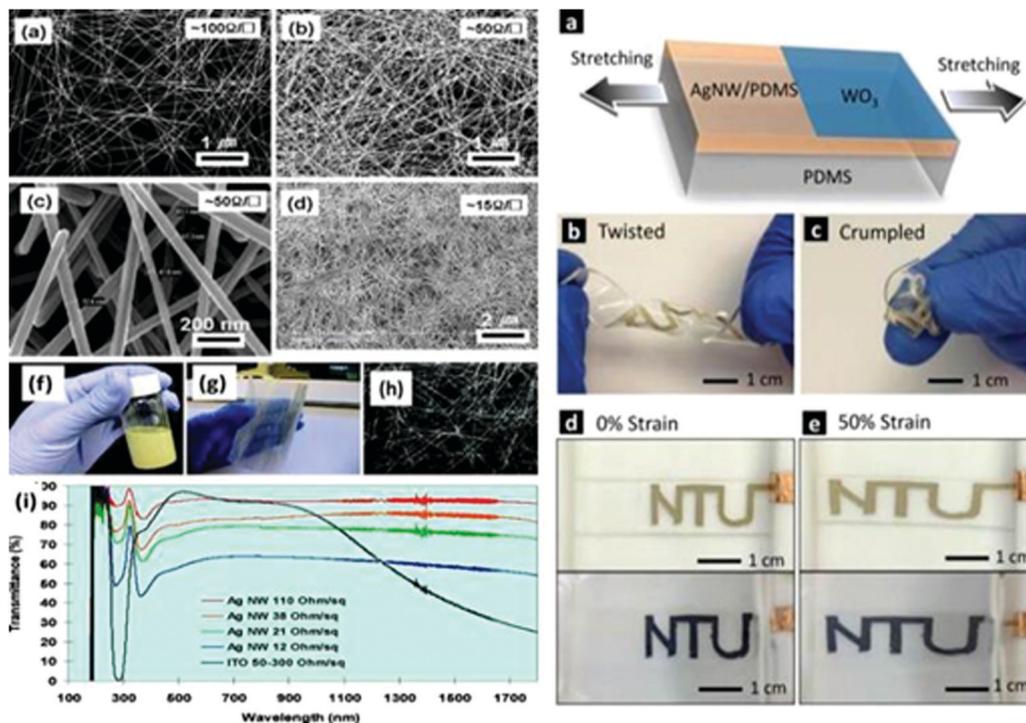


Fig. 27 (right) SEM images of Ag NW films produced from Ag NWs with a diameter of 40–100 nm and different densities lead to different sheet resistances of: (a) 100, (b and c) 50, and (d) 15 $\Omega \square^{-1}$. (f) 2.7 mg mL⁻¹ Ag NW ink in ethanol solvent. (h) SEM image of Ag NW coating shown in panel c. The sheet resistance is $\sim 50 \Omega \square^{-1}$. (i) Optical transmittance of transparent Ag NW electrodes measured with a UV-vis spectrometer, without including the substrates. Reproduced with permission.³⁸⁷ Copyright ©2010, ACS. (Left) (a) Schematic of the stretchable EC device. (b and c) Stretchable device is twisted and crumpled, showing excellent mechanical robustness. (d and e) Examples of the patterned device in the bleached and colored states at 0 and 50% strain, respectively. Reproduced with permission.³⁹² Copyright ©2014, ACS.

transmittance modulation after 1000 cycles) and remarkable mechanical flexibility (optical modulation decay of only 7.5% after 1200 compressive bending cycles). Consequently, a smart supercapacitor was developed as a conventional energy-storage device, which can simultaneously monitor the energy storage levels through rapid and reversible colour changes, even at high charge and discharge processes. The high-performance Ag-grid/PEDOT:PSS hybridized transparent films demonstrate a promising topology and a wide range of developing flexible electronics and photoelectronics. The high angle flexibility and bending nature of Ag nanowire-coated substrates maintain interconnectivity and can still illuminate light-emitting-diodes (LEDs) and bulbs after severe mechanical bending and deformation (Table 3).³⁷

(iii) *Carbon nanotubes (CNTs)*. Due to their distinctive 1D nanostructure and unique mechanical and electrical properties, CNTs have achieved potential in many types of electronic devices. In particular, their inherent high conductivity, excellent flexibility, and stability, and relatively easy and low-cost preparation make them a promising electrode candidate for next-generation smart electronics. When CNTs act as electrodes for EC devices, they can also be employed as promising sensing elements and stretchable/flexible electronics for the prospective development of soft and versatile ends. Moreover, because of their compatible functionalized structure, they can also

function as both an electrode candidate and EC smart material, which will further shorten the structure of EC devices and reduce their manufacturing costs.⁸⁴

