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# Flexible electrochromic devices based on tungsten oxide and Prussian blue nanoparticles for automobile applications†

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Smart windows, which control the amount of light entering buildings, houses, and automobiles, are promising in terms of energy conservation and their low environmental impact. Particularly, a next-generation smart window for automobiles will require high optical modulation, along with flexibility to adapt to various intelligent designs. We have previously fabricated electrochromic devices (ECDs) by wet coating glass substrates with nanoparticles (NPs), such as water-dispersive ink containing tungsten oxide (WO<sub>3</sub>), and Prussian blue (PB), and have evaluated and confirmed the various electrochromic (EC) properties, such as optical modulation, cyclic durability, and colouration efficiency, of the ECDs. However, glass substrates are heavy and difficult to adapt by deformation to meet the demand of next-generation automobiles. In this study, we aim to prepare complementary ECDs by wet coating WO<sub>3</sub> and PB thin films on indium tin oxide (ITO)-coated flexible polyethylene terephthalate (PET) substrates. Chromaticity and haze of ECDs were investigated in detail as evaluation indexes to verify specifications for practical use in automotive applications. Repeated switching, bending, and twisting did not degrade the ECD properties, thereby demonstrating its durability and mechanical robustness. These excellent electrochromic properties of the flexible ECDs suggest that they are promising materials for application in next-generation smart windows for automobiles.

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## Introduction

A “smart” automobile window is one that has flexibility to autonomously change its optical properties to admit or block solar energy in response to seasonal changes.<sup>1–3</sup> Recently, suspended particle devices (SPDs) have been developed for commercial automobile window applications.<sup>4,5</sup> However, due to the disadvantages of SPDs, such as poor thermal and electrical stabilities and poor memory effect, they cannot meet the demands of future automobile development. In contrast, electrochromic (EC) materials have high thermal and electrochemical stabilities as well as good memory properties, and are thus considered suitable candidates for automobile window applications. Upon application of a certain voltage, EC materials can switch their optical properties persistently and reversibly by balancing charges *via* simultaneous intercalation/deintercalation of cations (*e.g.*, H<sup>+</sup>, Li<sup>+</sup>, *etc.*) and electrons.<sup>6–8</sup> EC materials have been extensively investigated due to their potential for widespread application in information displays, automotive rear-view mirrors, and smart windows.<sup>9–11</sup>

A typical EC device comprises five superimposed layers: a transparent conductive electrode, an ion storage layer, an electrolyte layer, a EC layer, and a transparent conductive electrode.<sup>12,13</sup> The ion storage layer prevents ion accumulation on the electrode to maintain the neutrality of the electrolyte layer. The electrolyte layer is ion-conductive and provides a transport channel for the shuttling of ions between the EC and ion storage layers.

EC materials are mainly categorized as inorganic and organic materials. Inorganic materials include transition metal oxides (*e.g.*, tungsten oxide (WO<sub>3</sub>)<sup>14</sup>) and inorganic complexes (*e.g.*, Prussian blue (PB)<sup>15–17</sup>), while organic materials include  $\pi$ -conjugated organic molecules (*e.g.*, viologen<sup>18</sup>) and conductive polymers (*e.g.*, polyimide<sup>19</sup>). In particular, WO<sub>3</sub> and PB are representative EC materials that exhibit large optical modulation and high colouration efficiency in the visible and near-infrared regions. Interestingly, Wang *et al.*<sup>20</sup> reported not only a bi-functional device, but also a self-rechargeable transparent battery, using PB having EC characteristics.

EC layers for application in commercial smart windows can be prepared by thermal evaporation,<sup>21</sup> electrodeposition,<sup>22</sup> and sputtering.<sup>23</sup> In recent years, hydrothermal methods,<sup>24–26</sup> sol-gel methods,<sup>27</sup> along with electrodeposition,<sup>28</sup> printing,<sup>29</sup> and coating<sup>30</sup> techniques, have been investigated to reduce process cost for high productivity. These coating and printing techniques can be used to rapidly fabricate high-quality EC thin films on large substrates. Thus, these techniques show promise for the development of next-generation flexible ECDs.<sup>30,31</sup>

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However, this development relies on extensive basic research, because several issues must be resolved to enable mass production, ensure process safety (*e.g.*, minimise the use of substances that can harm the environment), and allow for easier handling.

We have previously developed a complementary coated ECD using  $\text{WO}_3$  and PB nanoparticles (NPs).<sup>14,32</sup> The  $\text{WO}_3$  and PB thin films were coated with an ink containing each of these NPs dispersed in water, thus ensuring processing safety. Slit coating was also evaluated for the fabrication of large-area thin films on a G2 (370 mm  $\times$  470 mm)-sized indium tin oxide (ITO)-coated glass substrate.<sup>33</sup> High-quality EC thin films can be fabricated in a short period at a low cost using wet coating techniques, such as slit-coating<sup>34</sup> and roll-to-roll<sup>35</sup> methods. However, suitability of the application of these  $\text{WO}_3$  and PB inks on flexible substrates has not yet been evaluated.

