ABSTRACT: Dextran-templating hydrothermal synthesis of monoclinic WO3 exhibits excellent specific surface area of ~110 m²/g and a monomodal pore distribution with an average pore diameter of ~20 nm. Dextran plays a crucial role in generating porosity on WO3. The role of supporting dextran has been investigated and found to be crucial to tune the surface area, porosity, and morphology. The photoluminescence and X-ray photoelectron spectroscopy studies reveal the existence of oxygen vacancies in substoichiometric WO3, which creates localized defect states in WO3, as synthesized through this templating method. The highly mesoporous WO3 has been further explored as an interfacial cathode buffer layer (CBL) in dye-sensitized solar cells (DSSCs) and perovskite solar cells (PSCs). A significantly enhanced photocurrent has boosted up the performance of the counter electrode used in traditional DSSC (as platinum) and PSC (as carbon) devices by ~48% and ~29%, respectively. The electrochemical impedance and incident photon to current conversion efficiency (IPCE) studies were also analyzed in order to understand the catalytic behavior of the WO3 interfacial CBL for both DSSCs and PSCs, respectively. The much higher surface area of WO3 enables rapid electron hopping mechanism, which further benefits for higher electron mobility, resulting in higher short circuit current. Through this study, we were able to unequivocally establish the importance of buffer layer incorporation, which can further help to integrate the DSSC and PSC devices toward more stable, reliable, and enhanced efficiency-generating devices. In spite of this, using WO3 constitutes an important step toward the efficiency improvement of the devices for futuristic photoelectrochromic or self-powered switchable glazing for low-energy adaptive building integration.

INTRODUCTION

In the recent past, there has been a great impetus to develop and design functional materials for energy-harvesting applications in the field of dye-sensitized solar cells (DSSCs) and perovskite solar cells (PSCs).1-4 While researching new ways to increase the efficiency of utilizing solar energy, the technology is constantly being developed into new and better-advanced products. DSSCs stand out as one of the front runners in view of the basic novelty of the concept derived from nature’s principles, which allows facile and cost-effective processing alternatives.3,4 PSCs are thus considered as an advancement over DSSCs, where the photoanode thickness is reduced to a few microns and the dye is replaced by a superior light-absorbing organometal halide perovskite supported by a hole-transporting layer, which can be deposited directly over this cell architecture. Among the research and development versatility of the third-generation solar cell photovoltaic (PV) area, PSCs are the best technology developed during the last few years that promise a cheaper and more efficient alternative to the existing technologies for converting light to electric power.5,6 Implementation of various structures and fabrication strategies such as nanostructured materials (core–shell, mesoporous, one-dimensional, composite),7-10 advanced light-harvesting materials (quantum dots, NIR sensitizers),11,12 solid-state electrolytes,13 elimination of hole-transport materials (HTM),14 flexible substrates,15 scattering layers,16 up-conversion materials,17 and encapsulation methods,18 therefore, is envisaged as a potential effort to achieve prolonged stability and high efficiency in DSSC and PSC devices. The motive of all these strategies is efficient collection and extraction of charges. Additionally, introducing a buffer layer can provide enormous support, depending on where it has been used in the device. In general, a cathode buffer layer (CBL) excels as a supportive or optional layer for the cathode or back electrode material, which protects the cell from air and moisture and hence induces enhanced perform-
WO₃ is an extensively studied potential candidate for PV devices. With a view to enhance the PCE, research focused on the exploration of WO₃ as a photoanode material retarding the efficiency because of high band gap and less electron mobility than that of DSSCs based on the other semiconducting metal oxides, such as TiO₂ and ZnO, without employing any surface modification. On the other hand, efforts to replace organic polymers with WO₃, as a hole injection layer (HTL) in PV cells leads to exhibit higher device performance, which are currently receiving increased attention. However, the methods such as sputtering, thermal evaporation, and pulsed laser deposition, involve depositing the WO₃ layer, which is incompatible with a low-cost and solution-processed layer for future scalable manufacturing.

In this study, our target is to integrate the mesoporous WO₃ as an n-type metal oxide-based CBL for DSSC and PSC devices. To achieve a high surface area and considerable mesoporosity features of WO₃, a dextran-templating hydrothermal synthesis process has been attempted. The synthesized WO₃ has been further characterized by various physicochemical techniques and finally introduced as a CBL for DSSC and PSC devices. Further, our subsequent agenda is to imply this WO₃-based PV devices for electrochromic (EC) switchable glazing applications. The glazing has the ability to tune its color in the presence of applied direct current electric power. By changing the transparency, an EC switchable window attenuates glare and undesired solar gains, which provides improved indoor comfort.