(iv) *Graphene*. The thinnest 2D carbon material has wide application prospects in many research fields because of its excellent electronic, optical, mechanical, and thermal properties.³⁸¹ Graphene electrodes made of mechanically cracked graphene, CVD-grown graphene, or mass-produced graphene derivatives from bulk graphite have been used in a wide range of applications as an alternative to ITO transparent electrodes such as light emitting-diodes, touch screens, solar cells, field-effect transistors, batteries, supercapacitors, and sensors.³⁸¹

(v) *MXenes*. Recently 2D transition metal carbides, nitrides, and carbonitrides known as MXenes have achieved wide research interest due to their flexibility, superior mechanical strength, physical/chemical properties, and multiplex tremendous functionalities.³⁸⁹ Because of their excellent electrochemical stabilities, mechanical properties, high ion adsorption capacities and charge transfer, metallic conductivities, and unique topology, MXenes have demonstrated innumerable potential applications in structural composites, electromagnetic interference (EMI) shielding materials, lithium-ion batteries (LIBs), supercapacitors (SCs), electrocatalysts, electronics, and topological insulators.^{256,389,412} Interestingly, 2D



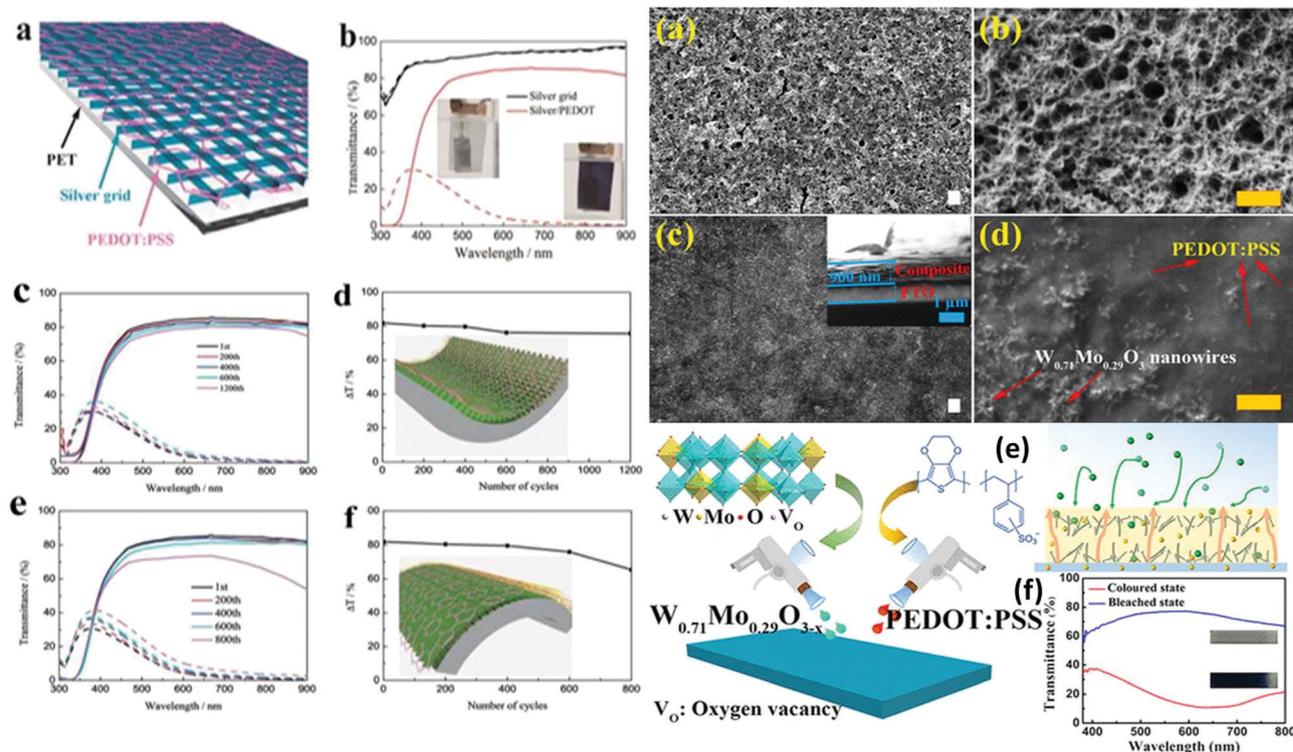


Fig. 28 Left (a) Schematic illustration of the structure of the silver grid/PEDOT:PSS hybrid film. (b) Transmittance spectra of WO_3 in the colored (-0.7 V) and bleached (0.1 V) state over the wavelength range of 300–900 nm. (c–f) Transmittance and optical modulation changes of WO_3 on the silver grid/PEDOT:PSS hybrid film in the bleached and colored state under repeated compressive bending (c and d) or tensile bending (e and f) with a curvature radius of 20 mm.¹¹⁴ Copyright©2016 Wiley-VCH. Right (a and b) FESEM images of the spray-LbL $\text{W}_{0.71}\text{Mo}_{0.29}\text{O}_3$ electrode and (c and d) $\text{W}_{0.71}\text{Mo}_{0.29}\text{O}_3$ /PEDOT:PSS electrodes. (e) Schematic illustration and (f) optical transmittance. Reproduced with permission.²⁹⁹ Copyright©2018 ACS.

Table 3 Transparent conductive substrates for EC devices

Electrode materials	Deposition method	Properties	Ref.