In this study, we aimed to fabricate a flexible ECD by coating  $\text{WO}_3$  and PB on ITO-coated polyethylene terephthalate (PET) as a substrate material. The ECD performance was evaluated in terms of EC properties, including optical density change, colouration efficiency, and response time, of the assembled device. Particularly, it is preferable to use glazing in automobile windows because it affords high transmittance in a transparent state with no excessive light scattering;<sup>36,37</sup> therefore, we investigated the haze and chromaticity of the coloured and transparent states of the ECD fabricated on a PET substrate in detail. ECDs fabricated on conventional glass and flexible substrates were also compared. The primary aim of the research described herein and future research is the development of a coated film for application in automobile smart windows.

## Experimental

### Water-dispersed $\text{WO}_3$ and PB NP inks

Water-dispersed PB and  $\text{WO}_3$  NP inks were prepared by following methods reported in our previous papers.<sup>14–16,32</sup> Water-dispersed PB NP ink was prepared by mixing  $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$  with  $\text{Na}_4[\text{Fe}(\text{CN})_6]$ .

$10\text{H}_2\text{O}$ , followed by washing *via* decantation. The dispersibility of the precipitate in water was improved by adding  $\text{Na}_4[\text{Fe}(\text{CN})_6] \cdot 10\text{H}_2\text{O}$ , and the suspension was stirred at room temperature for 1 week. Polyvinyl alcohol (PVA; 10 wt%) was added for increased viscosity and adhesion of the ink to ensure that a uniform PB film was formed.

$\text{WO}_3$  NPs (diameter  $\approx$  10 nm; Toshiba Materials Co.) were dispersed in purified water to a solid content of 25 wt%, followed by stirring at 1000 rpm for 48 h at 20 °C. PVA (1 wt%) was added for increased viscosity and adhesion of the ink to ensure that a uniform  $\text{WO}_3$  film was formed. The water-dispersed  $\text{WO}_3$  and PB NP inks were both filtered through a 7  $\mu\text{m}$  syringe filter (Kiriya Glass Co.) before use. The properties of the inks are listed in Table S1.†

### Surface treatment of substrates

ITO-coated glass and PET were both used as substrate materials (area = 25 cm<sup>2</sup>). Both substrates received surface treatments to improve ink coating efficiency and thin film adhesion. After setting the moving distance of the conveyor to 50 mm, the surfaces were repeatedly irradiated with ultraviolet (UV) light ( $\lambda = 365$  nm) for 25 min at a conveyor speed of 10 mm min<sup>-1</sup>, with the distance between the substrate and UV light emitting ceramic lamp being 10 mm (HLDL-375X10U6-SP, CCS Inc.; Fig. 1). The UV dose was adjusted between 0 and 2805 mW s cm<sup>-2</sup>. The temperature was controlled using a cooling fan (<80 °C), and temperature labels were used to avoid heating of the flexible substrate surface during UV treatment. The contact angles for water and the PB and  $\text{WO}_3$  NP inks dropped onto the ITO-coated glass and flexible substrate surfaces were measured after 60 s using a contact angle meter (DMS-400, Kyowa Interface Science Co., Ltd).

### Electrochromic thin films prepared using water-dispersible NP inks

$\text{WO}_3$  and PB thin films were deposited on surface-treated ITO-coated glass and flexible PET substrates by spin-coating

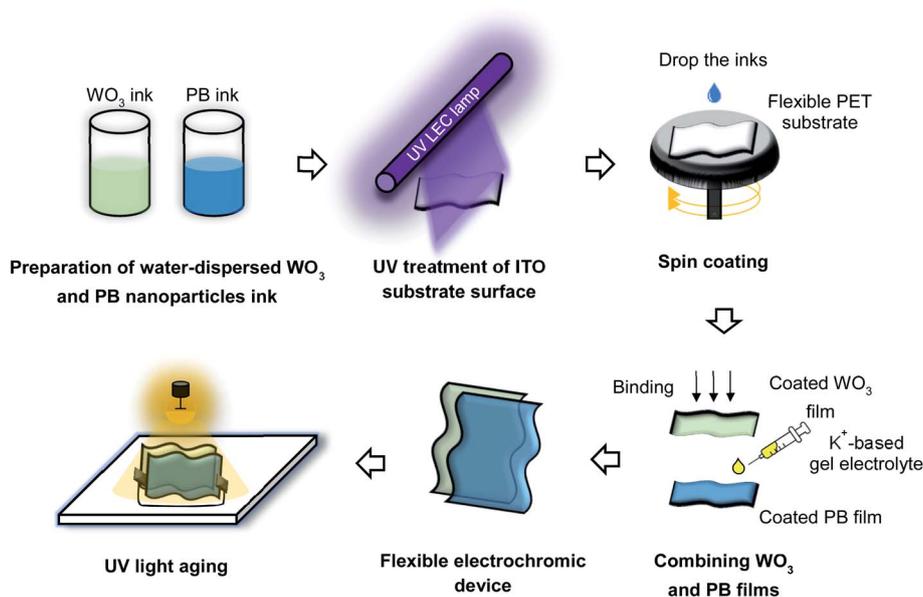


Fig. 1 Schematic of ECD preparation.