### MATERIALS AND METHODS

**Materials.** Sodium tungstate (Na₃WO₄·2H₂O), HNO₃, and HCl were purchased from Merck, and dextran (molecular weight ca. 75,000) was acquired from Alfa Aesar, Ward Hill, MA. Isopropanol and propylene glycol were purchased from Sigma-Aldrich, UK. All the chemicals were used without any further purification.

**Synthesis of Porous WO₃.** Hierarchically porous WO₃ assemblies were synthesized by the hydrothermal method. In a typical experiment, 1.98 g of sodium tungstate (Na₃WO₄·2H₂O) was first dissolved in 40 mL of distilled water, and then, 1.5 mL of HNO₃ aqueous solution was added dropwise into the above solution under constant stirring to prepare tungstic acid H₂WO₄·H₂O (8 mol L⁻¹, pH 2). To this solution, 10 g of dextran has been added to further prepare an aqueous viscous solution to control the growth, leading to porous textures at room temperature. To allow the dextran-formating network throughout the aqueous solution of sodium tungstate for texture formation and further porosity development, the entire viscous solution was further kept completely under an undisturbed condition overnight. After this, the solution was transferred into a 50 mL Teflon-lined autoclave for the hydrothermal treatment at 160 °C for 24 h. The yellowish product as obtained from the hydrothermal reaction was accordingly centrifuged at 10,000 rpm and successively washed with 0.1 M HCl to neutralize the solution pH to 7. Besides, HCl treatment further helps to remove any unreacted Na⁺ ion also. The collected product was dried under an infrared lamp at ~80 °C for 12 h followed by calcination at 300 °C for a period of 1 h to get the final product. The same synthesis process has been carried out without dextran to understand the role of dextran in controlling the morphology and porosity. The overall synthesis scheme has been represented in Figure 1. In this synthesis, dextran excels as a porosity developer and
over the WO3 Platisol (T/SP, Solaronix ref. 41250) paste was drop-casted another layer of a similar transparent TiO2 was deposited by testing. A scheme of the WO3 buffer layer in (a) DSSCs and (b) PSCs, respectively.

Figure 1. Schematic representation of the formation of hierarchically porous WO3 assembly by the hydrothermal reaction.

Followed for nonporous WO3 layer deposition as well. The mesoscopic carbon layer was finally screen-printed with the as-prepared carbon paste and sintered at 450 °C for 30 min. Finally, the perovskite precursor solution (CH3NH3PbI3) with an appropriate amount was infiltrated by drop-casting via the top of the carbon counter electrode and further spin-coating at a speed of 1000 rpm for keep at 50 °C for 1 h and finally used for further studies. The active area of the working electrode was selected as 0.25 cm2. A schematic configuration of the WO3—carbon-based PSC is described in Figure 2b.

Morphology-texturing agent, which is required to obtain a high surface area of WO3. It is anticipated that the dextran-formating network behaves like a mould where the WO3 precursor was attached. During the hydrothermal treatment, the removal of dextran further leads to the creation of pores on the particles.77

Fabrication of DSSCs. Pieces with an area of 2 × 2 cm2 of transparent conducting oxides [fluorine-doped tin oxide (FTO)] glass substrates were first ultrasonicated in deionized water and ethanol. Next, they were subjected to surface cleaning by using an UV-Ozone cleaner (Ossila UV Ozone Cleaner, UK) for 15 min to remove any dirt, patches on the surface of the FTO glasses. Next, a layer of transparent TiO2 paste (18NR-T, GreatCell Solar) was allowed for screen printing (120 T mesh/inch) on the UV-ozone-treated FTO glasses and dried for 6 min on a hotplate at 125 °C. After this, another layer of a similar transparent TiO2 was deposited by the same method. The screen-printed active area of the working electrode was selected as 0.28 cm2. Then, a final layer of opaque TiO2 paste (18NR-O, GreatCell Solar) was printed, which provides an approximate thickness of 10 μm for the TiO2 electrode. The prepared electrodes were heated on a hotplate at 450 °C for 30 min to allow the crystallization of the TiO2 layer and to remove the binders available in the paste. After cooling down to 50 °C, the TiO2 electrodes were immersed in a 0.1 mM ruthenium 535 bis-TBA (N719) dye from GreatCell Solar in a 1:1 acetonitrile and ethanol solution at room temperature for 24 h to assure complete uptake of sensitizer. The Pt CE (Dyesol) was placed on a spin coating unit (SPIN 120D, MIDAS System). A volume of 10 wt % (weight/volume) of a colloidal suspension of WO3 was prepared in a 2:1 isopropanol and propylene glycol medium. A volume of 1 mL of prepared WO3 colloidal solution was then spin-coated on the top of the FTO substrate at a speed of 3000 rpm for 30 s and dried at 200 °C for 30 min. Finally, the Platisol (T/SP, Solaronix ref. 41250) paste was drop-casted over the WO3 film and further calcined at 450 °C for 10 min to deposit a layer of Pt. The similar process has also been followed for nonporous WO3 layer deposition. The fabricated WO3—Pt CE was further placed over the dye-sensitized photoanode, and a plastic spacer was kept in between these two electrodes. The prepared redox electrolyte of I−/I3− was further infiltrated into the device and allowed for final testing.77 A scheme of the WO3 buffer layer-based DSSC device is given in Figure 2a.