ITO thin film	Sputtering	$R_s = 10 \Omega \text{ sq}^{-1}$, $T \approx 90\%$	396
FTO thin film	Spray pyrolysis	$R = \sim 7 \times 10^{-4} \Omega \text{ cm}$; $T = \sim 90\%$	397
Ag NWs/PI	Wet process	$R = 25 \Omega \text{ sq}^{-1}$, $T = 86\%$	398
Ag NWs/PDMS	Spray process	$R = 20 \Omega \text{ sq}^{-1}$, $T = 75\%$	399
Ag NWs/PET	Lithographic filtration	—	392
Ag NWs/glass	Spin coating	$R = 10 \Omega \text{ sq}^{-1}$, $T = 60\%$	85
Ag NWs/PDMS	Spray coating	$R = 20 \Omega \text{ sq}^{-1}$, $T = 85\%$	400
Ag NWs/PMMA/glass	Spin coatings	$R = 2 \pm 0.1 \Omega \text{ sq}^{-1}$, $T = 67\%$	401
Ag NWs/PDMS	Spray coating	$R = 16 \Omega \text{ sq}^{-1}$, $T = 87\%$	402
CNT/PDA nanofibers	Spinning coating	$R = 100\text{--}1000 \text{ S cm}^{-1}$	403
CNT/PDMS	Paving method	$R = 0.1\text{--}1 \text{ k S cm}^{-1}$, $T = 80\%$	404
Graphene/PET	Layer formation	$R = 1 \sim 2 \text{ k}\Omega \text{ sq}^{-1}$, $T = 97\%$	405
Graphene/PET-ITO	Electrodeposition	$R = \text{—}$, $T = 80\%$	406
N-Graphene/glass	LPCVD	$R = 1.1 \text{ k}\Omega \text{ sq}^{-1}$, $T = 93\%$	407
RGO/Cu NWs/glass	Spin coating/spray process	$R = 34 \pm 2.6 \Omega \text{ sq}^{-1}$, $T = 80\%$	408
Ag NWs/PEDOT:PSS/PET	Spray process	$R = 10 \Omega \text{ sq}^{-1}$, $T = 86\%$	409
CNT/PANI/PET	Spray process	$R = 364 \Omega \text{ sq}^{-1}$, $T \approx 94\%$	410
RGO/Ag NWs/Ag grid/PET	Bar rod coatings	$R = 0.714 \Omega \text{ sq}^{-1}$, $T = 90.9\%$	411

MXene nanosheets and their derivatives can be produced *via* wet-chemical synthesis, which makes it easy to self-assemble hybrid-films and unique structures.^{256,412,413} Thus, the effective combination of the mechanical and functional properties of MXenes emerges as a promising opportunity to design nano-scale flexible, foldable, and wearable EC devices on a large scale with superb durability.

6.2.2. Ion conducting electrolytes. An electrolyte is the conducting ion reservoir between the EC materials and electrodes. It can exist as a liquid, gel, polymer, or in the solid state, either as solid thin films or a polymer layer.^{104,414,415} Electrolytes play a crucial role in the electrochemical reactions, providing a path for ions between the electrodes and avoiding the flow of electrons due to the cation-host interaction.^{104,416}