(POLOS Spin150i, SPS Europe, Inc.). The spin parameters are listed in Table S2.† The speed (rpm) of the spin-coating process was controlled to yield a film thickness of  $\sim 1$   $\mu\text{m}$ . The film thickness ( $d$ ) was determined as follows:<sup>38</sup>

$$d = ml(A \times \rho), \quad (1)$$

where  $m$  is the weight of the coated  $\text{WO}_3$  or PB NP-water-dispersible ink,  $A$  is the area of the substrate, and  $\rho$  is the density of the ink. The weights of the ITO-coated substrates and  $\text{WO}_3$  and PB inks are listed in Table S3.†

### Fabrication of electrochromic devices (ECDs)

The gel electrolyte was prepared by mixing potassium bis(trifluoromethanesulfonyl)imide (KTFSI, 1.5 mol  $\text{kg}^{-1}$ ; FUJIFILM Wako Pure Chemicals Co.) and polycarbonate (PC, 99%; Kanto Chemicals Co., Inc.). The electrolyte viscosity was adjusted by adding 30 wt% polymethyl methacrylate (PMMA; FUJIFILM Wako Pure Chemicals Co.). The electrolyte was applied to the PB thin film, and a UV sealing material was applied to the four edges of the PB thin film to prevent electrolyte leakage. Next,  $\text{WO}_3$  thin film was bonded to the PB thin film with the electrolyte and UV seal material applied under vacuum. The effective area of the devices was 16  $\text{cm}^2$ . The ECDs produced in this study are referred to as either a glass-based ECD or a PET-based ECD.

A light-ageing treatment was performed to improve the performance of the ECDs, by irradiating the ECDs with a xenon lamp (Q-SUN XE-1 Xenon Test Chamber, Q-Lab Co.) operated at a power density of 80  $\text{mW cm}^{-2}$  for 60 min, placed 50 mm from the ECDs (Fig. 1). The electrodes at both ends of the prepared ECDs were short-circuited before light-ageing. Upon light-ageing, the appearance of the ECDs was switched from blue to transparent.

### ECD characterisation

The ECDs were evaluated in accordance with protocols used in our previous studies.<sup>14,32</sup> The chromaticity was measured using a UV-visible (UV-vis) spectrophotometer (SD 3000, Nippon Denshoku Industries Co. Ltd), while the haze was determined using a haze meter (NDH 7000, Nippon Denshoku Industries Co. Ltd). The chromaticity coordinates ( $a^*$  and  $b^*$ ) and luminance ( $L^*$ ) data were calculated from the transmittance spectra of PET-based and glass-based ECDs in their coloured and bleached states. The luminance ( $L^*$ ) value characterizes the brightness of the films and varies between 0 (black) and 100 (white). The colour of the films is defined by their  $a^*$  and  $b^*$  coordinates: positive and negative  $a^*$  values indicate that the colour tends toward red and green, respectively, while positive and negative  $b^*$  values indicate that the colour tends toward yellow and blue, respectively. The electrochemical properties were evaluated using an electrochemical analyser (6115D, ALS/HCH), in combination with a UV-visible-near-infrared (UV-vis-NIR) light source (DH-2000, Ocean Optics Inc.). Cyclic voltammetry (CV) was performed to estimate the peak potentials and peak currents of the reversible redox reactions at the anode and

cathode. Multiple potential step (MPS) measurements, transmission spectra, and time strip charts were obtained based on estimated response times. Chronocoulometry (CC) was conducted to estimate the transferred charge density ( $Q$ ), with respect to the transformation of ECDs from the transparent to the coloured states.

## Results and discussion

### Surface treatment of ITO substrates

The hydrophilization of a substrate surface is commonly achieved using the vacuum plasma method.<sup>39</sup> However, this approach can lead to plasma damage in organic substrates. Therefore, in this study, the surfaces were treated with UV light. UV-based hydrophilic treatments help remove contaminants from the ITO surface by breaking the organic molecular bonds and adding polar oxygen atoms. These modifications increase the surface energy of ITO.<sup>40–42</sup> The contact angles for water and the PB and  $\text{WO}_3$  inks dropped on the surface-treated substrates were evaluated after treatment at various UV irradiation intensities (Fig. 2). The contact angles of water,  $\text{WO}_3$ , and PB were found to decrease from  $\sim 90^\circ$  to  $32^\circ$ ,  $48^\circ$ , and  $47^\circ$ , respectively, with increasing UV dose; the optimal effect without any damage to the flexible substrate was achieved at a UV dose of 2805  $\text{mW s cm}^{-2}$ . The UV surface treatment reduced the content of non-polar groups, while increasing the content of polar components, and the formation of various photo oxides on the ITO surface led to increased hydrophilicity.<sup>40–42</sup>