Fabrication of PSCs. The PSC devices were fabricated as per our previous report. However, for this case, 1 mL of the prepared WO3 colloidal solution, as used for DSSC fabrication, was spin-coated at a speed of 3000 rpm for 30 s on the top of Al2O3 layer deposition and further heated at 200 °C for 30 min during the PSC fabrication. The similar process has been also followed for nonporous WO3 layer deposition as well. The mesoscopic carbon layer was finally screen-printed with the as-prepared carbon paste and sintered at 450 °C for 30 min. Finally, the perovskite precursor solution (CH3NH3PbI3) with an appropriate amount was infiltrated by drop-casting via the top of the carbon counter electrode and further spin-coating at a speed of 1000 rpm for keep at 50 °C for 1 h and finally used for further studies. The active area of the working electrode was selected as 0.25 cm2. A schematic configuration of the WO3—carbon-based PSC is described in Figure 2b.
range from 0.1 Hz to 100 kHz. All the devices were measured at their corresponding open-circuit voltage. Incident photon to current efficiency (IPCE) was carried out using a Bentham PVE300 Photovoltaic EQE (IPCE) and IQE solution under 300–800 nm wavelength using a tungsten halogen lamp source.

■ RESULTS AND DISCUSSION

Structural, Optical, and Surface Area Analysis of WO₃. X-ray diffraction pattern, as shown in Figure 3, reveals the polycrystalline monoclinic WO₃ structure as it matches well with the reference diffraction pattern of WO₃ (JCPDS card number 83–0950). The main diffraction peaks at 23.1, 23.7, and 24.4° ascribed to Miller indices (002), (020), and (200), respectively, correspond to the monoclinic WO₃ phase. This also rationalizes the synthesized WO₃ free of orthorhombic WO₃ and WO₃·1/3H₂O phases showing that the products are pure phase. High and sharp peak intensities indicate an excellent crystallinity of the sample. It is interesting to mention here that the orthorhombic phase of WO₃ or WO₃·1/3H₂O is quite often to be obtained via the hydrothermal treatment of tungstic acid H₂WO₄·H₂O, although upon further heating >300 °C, all these metastable oxides lead to the stable monoclinic WO₃ phase. Here, through this dextran-templating synthesis, the WO₃ attained the monoclinic phase directly.