The ion transfer speed and efficiency have a great impact on the switching performance, including response speed, coloration efficiency, and cyclic stability. Granqvist well explained the corresponding types of glass-based rigid electrolytes and polymer-based flexible electrolytes.⁴¹⁷ Lee's team reviewed solid-state and polymer-based flexible electrolytes,^{418–420} whereas, Thakur *et al.*¹⁰⁴ reviewed the development of polymer electrolytes, and Li *et al.*³⁸⁰ also reviewed the solid-state gel-electrolytes for flexible/stretchable devices. Similarly, Yun *et al.*⁴²¹ also demonstrated non-volatile, Li ion-based gel electrolytes for WO₃-based flexible EC devices. In practical applications, solid-state electrolytes, for example polymers, ions, gels, and ion-conducting ceramic-based electrolytes are required to avoid certain limitations⁴¹⁹ caused by the use of liquid electrolytes, such as leaks, fire, and even deformation/breaking of the glass, especially in large-area EC window applications. Monovalent cationic-electrolytes are used in WO₃-based EC applications, in which tiny size ions can easily enter/exit the host-lattice. However, they have some disadvantages. The cyclic stability of the EC devices composed of amorphous WO₃ films is generally lowered in acidic electrolytes. This is because of the corrosion effect of H⁺ ions in acidic media. Due to the low electrode proficiency, the production of H₂ bubbles has also been observed on the surface of electrodes. The best choice is lithium-salt-based electrolytes, which can be used alternatively to achieve tremendous EC stability and prolonged switching kinetics. Polymer electrolytes have been employed for advanced electrochemical and EC devices.^{415,420,422–426} However, it is necessary to develop and select an electrolyte with an ideal performance for EC smart windows.

6.2.3. Ion storage/conductive oxide counter electrodes (anode). Complementary electrodes in EC devices play a key role in balancing the charge and ion storage properties. The electrode with ion-storage properties is generally referred to as the counter electrode, where the electrode with the EC layer in smart devices is known as the working electrode. When an appropriate electric current is applied, the EC materials on the working electrode are oxidized/reduced, and similarly, the equivalent reduction/oxidation of the electroactive materials occurs simultaneously on the counter electrode (Fig. 1). Next-generation smart windows generally require a counter electrode with (i) high transmittance, (ii) outstanding charge capacity, and (iii) low interface resistance, matching the working electrode. NiO, IrO₂, CeO₂, and V₂O₅ can be used as anodically chromic and/or charge storage materials, which are termed the counter electrode. The conventional EC windows show colour modulation in a complementary fashion to the primary material. Among them, NiO is the most popular, which shows a grey colour in the oxidized state. In NiO, the EC coloration is due to the surfaces and grain interfaces by the conversion of hydrated Ni oxide into hydroxide and then oxy-hydrate phases. The overall electrochromism occurs under the Bode-reaction scheme⁴²⁷ between Ni-hydroxide and oxy-hydroxide, as explained earlier.

Choi *et al.*⁴²⁸ deposited different size NiO particles on ITO glass through dry deposition, as shown in Fig. 29a–c. The

electrochemical results showed that the nanoporous NiO film had a high charge storage capacity (49.49 mC cm⁻²) and lower charge transfer resistance (0.49 Ω). The porous structure and large active surface area enabled better contact with the electrolyte and ion diffusion in the nanoscale NiO fabricated film. The film showed an excellent transmittance modulation (42%), with a stable switching speed and electrochemical cycles. Kim *et al.*⁴²⁹ fabricated N-doped carbon-based porous NiO films *via* a sol-gel process with the assistance of oleylamine and amide condensation. The film showed the coloration efficiency of 48.5 cm² C⁻¹, color/bleach speed of 3.2/2.7 s, and specific capacitance of 235.8 F g⁻¹ with the applied current density of 2 A g⁻¹ (Fig. 29d–h). Wang *et al.*⁴³⁰ reported the electrochemical deposition of H₃PW₁₂O₄₀ (PW₁₂) on a TiO₂ nanoparticle film coated on FTO glass, which functioned as an electron acceptor and redox shuttle for charge balancing of the TiO₂ nanoparticles as the counter electrode (Fig. 29i–l). The hybrid film achieved 68% optical modulation at 1.5 V to –2.0 V applied voltage, cyclic stability (2500 cycles), and good coloration contrast.

6.2.4. EC material cathode (WO_{2.72}). The EC layer is the main component in smart windows technology (Fig. 1), which consists of EC materials (Sections 2 and 3). The reversible electrochemical redoxation occurs with the insertion of ions through an applied electric potential and EC materials change their optical absorption properties dynamically. Among the EC materials, nonstoichiometric WO_{3–x} has received wide research interest due to its unique properties, as has been explained in detail in Sections 2–5.

7. Challenges and perspective outlooks

7.1. Challenges

Chromogenic materials have been used for dynamic switchable smart windows; however, the commercialization of EC windows in the market is still hindered because of major problems including high manufacturing costs, material implementation, durability, and reliability. The significant challenges faced in the synthesis of WO_{3–x}/WO₃ nanostructures on a large scale and efficient film preparation are as follows:

(i) Lack of standard synthetic methods for ultra-small size WO_{2.72}/WO₃ particles with a high surface area and EC electrode fabrication. It is very difficult to evaluate/compare the EC performance (coloration efficiency, switching speed, cycling life, cell stability, durability, *etc.*) reported in the literature.