### Effect of light-ageing on ECD properties

Light-ageing of the ECDs with short-circuited electrodes afforded a transparent appearance due to presence of oxidised  $\text{WO}_3$

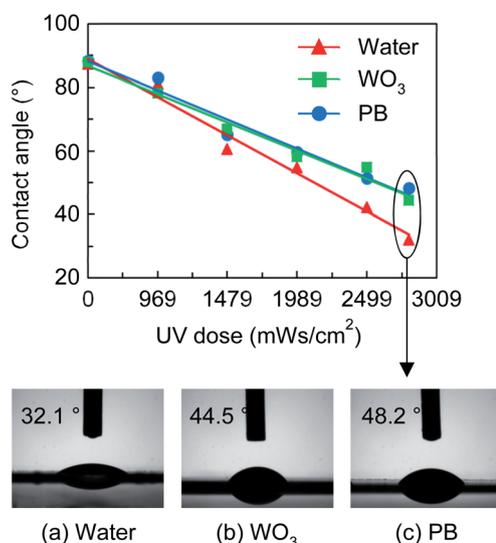
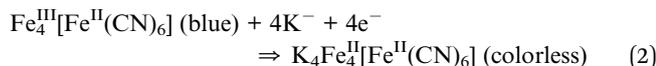


Fig. 2 Contact angles for water and the PB and  $\text{WO}_3$  inks dropped on the ITO/PET substrate after UV treatment at various UV irradiation intensities. The photos show the contact angle of (a) water, (b)  $\text{WO}_3$ , and (c) PB on the surface of ITO-coated PET substrate treated with a UV dose of 2805  $\text{mW s cm}^{-2}$ .



and reduced PB. UV irradiation facilitated the injection of  $K^+$  into the electronic states of PB for the reduction of  $Fe^{III}\text{-N}$  to  $Fe^{II}\text{-N}$  as follows:<sup>43,44</sup>

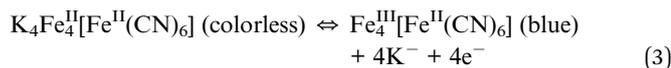


The PB thin film in the sealed ECD was protected from oxidative degradation, thereby minimising its participation in the oxidation of PB.<sup>44,45</sup>

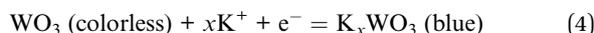
### Electrochemical and EC properties

The CV curve recorded at an applied potential ranging from  $-1.2$  to  $+1.0$  V at a sweep rate of  $5 \text{ mV s}^{-1}$  is shown in Fig. 3(a). The PET-based ECD exhibited a significantly smaller CV area than the glass-based ECDs, probably because the sheet resistance of the ITO/PET substrate ( $\sim 50 \text{ } \Omega \text{ sq}^{-1}$ ) was about four times higher than that of the ITO/glass substrate ( $\sim 12 \text{ } \Omega \text{ sq}^{-1}$ ). The charges of the glass and PET-based ECDs were  $0.44$  and  $0.23$  C, respectively, when a potential of  $-1.2$  and  $+1.0$  V was applied for  $60$  s (Fig. 3(b)). Thus, the glass-based ECD with a low electrical resistance had double the charge for oxidation and reduction reactions compared to the PET-based ECD. However, the PET-based ECD had an estimated charge balance ( $C_{\text{transparent}}/C_{\text{coloured}}$ ) of over 98%, indicating that reversible  $K^+$  charge/discharge processes occurred smoothly in both ECDs, as described in eqn (3) and (4):<sup>46,47</sup>

Anodic reaction:



Cathodic reaction:



The changes in optical transmittance of the coloured and transparent ECDs were examined using UV-vis spectroscopy during CC analysis, and the corresponding spectra are shown in Fig. 4(a) and (b). The glass and PET-based ECDs exhibited optical modulations of 80% and 79%, respectively, at a wavelength of  $633$  nm. Both the ECDs exhibited excellent optical properties. MPS measurement were conducted to determine the switching responses of the films. The changes in optical transmittance with time at a wavelength of  $700$  nm upon

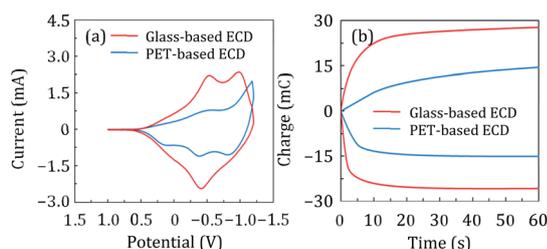


Fig. 3 (a) CV and (b) CC profiles of the glass-based (red) and PET-based (blue) ECDs acquired at a potential scan rate of  $5 \text{ mV s}^{-1}$ .