In order to ascertain the effect of dextran addition on porosity improvement, the TEM investigation was performed with both the dextran-added and without dextran samples of WO₃, respectively. Figure 4a,b indicate the TEM bright-field images of WO₃ in the absence of dextran. It has been clearly observed that there are, as such, no porous characteristics formed on the WO₃ surface. Moreover, without dextran employment, it leads to form a densely populated large hierarchical structure, whereas the addition of dextran assists to render the porous nature throughout the surface of WO₃, as shown in the TEM bright-field images of Figure 4c,d at different magnifications. Besides, dextran addition influences to attain the morphology of WO₃ toward a separated hierarchical platelet-like structure. During the hydrothermal treatment, the chainlike structure of the dextran preferentially combines with the hydroxide nanocrystals of WO₃ anisotropically as building blocks thereby able to create patterned slits, which also facilitates the etching of the sites without the adsorption of dextran. This may cause the pore formation of WO₃ allowing higher surface energy. In addition, Figure 4e shows the nitrogen adsorption—desorption isotherm of the WO₃ sample in the absence of dextran and appeared as type III isotherm. It exhibits a BET surface area of 30.28 m² g⁻¹, involving no mesoporous characteristics, whereas the isotherm indicated mesoporous characteristic of the dextran-assisted WO₃ samples and appeared as type IV isotherm as reported in Figure 4f. The appearance of type H-3 hysteresis indicating a stacked-plane (slit)-like pore shape. This exhibits comparatively a high surface area of 110.68 m² g⁻¹. To the best of our knowledge,
the obtained surface area of this WO₃ is quite higher compared to the other similar reports of WO₃ as shown in Table S₁.⁴²−⁵¹ The pore diameter was found to be about 3.7 nm using the BJH method adsorption method (Figure S₂). This value was artificial (pseudo peak) calculated from the BJH computation method. Further, the pore diameter was found to be about 20.2 nm using the BJH desorption method, which was further supported by TEM investigation, as shown in Figure 4f (inset). The corresponding pore volume was found to be 0.46 cc g⁻¹. Again, stacked-plane pore geometry throughout the plate-like microstructural framework was fairly visible under a transmission electron microscope evidencing H₃ hysteresis caused by the tensile strength effect during the gas condensation process and therefore resulted in a monomodal pore-size distribution as obtained from the BJH desorption method.⁵²,⁵³ Therefore, considering the high surface area and mesoporous characteristics of the WO₃ sample, achieved by dextran addition, was selected for further studies in this manuscript.

To further evaluate the morphology formation of the porous WO₃ sample, the various microstructural analysis was investigated, as shown in Figure 5. During the gel formation, the dextran-added precursor solution creates amorphous networking of the sol as obtained from Figure 5a. The gel collapses into particles that precipitate from the solution during the hydrothermal treatment. This may look like the precursor solution is getting embedded inside the dextran polymeric network. On the other hand, Figure 5b,c,d,e exhibit the SEM images of the synthesized WO₃ sample at different magnifications, respectively. The morphology exhibits as the hierarchical platelet-type shapes as obtained by the hydrothermal synthesis. Besides, the TEM bright-field images further reveal that the platelet-type morphology contains a wide range of size of 80–300 nm, having a predominate order of large distinct porosity (Figure 5f,e). The lattice fringes in high-resolution TEM (HRTEM) image (Figure 5f) corresponds to the (020) reflection with an inter planer spacing of 0.379 nm, which confirms the single-crystalline nature of the monoclinic WO₃ phase. The EDS analysis was also performed to confirm the elements of WO₃ and the complete removal of Na, as shown in Figure S₁.

X-ray Photoemission Spectroscopic and Photoluminescence Analysis. The valence states of the fundamental elements W and O of the synthesized mesoporous WO₃ have been identified through XPS measurements (Figure 6). The survey spectrum of the WO₃ is shown in Figure 6a. First, two characteristic peaks at 36.0 ± 0.2 eV (Figure 6b) has been ascribed to the typical doublet of W⁶⁺ with the binding energy of W (4f7/2 and 4f5/2). The peak ratio of W (4f7/2) to W (4f5/2) was found to be 4:3 with a spin–orbit splitting of a doublet value of 2.12 eV. A third broad peak of W (5p3/2) locates at ∼41.8 eV. This result reveals only W for its +6 oxidation state.⁵⁴ However, the asymmetry of the W 4f7/2 indicates that WO₃ readily becomes oxygen-deficient to form WO₃₋ₓ with the variable oxygen composition parameter x. On deconvolution, the asymmetric O 1s XPS spectrum evolves into two components centered at 529.21 (O₁) and 530.65 eV (O₂) as shown in Figure 6c. The dominant O₁ component is a characteristic of O²⁻ ions, whereas the O₂ component at the higher binding side may be associated with the presence of chemisorbed oxygen in the grain boundaries or the existence of the hydroxyl group (−OH) because of the adsorption of water onto the surface of the WO₃ during synthesis through the aqueous medium route. This oxygen deficiency greatly

Figure 5. FESEM microstructural images of (a) the as-prepared sol, (b,c) mesoporous WO₃ at different magnifications, respectively; (d,e) TEM bright-field images of mesoporous WO₃ at different magnifications, respectively and (f) the corresponding HRTEM image.