(ii) Considering industrial applications, the synthesis methods are still far from market penetration due to the technical aspects and factors including quality, safety, cost, environmental hazard, mass production, and processability.

(iii) The phase-pure growth of WO_{2.72}-based materials on the surface of electrodes with a controllable morphology and dimensions can achieve excellent EC performances due to fast electron/ion transportation, but this has been largely ignored. In the meantime, the loading (density/%) of WO_{2.72}-based



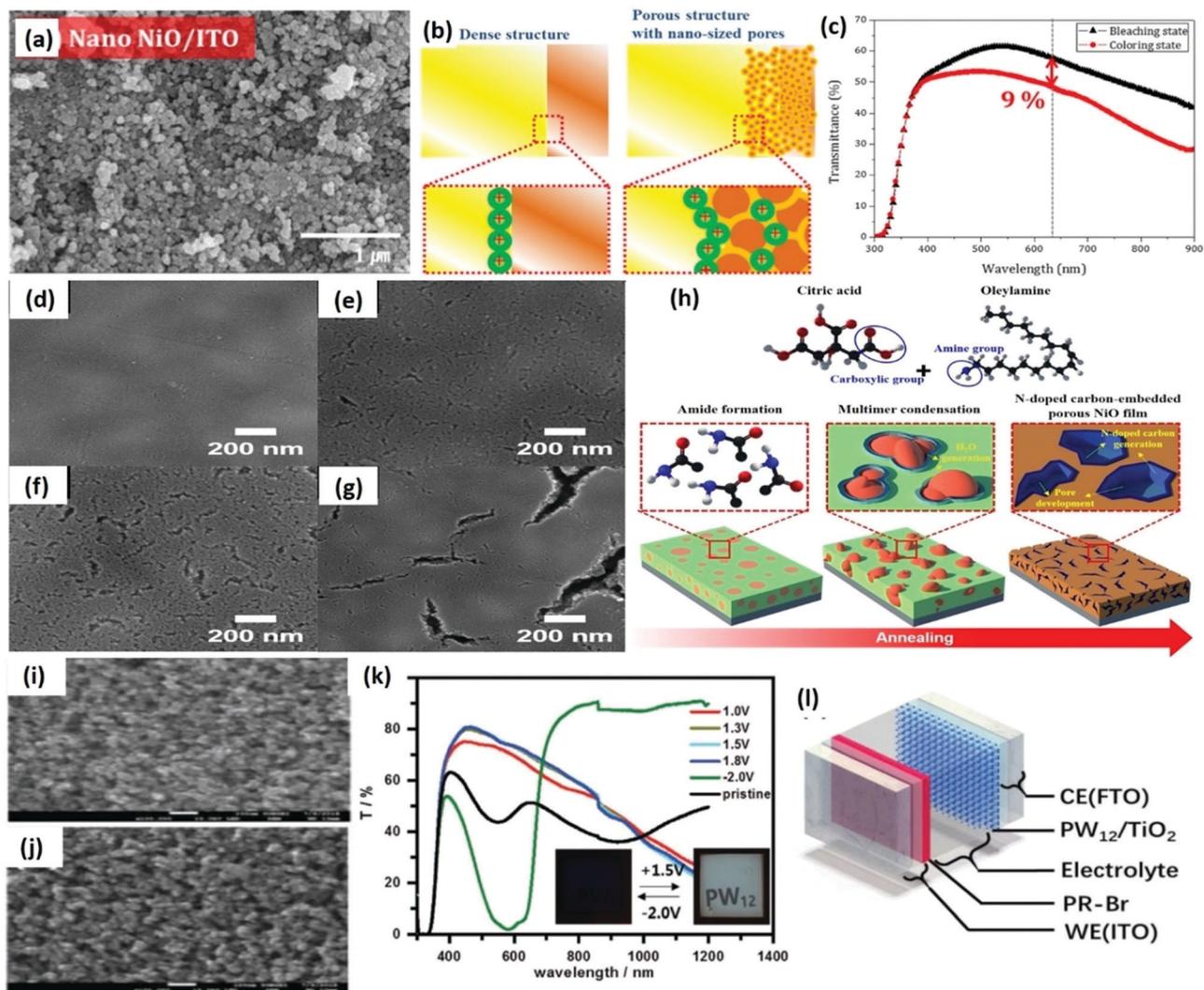


Fig. 29 SEM image of NiO film. (b) Schematic illustration of the ion storage and charge capacity on ITO film. (c) Transmittance of the film as a counter electrode. Reproduced with permission.⁴²⁸ Copyright@2018, Elsevier. (d–g) FESEM images of N-doped carbon-based NiO in the presence of different amounts of oleylamine, followed by annealing. Reproduced with permission.⁴²⁹ Copyright@2021, Elsevier. (i) SEM images of TiO₂ and (j) PW₁₂-TiO₂ film. (k) Transmittance of PW₁₂-TiO₂ at different voltages with photographs of the colored and bleach states. (l) Schematic of the various layers in the device. Reproduced with permission.⁴³⁰ Copyright@2019 WILEY-VCH.

materials, electrode area, EC performance parameters, and other conditions should be stipulated.

(iv) The active WO_{2.72} nanostructure, plasmonic-doped or plasmonic-nanocomposite powders are considered ideal for practical applications due to the presence of conductive agents. However, smart techniques are required to overcome the inherent disadvantages and phase purity of each component.

(v) For generating multifunctionality, the shape, size and tunable porous film coatings of WO_{3-x}-based plasmonic composites on wearable substrates acquire lightweight, high active surface area, and excellent conductivity. However, the charge/discharge capacity and expenses utilizing stretchable/deformable substrates are a key challenge for global researchers. Similarly, the optical modulation in white light and photothermal instability also need to be optimized.

(vi) When WO₃/WO_{3-x} tailored structures are deposited on a conductive substrate support, they can achieve relatively high electrochemical efficacy. However, at the device level, the cost may be quite high and the performance poor, with failure to reach the optical saturation threshold limit. Moreover, fast redox reactions/large electrochemical cycles result in EC dysfunction, and thus yellowing appears.

(vii) The mechanism through which oxygen vacancies/crystal defects and the electron/hole coupling in sub-stoichiometric WO_{3-x} affect EC and electrochemical properties is not well understood experimentally and/or theoretically.

(viii) Self-chargeable, foldable/flexible smart windows with unique structural designs, non-sacrificing optical properties, and cost-effective materials are a growing future demand and can be implemented first in unique wearable applications, as part of energy storage systems.



7.2. Outlook