applying constant voltages of  $+1.0$ ,  $-1.2$  and  $+1.0$  V, as  $V_{\text{initial}}$ ,  $V_{\text{step1}}$ , and  $V_{\text{step2}}$ , for  $30$ ,  $30$ , and  $60$  s, respectively, are shown in Fig. 4(c) and (d). The switching times of the ECDs between the coloured ( $t_{\text{coloured}}$ ) and transparent ( $t_{\text{transparent}}$ ) states was evaluated using the time in which 90% change in transmittance was achieved. The PET-based ECD exhibited  $t_{\text{coloured}}$  and  $t_{\text{transparent}}$  values of  $16.79$  and  $19.03$  s, respectively, which were higher than those of the glass-based ECD. However, there was no significant difference in the response time between the two types of ECDs, as obtained in the CV and CC analyses (Fig. 3), despite the lower current values. This result shows promise for lower current drive in future applications, which could reduce the power consumption of a device. A shorter switching time is also expected if a substrate with a lower sheet resistance than ITO-coated substrates is used.

Fig. 5 shows the chromaticity and haze of the ECDs, aspects that are important for meeting the regulations governing window materials for automobiles (the details of the haze and chromaticity values are described in Table S4†). In the CIELAB colour space designed to approximate the human vision, the colour coordinate values ( $L^*$ ,  $a^*$ , and  $b^*$ ) of the ITO-coated PET- and glass-based ECDs were determined in transparent and coloured states (Fig. 5(a) and (b)). When  $-1.2$  V was applied, the  $a^*$  and  $b^*$  values of both types of ECDs suddenly dropped below  $-15$ , corresponding to blue colour. Conversely, upon the application of  $+1.0$  V, both types of ECDs changed from blue colour to transparent ( $L^* > 85$ ). In addition, the haze values are low ( $<5\%$ ) in the transparent states of PET-based and glass-based ECDs. For application in windows, visual transmittance must be high, desirably  $>70\%$ , and haze must be low, desirably  $<2\%$ .<sup>48</sup> The PET-based ECD appears remarkably clean with haze  $<3\%$  in the transparent state, which is adequate for use as ECDs. Moreover, the ECDs maintained their coloured and transparent states even when disconnected from the power source, demonstrating it as a promising energy-efficient device.

The current and transmittance at  $633$  nm were evaluated as a function of the number of continuous colouration/

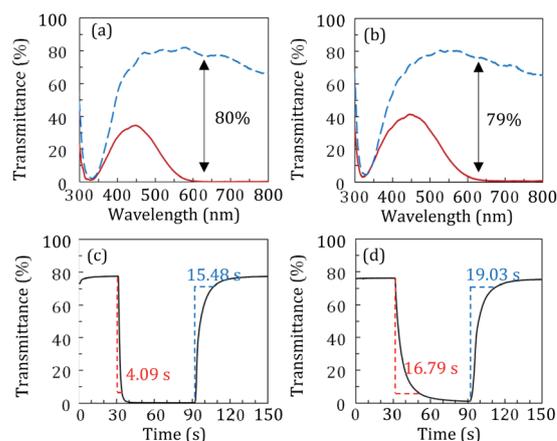


Fig. 4 Transmittance spectra of the transparent (blue lines) and coloured (red lines) states of (a) PET-based and (b) glass-based ECDs. Optical switching speed between the transparent (blue lines) and coloured (red lines) states of (c) PET-based and (d) glass-based ECDs.



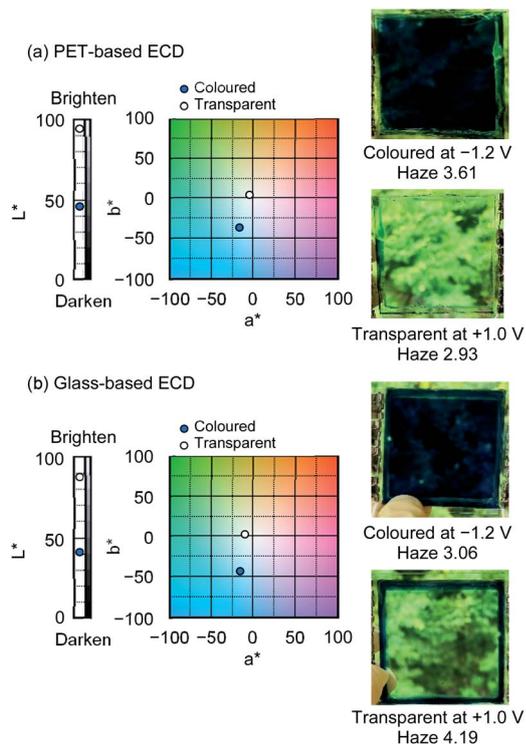


Fig. 5 Variation in CIELAB colour coordinates ( $L^*$ ,  $a^*$ ,  $b^*$ ) and photographs of the (a) PET-based and (b) glass-based ECDs in the colored and transparent states.

discolouration cycles up to 100 (Fig. 6). Both ECDs exhibited no change in the initial and subsequent electrochemical properties, and no notable degradation was observed in EC performance. Thus, the ECDs were electrochemically very stable. In addition, the PET-based ECD was mechanically robust during both bending and twisting (Movie S1†).