Figure 6. (a) XPS survey spectrum of the mesoporous WO₃ sample, (b) XPS of spin–orbit split peaks of W 4f, and (c) spectral decomposition of the O 1s spectrum results in two different types of oxygen environment denoted as the O₁- and O₂-fitted component in the WO₃.
influences the bulk of the electronic transport properties by introducing donor electronic states.55

Further, the photoluminescence (PL) spectrum of the aqueous dispersion of the WO$_3$ sample was monitored at room temperature, as shown in Figure S3. A blue emission at 467 nm was observed against an excitation wavelength of 350 nm. It is anticipated that the reason for the visible emission exhibited by WO$_3$ could be due to the presence of oxygen-related defects as observed from the XPS analysis. The aqueous solution of synthesized WO$_3$ resulted in a quantum yield (QY) of 3.25% at room temperature.

Microstructural and Optical Measurement of WO$_3$ Films. The SEM line-scale mapping was recorded on the WO$_3$–Pt film, as shown in Figure S4a. This indicates a homogeneous distribution of W and O throughout the Pt layer. Further, the distribution of W and O over the Pt layer is very clear from the elemental mapping diagram, as shown in Figure S4b. Besides, in the case of WO$_3$–carbon-based PSC film, a thin layer (<200 nm) of mesoporous WO$_3$ has been formed as shown in Figure S4c, and the corresponding elemental mapping has been given in Figure S4d. The EDS result indicates that despite the different approach of the buffer layer formation for DSSCs and PSCs, both of them showed a proper distribution of W and O along with their respective back electrode material, indicating that the formulation of WO$_3$ synthesis is precise and efficient in preparing the well-dispersed sample.

Table 1. PV and Impedance Spectroscopic Parameters for Different DSSC Devices

| cell                          | $J_{SC}$ (mA cm$^{-2}$) | $V_{OC}$ (mV) | FF   | PCE ± 0.05 (%) | $R_s$ (Ω cm$^{-2}$) | $R_{CT1}$ (Ω cm$^{-2}$) | $C_p$ (F) |
|------------------------------|--------------------------|----------------|------|----------------|---------------------|------------------------|------------|
| without WO$_3$               | 11.49                    | 722            | 0.59 | 5.01           | 18.52               | 32.68                  | 1.52 × 10$^{-6}$ |
| Pt–WO$_3$ (nonporous)        | 14.68                    | 753            | 0.55 | 6.12           | 18.65               | 26.86                  | 3.57 × 10$^{-4}$ |
| Pt–WO$_3$ (mesoporous)       | 16.55                    | 753            | 0.64 | 7.43           | 18.23               | 27.12                  | 2.48 × 10$^{-3}$ |

Figure 7. (a) $J$–$V$ characteristic and (b) corresponding Nyquist plot (inset: corresponding equivalent circuit) for Pt and WO$_3$–Pt-based DSSC devices, respectively.

PV Performance of WO$_3$ as a Cathode Buffer Layer in DSSCs. In order to evaluate the effect of mesoporous WO$_3$ as a CBL in PV applications, the PV performances have been further investigated. An evaluation of the DSSC performances of the fabricated WO$_3$-based devices were executed under 1SUN 1.5 AM (Figure 7). The performances of the $J$–$V$ characteristic plot for the DSSC devices with Pt CE and WO$_3$–Pt CE are shown in Figure 7a. The WO$_3$–Pt device exhibited a maximum PCE of 7.43% with a short circuit current ($J_{SC}$) of 16.25 mA cm$^{-2}$, an open-circuit voltage ($V_{OC}$) of 0.75 V, and a fill factor (FF) of 0.59. On the contrary, with only Pt CE-based devices exhibited a PCE of 5.01% with a $J_{SC}$ of 11.49 mA cm$^{-2}$, an $V_{OC}$ of 0.72 V, and a FF of 0.64. The digital image of WO$_3$–Pt-based DSSCs has been given in the inset of Figure 7a. The effect of mesoporous nature also influences the efficiency of the device. In the absence of the mesoporous WO$_3$ layer, the maximum average PCE was achieved to be ∼6.12% (Figure S5a), which is ∼21% lesser than compared to mesoporous WO$_3$ layer-added devices. These results strongly support the benefits of porous layer incorporation in such a DSSC device. Further, the recorded $J$–$V$ characteristics parameters are compared in Table 1. In spite of this, using WO$_3$ on Pt CE, the corresponding device exhibits ∼48% higher efficiency than only Pt CE. The enhancement of PCE was resulted by keeping insignificant alteration in $V_{OC}$ except for the $J_{SC}$ and FF, which signifies the presence of mesoporous WO$_3$ gets an advantage as a catalyst for Pt CE and, therefore, accelerate the rate of high electron mobility and hole-extraction leading toward high electron mobility and hole-extraction.
efficiency. The EIS measurements were carried out to understand the transport properties at different interfaces in the DSSC assembly. The EIS spectra (Nyquist plot) and the corresponding equivalent circuit diagram (inset) of cells for the DSSC devices, as shown in Figure 7b. As shown in Table 1, the DSSCs show almost closure of sheet resistance ($R_s$), which is due to the use of the same Pt counter electrodes and $I/Re$ electrolyte in all the devices. On the other hand, resistance at the Pt/electrolyte interface has been found significantly higher for the only Pt CE device compared to WO$_3$-Pt-based devices. The CT resistance of 32.68 $\Omega$ cm$^{-2}$ at the electrolyte/dye/TiO$_2$ interface for the Pt CE device is considerably higher than the WO$_3$-Pt device (27.12 $\Omega$ cm$^{-2}$). This result clearly shows that the catalytic activity of WO$_3$ reduces the CT resistance of Pt producing higher efficiency. The EIS spectrum of the without mesoporous WO$_3$-based DSSC device has been given in Figure S5a (inset), and the corresponding data are shown in Table 1.