7.2.1. Energy savings and multifunctional features. Due to their integration with multifunctional features such as energy storage, energy harvesting, and environmental sensors, the next-generation EC smart windows will be promising for novel diverse applications.^{33,34,40,66,103,196,372} Tungsten oxide-based coated films achieve dark blue color with an applied voltage and also become absorbing and store electric current. In self-powered configurations, solar energy is absorbed and converted to electrical energy. The stored energy can be used to power EC windows to initiate the redox reaction which dynamically changes the color of the windows, thus effectively regulating the solar loads in buildings.

Due to their similarities with energy storage devices, *e.g.*, supercapacitors and batteries, EC smart windows are known as transparent electrodes with reversible color states. Electrochromic supercapacitor devices (ESDs) with multicolor changes, energy harvesting and energy storage capacity, cyclic stability, and the realization of the quantitative monitoring of energy storage level of an ESD are very important to realize the practical application of the device.^{431,432} Conducting polymers have been studied as electrode materials for ESDs due to their rich color change and good energy storage performance.⁴³¹ Pseudocapacitive materials simultaneously enable a high rate, high energy density, and energy storage systems based on ion intercalation.^{433,434} They can be incorporated into electrodes to minimize the mass-transfer limitations due to thickness effects, and electrolyte starvation/low particle-to-particle electron transport. For this purpose, WO_{3-x} , NbO_x , MoO_2 , and TiO_2 with Mn, Ni, and Co oxides are appealing because of their properties related to storing electric charges and dynamic modulation of LSPR-absorption through capacitive charging/discharging.³⁴⁷ The 2D-layered materials known as MXenes have attracted much attention for applications in energy storage, optoelectronics, and photocatalysis.³⁷² MXenes demonstrate excellent flexibility, are easy to process, with an ultrafine thickness of high conductivity, and possess the highest electromagnetic interference shielding effect. Thus, self-assembled MXene- WO_{3-x} heterostructures should be employed as flexible and foldable energy storage EC films. The MXenes as a flexible transparent electrode and WO_{3-x} as the flexible EC layer are expected to achieve excellent performances in flexible EC windows on a large scale. The charge-storage mechanism should be better understood based on pseudocapacitive/battery-type materials linked with $\text{WO}_{2.72}$ to enrich the capacitive behavior for achieving self-powered multifunctionality. In addition, there is no clear quantitative relationship between the color change parameters and the energy storage level to achieve quantitative monitoring. This is because the constructed ESD is used to estimate the energy storage state through a color change. Under different charge-discharge voltages, the ESD displays different color.