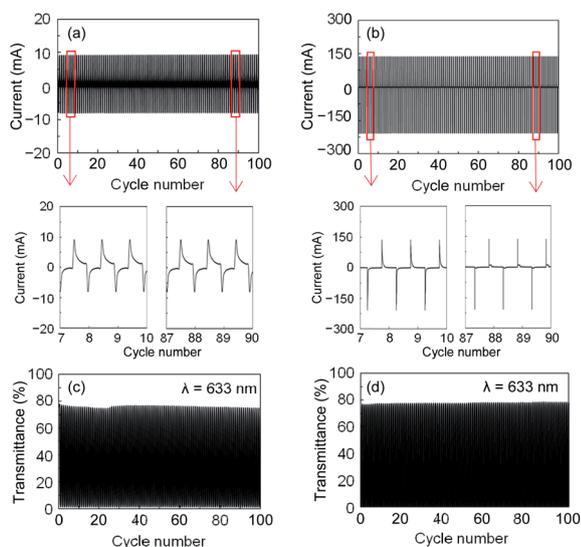


Fig. 6 Cycle stability up to 100 cycles of (a and c) PET-based and (b and d) glass-based ECDs illustrated as (a and b) MPS measurements of electrochemical stability at constant voltages of  $-1.2$  and  $+1.0$  V, and (c and d) transmittance at a wavelength of  $633$  nm.

Optical density ( $\Delta OD$ ) was used to calculate the EC colouration efficiency (CE), which is an important parameter governing the electrochemical performance of ECDs. CE and  $\Delta OD$  are defined as follows:<sup>49</sup>

$$CE = \Delta OD/Q = \log(T_t/T_c)/Q, \quad (5)$$

where  $T_t$  and  $T_c$  are the transmittance ratios of the transparent and coloured states, respectively, and  $Q$  is the amount of injected charge (Table S5†). Fig. S1† shows the plots of *in situ*  $\Delta OD$  at  $633$  nm versus colouration charge density at  $-1.2$  V. The CE is extracted as the slope of the line fitting the linear region of the curve. The glass-based ECD exhibited a higher  $\Delta OD$  of  $2.40$  than PET-based ECD due to its larger optical modulation. However, the PET-based ECD exhibited a higher CE of  $123.32$   $\text{cm}^2 \text{C}^{-1}$ , indicating high efficiency, even with a small amount of charge. The EC properties of PET-based ECDs deposited using various techniques and cycled in different electrolytes have been previously reported. Cai *et al.* reported that a  $\text{WO}_3$  PET-based ECD prepared by ink-jet printing had a CE of  $68.8$   $\text{cm}^2 \text{C}^{-1}$ ,<sup>50</sup> while Li *et al.* reported that a ECD of structure  $\text{WO}_3/\text{PEDOT:PSS}/\text{ITO}/\text{PET}$ , prepared by spray deposition, exhibited a CE of  $83.11$   $\text{cm}^2 \text{C}^{-1}$ .<sup>51</sup> Further, Yun *et al.* reported that the CE values of  $\text{WO}_3$  PET-based ECDs prepared by spin-coating varied between  $50.1$  and  $57.4$   $\text{cm}^2 \text{C}^{-1}$ .<sup>52</sup> On the other hand, the CE value of the PET-based ECD prepared in this study ( $123.32$   $\text{cm}^2 \text{C}^{-1}$ ) was higher than these previous reported values. Therefore, we strongly believe that the results of our study provide a new comprehensive foundation for the development of next-generation flexible ECDs for application in automobiles.

## Conclusions

A novel flexible ECD was prepared by coating water-based inks containing  $\text{WO}_3$  and PB NPs onto ITO-coated PET substrates. UV-treatment of ITO-coated substrates enhanced the wettability of the surface, which facilitated the adhesion of the  $\text{WO}_3$  and PB inks to the films. A complementary flexible ECD was produced using  $\text{K}^+$ -based gel electrolyte. Light-ageing was conducted to improve the EC performance of the PET-based ECD. To illustrate the colour change, the chromaticity diagram is plotted in terms of  $L^*$ ,  $a^*$ , and  $b^*$ , and the PET-based ECD showed a dramatic colour change from blue to transparent. Furthermore, the haze of the PET-based ECD in the transparent state was less than 3%, indicating clear transparency; therefore, transparent PET-based ECD is considered suitable for automobile applications. The ECDs exhibited excellent EC performance, including a large optical modulation of 79% at  $633$  nm, high CE ( $123.32$   $\text{cm}^2 \text{C}^{-1}$ ), and good cycling stability. These excellent EC properties of the flexible  $\text{WO}_3$ -based and PB-based ECDs demonstrate their promise as candidate materials for application in next-generation smart windows. However, when considering the application of this device as a window material for automobiles, several issues remain, including environmental stability (*e.g.*, resistance to temperature, humidity, UV) and introduction methods. Future research and development in collaboration with our partner companies is required to



produce scaled-up samples for field testing, thus facilitating rapid practical implementation.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## References