PV Performance of WO$_3$ as a Cathode Buffer Layer in PSCs
In addition, incorporation of the synthesized WO$_3$ as a cathode buffer layer, it was further evaluated in the case of PSC devices. The corresponding $J-V$ characteristics plot is shown in Figure 8a. It was noted that the addition of WO$_3$-carbon back contact exhibited the highest PCE of 12.42%, which is $\sim$17 and $\sim$29% higher than the without mesoporous WO$_3$ (PCE: $\sim$10.58%) and only carbon-based devices ($\sim$9.65%), respectively. The $J-V$ characteristics plot for the without mesoporous WO$_3$-added device has been given in Figure S5b (inset), and the corresponding data are shown in Table 1.

The Nyquist plots derived from the impedance data, Figure 8b, revealed a clear difference between the two devices. The WO$_3$-carbon-based PSC shows the smallest radius of the semicircle in the high-frequency region, which indicates the lowest CT resistance at the CH$_3$NH$_3$PbI$_3$/carbon interface. This result implies that the film addition of WO$_3$ plays an important role in DSSC electron recombination. All the measured parameters have been summarized in Table 2. $R_s$ of PSCs with WO$_3$ is less than the device without WO$_3$, which suggests that the hole-extraction process at the carbon electrode with WO$_3$ is more efficient. Moreover, a lower charge exchange resistances ($R_{CT}$) at a bias of $-0.90$ V was observed for WO$_3$-additive PSCs compared to the devices without additive. However, the WO$_3$ additive layer may play the role of a hole collector along with the appropriate energy level to the perovskite CH$_3$NH$_3$PbI$_3$ sensitizer and lead to an acceleration of the hole extraction efficiency. The WO$_3$ layer can also significantly reduce the series resistance and significantly improve the $J_{SC}$ of the PSC device. Besides, the EIS measurement was also performed for the absence of mesoporous WO$_3$-based PSC devices, as shown in Figure S5b (inset), and the corresponding data are summarized in Table 2.