Besides solar loads, human motions, and daily activities such as talking, walking, and running can produce a large amount of mechanical energy. Sensors that can harvest energy produced by human activities have been developed.

Self-powered Wi-Fi-connected smart windows integrated with environmental sensors can read room occupancy and detect changes in the weather, gases, light, temperature, and mechanical deformations. These all-in-one self-driven, shape-adaptive versatile smart windows can be self-chargeable with solar energy/heat and gather energy by human interaction/motions, and various other types of sustainable mechanical energy conversion resources would be promising improvements.

7.2.2. Reducing toxicity. LSPR nanocrystals anchored/doped with WO_{3-x} for the optical transmission of electrochromic-photothermal conversion of visible and NIR-light can effectively reduce the risk of microorganisms and are expected to be used as sterile smart windows. Au, Ag, Rh, Pd, and Pt LSPR have been widely used for cancer therapy. Thus, the broadband non-radiative plasmon decay of LSPR nanocrystals is assumed to be highly tunable as an electric field on the surface of WO_{3-x} . Surface-decorated WO_{3-x} plasmonic composites are very useful for reducing the thermal damage threshold due to electron-phonon and phonon-phonon coupling. MXenes are also promising antibacterial agents, with higher efficiency than graphene in diminishing bacterial cell viability.³⁷² Thus, Au-decorated/self-assembled Au-MXene- WO_{3-x} heterostructure-based films exhibit promising transparency, flexibility, EC-photothermal sterility, and effective modulation of visible and NIR light. The photothermal transformation will be attained in the colored state, which is ascribed to the coupling of traditional visible-band optical switching with the NIR-LSPR extinction, and thus can act as high-tech sterile EC smart windows. These multifunctional sterile EC smart windows will be particularly useful in buildings, hospitals, high-latitude zones, and aircraft.

7.2.3. Coloration enhancement/poly-electrochromism. Due to electron/ion intercalation, tungsten oxide films achieve dynamic controllable blue color, which has a great influence on light absorptivity. The color preferences are closely associated with the cultural background, where the deep blue color is very suitable in China and Pakistan. The color development choices may vary from country to country. Doping can pave a path to change the colors by shifting the adsorption edges. Thus, doping and co-/multi-doping must be tested at the atomic level through experiments and simulation studies, together with further structural design at the nano-level. Hybridization with either inorganic/organic candidates is an efficient way to modulate colors/multicolor; however, the complicated fabrication strategies should be exploited further with more creative ideas.

In recent years, DMD (dielectric/metal/dielectric) blends have attracted increasing attention, in which the WO_3 membrane can play a dual role of reflecting light energy in the bleaching state and absorbing in the colored state. In addition, Ag is an excellent heat-regulating material, which can also replace expensive substrates, *e.g.*, FTO and ITO. The presence of extreme transmissive states in $\text{WO}_{3-x}/\text{Ag}/\text{WO}_{3-x}$ allows the combination of numerous EC materials (hybrids) with controlled redox reactions to produce a variety of colors and bleach states with color coordination. In addition, the coloration efficiency and color coordination for $\text{WO}_{2.72}$ electrochromism have not been investigated systematically, which



complicates further advanced research. In the DMD architecture, the Cu and/or Al metals sandwiched between EC material layers are reliable and thermally reflective. Moreover, they can be promising in terms of cost, durability, and ease of processing. Researchers found that WO_3 , TiO_2 , SnO_2 , ZnO , CuS , ZnS , AlN , MoO_2 , and Nb_2O_5 can form functional multilayer structures that can functionalize conductive metals, for multicoloration in EC smart windows.

7.2.4. Reversibility, durability, and inadequate switching. It is worth mentioning that $\text{WO}_{2.72}$ films suffer from degradation due to prolonged switching (ion-intercalation/deintercalation), thus affecting the optical modulation and cyclic stability, and consequently the device performance. Continuous current-driven ion de-trapping and self-healing are a successful way to eliminate the ion-trapping resulting in degradation and reduction in EC properties. Therefore, the film can be refreshed and attain its initial state, which is a successful effort to design super EC substrates for smart windows and other powered optoelectronics. Self-healing EC materials (*e.g.*, viologens/polymers) can perform *in situ* redox reactions with excessive Li^+ ions to reduce resistance transfer and avoid interface obstacles. The reversible Diels–Alder cross-linking network⁴³⁵ structure can heal the cracks and improve the reliability of the WO_{3-x} EC layer. The formation of an inert oxide shell separating WO_{3-x} from oxygen also induces a characteristic effect, but the controllable formation of the core–shell nanoarchitecture is not very easy. Other essential variables are the $\text{WO}_{2.72}$ response time under various film thicknesses, material loading per unit area, and long-term environmental stability. For additional protection and to avoid unnecessary physical/chemical reactions or environmental strain (damage, ultraviolet, heat, water vapor, and/or oxygen), smart window devices can be well sealed by covering curtains, as shown in Fig. 1, which helps improve their stability, performance, and reliability.