- 1 F. M. Andersen, H. K. Jacobsen and P. A. Gunkel, *Int. J. Electr. Power Energy Syst.*, 2021, **130**, 106900.
- 2 S. Fd da Silva, J. J. Eckert, F. L. Silva, L. C. A. Silva and F. G. Dedini, *Energy Convers. Manage.*, 2021, **234**, 113909.
- 3 G. Doluweera, F. Hahn, J. Bergerson and M. Pruckner, *Appl. Energy*, 2020, **268**, 114961.
- 4 F. Ren, S. Huang, F. Yang, A. Yurtsever and D. Ma, *J. Mater. Chem. A*, 2018, **6**, 24157–24165.
- 5 S. Huang, Q. Zhang, F. Yang, D. T. Gangadharan, P. Li, F. Ren, B. Sun and D. Ma, *J. Mater. Chem. A*, 2020, **8**, 8620–8628.
- 6 A. Llordes, G. Garcia, J. Gazquez and D. J. Milliron, *Nature*, 2013, **500**, 323–326.
- 7 X. Xie, K. Kretschmer and G. Wang, *Nanoscale*, 2015, **7**, 13278–13292.
- 8 R.-T. Wen, C. G. Granqvist and G. A. Niklasson, *Nat. Mater.*, 2015, **14**, 996–1001.
- 9 M. Gratzel, *Nature*, 2001, **409**, 575–576.
- 10 C. G. Granqvist, *Sol. Energy Mater. Sol. Cells*, 2000, **60**, 201–262.
- 11 C. Y. Yan, W. B. Kang, J. X. Wang, M. Q. Cui, X. Wang, C. Y. Foo, K. J. Chee and P. S. Lee, *ACS Nano*, 2013, **8**, 316–322.
- 12 C. G. Granqvist, *Thin Solid Films*, 2014, **564**, 1–38.
- 13 C. G. Granqvist, P. C. Lansåker, N. R. Mlyuka, G. A. Niklasson and E. Avendano, *Sol. Energy Mater. Sol. Cells*, 2009, **93**, 2032–2039.
- 14 C. Y. Jeong, H. Watanabe and K. Tajima, *Electrochim. Acta*, 2021, **389**, 138764.
- 15 M. Ishizaki, K. Kanaizuka, M. Abe, Y. Hoshi, M. Sakamoto, T. Kawamoto, H. Tanaka and M. Kurihara, *Green Chem.*, 2012, **14**, 1537–1544.
- 16 M. Ishizaki, Y. Sajima, S. Tsuruta, A. Gotoh, M. Sakamoto, T. Kawamoto, H. Tanaka and M. Kurihara, *Chem. Lett.*, 2009, **39**, 1058–1059.
- 17 A. Kraft, *Ionics*, 2021, **27**, 2289–2305.
- 18 K. Madasamy, D. Velayutham, V. Suryanarayanan, M. Kathiresan and K.-C. Ho, *J. Mater. Chem. C*, 2019, **7**, 4622–4637.
- 19 H.-M. Wang and S.-H. Hsiao, *J. Mater. Chem. C*, 2014, **2**, 1553–1564.
- 20 J. Wang, L. Zhang, L. Yu, Z. Jiao, H. Xie, X. W. D. Lou and X. W. Sun, *Nat. Commun.*, 2014, **5**, 4921.
- 21 P. H. Yang, P. Sun, Z. S. Chai, L. H. Huang, X. Cai, S. Z. Tan, J. H. Song and W. J. Mai, *Angew. Chem., Int. Ed.*, 2014, **53**, 11935–11939.
- 22 Z. Z. He, B. B. Gao, T. Li, J. L. Liao, B. Liu, X. J. Liu, C. Y. Wang, Z. Q. Feng and Z. Z. Gu, *ACS Sustainable Chem. Eng.*, 2019, **7**, 1745–1752.
- 23 K. S. Usha, R. Sivakumar, C. Sanjeeviraja, V. Sathe, V. Ganesan and T. Y. Wang, *RSC Adv.*, 2016, **6**, 79668–79680.
- 24 D. Ma, T. Li, Z. Xu, L. Wang and J. Wang, *Sol. Energy Mater. Sol. Cells*, 2018, **177**, 51–56.
- 25 J. Qian, D. Ma, Z. Xu, D. Li and J. Wang, *Sol. Energy Mater. Sol. Cells*, 2018, **177**, 9–14.
- 26 J. Wang, E. Khoo, P. S. Lee and J. Ma, *J. Phys. Chem. C*, 2008, **112**, 14306–14312.
- 27 Y. Ren, T. Fang, Y. Gong, X. Zhou, G. Zhao, Y. Gao, J. Jia and Z. Duan, *J. Mater. Chem. C*, 2019, **7**, 6964–6971.
- 28 G. Cai, P. Darmawan, M. Cui, J. Chen, X. Wang, A. L.-S. Eh, S. Magdassi and P. S. Lee, *Nanoscale*, 2016, **8**, 348–357.
- 29 X. Li, K. Perera, J. He, A. Gumyusenge and J. Mei, *J. Mater. Chem. C*, 2019, **7**, 12761–12789.
- 30 S. Lin, X. Bai, H. Wang, H. Wang, J. Song, K. Huang, C. Wang, N. Wang, B. Li, M. Lei and H. Wu, *Adv. Mater.*, 2017, **29**, 1703238.
- 31 S.-H. Park, S.-M. Lee, E.-H. Ko, T.-H. Kim, Y.-C. Nah, S.-J. Lee, J. H. Lee and H.-K. Kim, *Sci. Rep.*, 2016, **6**, 33868.
- 32 K. Tajima, H. Watanabe, M. Nishino and T. Kawamoto, *RSC Adv.*, 2020, **10**, 2562–2565.
- 33 K. Tajima, C. Y. Jeong, T. Kubota, T. Ito, K. Araki, T. Kamei and M. Fukui, *Sol. Energy Mater. Sol. Cells*, submitted.
- 34 H. W. Ro, J. M. Downing, S. Engmann, A. A. Herzing, D. M. DeLongchamp, L. J. Richter, S. Mukherjee, H. Ade, M. Abdelsamie, L. K. Jagadamma, A. Amassian, Y. Liue and H. Yan, *Energy Environ. Sci.*, 2016, **9**, 2835–2846.
- 35 J. Kim, J. Ahn, J. Shin, K. J. Yoon, J.-W. Son, J.-H. Lee, D. Shin, H.-W. Lee and H.-I. Ji, *J. Mater. Chem. A*, 2019, **7**, 9958–9967.
- 36 C. G. Granqvist, S. Green, E. K. Jonson, R. Marsal, G. A. Niklasson, A. Roos, Z. Topalian, A. Azens, P. Georén, G. Gustavsson, R. Karmhag, J. Smulko and L. B. Kish, *Thin Solid Films*, 2008, **516**, 5921–5926.
- 37 A. Jonsson, A. Roos and E. K. Jonson, *Sol. Energy Mater. Sol. Cells*, 2010, **94**, 992–997.
- 38 H.-F. Xiang, Z.-X. Xu, V. A. L. Roy, C.-M. Che and P. T. Lai, *Rev. Sci. Instrum.*, 2007, **78**, 034104.
- 39 Y. Takata, S. Hidaka, A. Yamashita and H. Yamamoto, *Int. J. Heat Mass Transfer*, 2004, **25**, 320–328.
- 40 Z. Xu, Z. Ao, D. Chu, A. Younis, C. M. Li and S. Li, *Sci. Rep.*, 2014, **4**, 6450.
- 41 E. Bormashenko, R. Pogreb, G. Whyman, Y. Bormashenko, R. Jager, T. Stein, A. Schechter and D. Aurbach, *Langmuir*, 2008, **24**, 5977–5980.
- 42 H. Y. Nie, M. J. Walzak, B. Berno and N. S. McIntyre, *Appl. Surf. Sci.*, 1999, **144**, 627–632.
- 43 C. Gervais, M. A. Languille, S. Reguer, M. Gillet, E. P. Vicenzi, S. Chagnot, F. Baudalet and L. Bertrand, *Appl. Phys. A: Mater. Sci. Process.*, 2013, **111**, 15–22.