**Probable Mechanism of the WO$_3$ Buffer Layer in Solar Cell Devices.** The efficiency was measured for 10 numbers of cells for DSSCs and PSCs with an error bar plot, as shown in Figure 9a. This result implies that the film addition of WO$_3$ plays an important role in DSSC electron recombination. Therefore, we suggest an apparent mechanism to understand the effect of the porous WO$_3$ structure as an interfacial cathode buffer layer on the improved crystalline and enhanced PCEs of the solar cell devices. WO$_3$ treatment may also play a similar role in affecting the electronic structure of carbon and can be explored as a hole-extracting layer for PSCs. It is assumed that a small polaron hopping conduction mechanism can be influenced to operate in WO$_3$ as a cathode buffer layer for solar cell devices. This mechanism possibly suggests hopping can be a faster way compared to the traditional way of hole diffusion with an extremely fast rate constant that leads to exhibit enhanced device performance. Besides, the photoluminescence spectrum of WO$_3$ predicts partial oxygen vacancies in substoichiometric WO$_3$ which creates localized defect states: W$^{4+}$ or W$^{5+}$ and W$^{6+}$. These mixed valance states may further promote the CT and enhance electrical conductivity. The high surface area facilitates to collect and regenerate a higher number of holes, which is ultimately
beneficial for rapid and smooth electron–hole mobility for the devices. At the same time, it is also assumed that because of distinct porous nature of \( \text{WO}_3 \) that it may be suitable to entrap/adsorb any moisture or oxygen molecule, which may retard the overall performance of the device. Pt counter electrodes are usually affected by high corrosion because of the use of liquid \( \Gamma^-/I^- \) electrolyte, which can retard the hole conduction process, although using the \( \text{WO}_3 \) buffer layer may restrict the rate of the corrosion and facilitate the hole conduction process for the device. Therefore, the high porosity could be beneficial for the \( \text{WO}_3 \) and favor as a buffer layer for the solar cell devices. The overall concept has been schematically illustrated in Figure 9b,c. The enhanced PCE has been mainly influenced by the enhancement of the \( J_{SC} \) factor of the device. The increased \( J_{SC} \) of the porous \( \text{WO}_3 \) probably resulted in a thicker active layer, which is helpful to improve optical-trapping properties.

**Incident Photon to Current Conversion Efficiency Measurements.** IPCE stands for the external quantum efficiency of the DSSC and PSC devices including the effects of optical losses caused by transmission and reflection.\(^{37}\) The IPCE measurements were also performed as a function of wavelength by comparing the \( \text{WO}_3-\text{Pt} \) and Pt-based devices for DSSCs and \( \text{WO}_3-\text{carbon} \) and carbon-based devices for PSCs as shown in Figure 10. The IPCE curve for all the DSSCs shows a broad peak over the range of 300–800 nm with a maximum value of \( \sim 83\% \) at 536 nm for the \( \text{WO}_3-\text{Pt-based} \) device, whereas the Pt-based DSSC exhibits an IPCE value of \( 61\% \) at the same wavelength as shown in Figure 10a. In the case of PSCs, incorporation of the \( \text{WO}_3 \) layer exhibits predominant incident photon to current conversion efficiency (IPCE) value (\( \sim 90\% \) at 400 nm and \( 76\% \) at 560 nm wavelength) than without \( \text{WO}_3 \) layer device value (\( \sim 76\% \) at 400 nm and \( 60\% \) at 560 nm wavelength) as shown in Figure 10b. This also strongly recommends that the incorporation of a \( \text{WO}_3 \) layer improves the IPCE in a great manner to minimize the electron recombination or electron entrapment within surface trap states or electron loss within the hole transporting section of the device.

Therefore, the EIS and IPCE results predict that the CT process is taking place at the interface of Pt/carbon and \( \text{WO}_3 \), leading to an acceleration of the hole mobility, and as a result, the short circuit current increases remarkably. This is quite competitive with organic interfacial materials where it cannot be readily achieved with the later. All the above-discussed results and the fundamental insights could be useful in the development of achieving solution-processed successful incorporation of the \( \text{WO}_3 \) cathode buffer layer for high-efficiency devices.\(^{36,38,59}\) Although the use of \( \text{WO}_3 \) presented here are far from being fully optimized, but the profound advantages along with ease of preparation and applicability for potential performance make it a good candidate for the cathode buffer layer for solar cell devices.\(^{50,61}\) This work also provides insights into the design of the interfacial hole transport layer for the fabrication of solution-processed solar cell devices. The obtained results are quite encouraging and open up ample scope to develop new and economically viable synthetic approaches to enhance the performance of the solution-processed solar cell devices.

In spite of this, there are very few reports available on the buffer layer-incorporated DSSC and PSC devices. In comparison to the reported results on the performance of various materials as incorporated buffer layer in solar cell devices (Table 3), the results obtained in this work are quite competitive and interesting to execute dextran template mesoporous \( \text{WO}_3 \) as a promising buffer layer candidate for both DSSC and PSC devices, respectively.