7.2.5. Dual/multifunction. Self-powered EC windows can offer numerous ways to reconfigure sustainable energy resources in smart windows, thus reducing energy consumption by minimizing the heating loads, cooling loads, lightings, and also protecting the privacy of users. Smart windows can also store energy during the electrochemical process through a rational circuit in the same electrolyte. Consequently, the stored energy can be utilized to derive other functional devices. Tremendous progress has been made in smart electronics and dual/multifunctional windows for practical applications, in which the WO_3 and NiO ⁴³⁶-based EC supercapacitors have been widely demonstrated.⁴³⁷ Dual-function EC supercapacitors serve as an energy storage device and electrical power source simultaneously, displaying real-time capacity in the color state.^{431,432} The unique characteristics achieved include coloration/bleaching and charging/discharging, large transmittance contrast ($\sim 91\%$), high coloration efficiency ($\sim 61.9 \text{ cm}^2 \text{ C}^{-1}$), high areal capacitance ($\sim 13.6 \text{ mF cm}^{-2}$), and good charging/discharging cyclic stability.⁴³⁷ A low doping amount of less than 10 mol% can significantly enhance the EC performance.⁴³⁶

Moreover, smart windows with unique novel multifunctional features extend their utility. Metal oxides become opaque

upon ion extraction such as oxides of nickel and iridium or PEDOT/PSS materials. Among them, Ni is more functional, relatively common, and less expensive, which make it preferable for large-scale production. A device that can combine two types of EC functional materials, *e.g.*, WO_{3-x} and NiO , will be more advantageous, where the charges transfer from Ni to W oxide. In the opposite direction, the EC materials will be bleached (Fig. 1). It should be stated that the EC properties of NiO are different from that of WO_3 in the case of Li^+ ion-based electrolytes. The reason for this is that NiO involves surface sites for the exchange of both cations and anions.⁴³⁸ However, these novel multifunctional EC smart windows should be designed to enrich various EC materials coatings to convert solar loads into useful energy simultaneously,³⁴ or to allow the transmission of visible light but reflect NIR light. Hybrid functional compatible electrochromics in a single platform are significant alternatives in balancing the performance limitations of one type of EC material by supplementing with the required activity of the competent material. Thus, enriching versatility in a smart way and minimizing the overall device size with superb service life will be fundamental approaches in the future.

7.2.6. Advancements in $\text{WO}_{2.72}$ nanoparticles and film formation strategies. $\text{W}_{18}\text{O}_{49}$ nanoparticles should be considered for commercialization by controlling their dimensions, size, dispersibility, phase purity, synthesis routes, and also coating strategies, such as annealing the coated film, surface porosity and adhesion of films with substrates, nature of the conductive substrates, nature of electrolytes, *etc.*, and quantity/quality regulation of all components. Plasmonic metal-doped WO_{3-x} and its composites in the desired shape, size, and tunable porous film coatings with a high active surface area, excellent conductivity, and lightweight on stretchable/deformable substrates should be the focus for scaling up. PED, PLD, and HiPIMS can be effective deposition methods for the coating of $\text{WO}_{2.72}$ -based EC materials.^{37,86,114,309,310,392} The wet-deposition techniques require the preparation of suitable inks from nanostructured EC materials, and thus the ink preparation can be the main components in assessing the coating strategies. Inkjet printing is a cost-effective and highly attractive industrial technology that can be used to manufacture EC materials on a variety of substrates. Moreover, it allows the coating of EC materials in a precise location with a controllable thickness.

The traditional vacuum evaporation, magnetron sputtering, and spin-coating methods are not suitable to achieve large-area coatings, controllable thickness, and low cost. Recently, our team has focused on different ways to produce ultra-thin films by rolling, spraying, casting/screen printing, and spin coating on various substrates, which shall remarkably reduce costs and readily achieve high-quality EC/TC smart responsive coatings to fulfill the huge market demands. However, the performance optimization and the commercial production still need more effort, as the researchers require cost-effectiveness, manufacturability, and market demand user willingness and policies. Thus, suitable and sustainable solutions to the problems



addressed in this review regarding EC smart windows need to be found in the future.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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