- 44 C. Gervais, M. A. Languille, S. Reguer, C. Garnier and M. Gillet, *Heritage Sci.*, 2014, **2**, 26.
- 45 C. Y. Jeong, T. Kubota, K. Tajima, M. Kitamura and H. Imai, *Mater. Chem. Phys.*, submitted.
- 46 F. Ricci, F. Arduini, A. Amine, D. Moscone and G. Palleschi, *J. Electroanal. Chem.*, 2004, **563**, 229–237.
- 47 M. Hepel, L. I. Dela-Moss and H. Redmond, *J. Solid State Electrochem.*, 2014, **18**, 1251–1260.
- 48 R. Koua, Y. Zhonga, J. Kim, Q. Wang, M. Wang, R. Chen and Y. Qiao, *Energy Build.*, 2019, **193**, 69–77.
- 49 C. G. Granqvist, *Handbook of Inorganic Electrochromic Materials*, Elsevier, Amsterdam, 1995.
- 50 G. Cai, X. Cheng, M. Layani, A. W. M. Tan, S. Li, A. L.-S. D. Gao, S. Magdassi and P. S. Lee, *Nano Energy*, 2018, **49**, 147–154.
- 51 W. Li, J. Suna, J. Zhang, O. A. Ganiyata and Y. Cui, *Surf. Interfaces*, 2021, **2**, 100002.
- 52 T. Y. Yun, X. Li, J. Bae, S. H. Kim and H. C. Moon, *Mater. Des.*, 2019, **162**, 45–51.