| CBL | type of solar cell | PCE (%) | reference |
|-----|-------------------|--------|-----------|
| PAMPS-Na nonconjugated polymer/Al | OSC | 9.16 | 22 |
| PCBM/bis-C_{60}/Ag | PSC | 15.44 | 23 |
| C_{60}/bis-C_{60}/Ag | PSC | 13.37 | 23 |
| TiO_{2}/Pt | DSSC | 2.84 | 24 |
| CdSe QDs/LiF/AuAg | inverted-PSC | 15.1 | 27 |
| SnO_{2}/Ag | inverted-OSC | 2.89 | 30 |
| MoO_{3}/Au | all inorganic PSC | 5.52 | 31 |
| ZnO-DOL/Ag | ISC | 3.48 | 62 |
| Zr-isopropanoxide/Al | BHJ-PSC | 8.07 | 63 |
| PCBB-2CN-2C_{6}/Au | PSC | 17.35 | 64 |
| a-ZrAcac/Al | poly-SC | 8.75 | 65 |
| PCBM, C_{60} and LiF/Al | planar-PSC | 14.69 | 66 |
| ZnO/PEI/Al | OSC, PSC | 11.76 | 67 |
| ZnO/Ag | OSC | 9.2 | 68 |
| tris-(8-hydroxyquinoline)/Al | planar-PSC | 14.22 | 69 |
| PCBDANI/LiF/Al | planar-PSC | 15.71 | 70 |
| bathocuproine/Ag | inverted OSC | 3.28 | 71 |
| PCMP/BCP/Ag | planar PSC | 14.47 | 72 |
| Julolidine | inverted OSC | 8.07 | 73 |
| mesoporous WO_{3}/Pt | DSSC | 7.43 | current work |
| mesoporous WO_{3}/carbon | mesoscopic PSC | 12.42 | |

*OSC: organic solar cells; PSC: perovskite solar cells; DSSC: dye-sensitized solar cells; poly-SC: polymer solar cells; ISC: inverted solar cell; BJH: bulk-hetero junction.*

\[ \text{Figure 10. IPCE spectra for (a) WO}_3-\text{Pt and Pt-based DSSC and (b) WO}_3-\text{carbon- and carbon-based PSC devices, respectively.} \]
In our previous report, different concentrations of commercial WO₃ nanoparticles as a HTM incorporated in carbon counter electrodes were tested for PSCs. However, the current study mainly signifies about the mesoporous layer involvement of WO₃ as a CBL for both the DSSC and PSC devices. Of course, controlled synthesis, optimum pore size distribution, various surface treatments, and optimization of film fabrication and device fabrication techniques are required, which may further lead to the enhancement of efficiency of the device for futuristic applications. We believe this study seems to be applicable for the synthesis of other related materials with controllable morphology and tunable textural properties for various applications such as catalysis, adsorption, gas sensing, separation, and so forth.32−45,48,51

CONCLUSIONS

Hydrothermal synthesis of mesoporous WO₃ using the dextran templating process has been performed in this study. Dextran can effectively help to generate pore on the surface of WO₃. Hierarchical platelet-like porous morphology of WO₃ exhibited a high surface area of ∼110 m² g⁻¹ and an average monomodal pore size diameter of 20 nm. Taking advantage of this mesoporous WO₃, it has been further explored as a solution-processed cathode buffer layer on Pt in the case of DSSCs and on carbon for the PSC device performance evaluation. Incorporation of WO₃ interestingly enhanced the PCE of a Pt-based DSSC device by 48%, whereas in the case of PSCs, the PCE enhancement was recorded ∼29% compared in the absence of buffer layer-employed devices. It is anticipated that the WO₃ layer significantly boosts up the short-circuit current of the PV devices, leading to enhance the PCE. Besides, an apparent hole hopping mechanism associated with the porous WO₃ layer structure credit some added benefits to the devices. On the basis of our results, we are able to establish porosity and buffer layer incorporation as an important factor for PV efficiency enhancement for DSSC and PSC devices. Moreover, the outcome from this work is highly promising and renders a potential for future self-powered electrochromic device integration, where WO₃ will enhance the power generation and switching color in a tandem structure concomitantly.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.jpca.0c02912. Reported surface area and porosity results of different morphology-based WO₃ for various applications; EDS spectrum of the mesoporous WO₃ sample; pore size distribution plot of WO₃ as obtained from the BJH adsorption model; PL spectrum of the aqueous solution of the mesoporous WO₃ sample at room temperature; SEM elemental and line scale mapping of WO₃−Pt and WO₃−carbon films; and J−V characteristics plot along with corresponding fitted EIS plot of the nonporous (without dextran added) WO₃ buffer layer for DSSC and PSC devices (PDF).

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Notes

The authors declare no competing financial interest. In support of open access research, all underlying article materials can be accessed upon request via email to the corresponding author.

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