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Prospects of e-beam evaporated molybdenum oxide as a hole transport layer for perovskite solar cells


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Perovskite solar cells have emerged as one of the most efficient and low cost technologies for delivering of solar electricity due to their exceptional optical and electrical properties. Commercialization of the perovskite solar cells is, however, limited because of the higher cost and environmentally sensitive organic hole transport materials such as spiro-OMeTAD and PEDOT:PSS. In this study, an empirical simulation was performed using the Solar Cell Capacitance Simulator software to explore the MoOx thin film as an alternative hole transport material for perovskite solar cells. In the simulation, properties of MoOx thin films deposited by the electron beam evaporation technique from high purity (99.99%) MoO3 pellets at different substrate temperatures (room temperature, 100°C and 200°C) were used as input parameters. The films were highly transparent (>80%) and have low surface roughness (≤2 nm) with bandgap energy ranging between 3.75 eV and 3.45 eV. Device simulation has shown that the MoOx deposited at room temperature can work in both the regular and inverted structures of the perovskite solar cell with a promising efficiency of 18.25%. Manufacturing of the full device is planned in order to utilize the MoOx as an alternative hole transport material for improved performance, good stability, and low cost of the perovskite solar cell. Published by AIP Publishing.

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I. INTRODUCTION

Perovskite solar cells (PSCs) are one of the promising technologies in photovoltaics that have encouraged the world’s ambition towards solar energy utilization at low cost. Due to the low cost and simple solution procedure, PSCs have revolutionized the field of photovoltaic research. The efficiency of PSC has increased from 3.8%1 in 2009 to more than 20% in 2016.2 Despite the high efficiency of the solar cell, the hole transport materials (HTM) used are mostly limited to organic compounds including 2,2',7,7'-tetrakis(N,N-di-p-phenylphénylamine)-9,9’-spirobifluorene (spiro-OMeTAD)3 and poly(3,4-ethyleneoxythiophene) polystyrene sulfonate (PEDOT:PSS). These materials are expensive and have poor stability upon exposure to the environment.4,5 The acidic nature of PEDOT:PSS6 and the hygroscopic nature of the additives used in the spiro-OMeTAD decreases the long term stability of PSCs.7 In practice, high cost and poor environmental stability are the main limitations for large scale applications of the PSCs. Metal oxides will be a better replacement to overcome these challenges. Metal oxide thin films have been widely used in various applications such as display devices, optical small window, gas sensors, and photovoltaics8 and they can be a better candidate as HTM for solar cells.9,10 Inorganic hole transport materials such as NiO, Cu2O, CuOx, and VOx have been incorporated into PSC as a replacement of the existing organic HTMs8,11 due to their higher stability and lower materials cost than organic materials.12 MoOx is one of the potential materials not only for solar cells but also gas sensors and light-emitting diodes.13 Recently, MoO2 nanoparticles were used as a hole transport layer with efficiency of 15%, with very low hysteresis and high device stability.14 Similarly, MoOx has been used as a hole selective contact for silicon solar cells.15 Xiao et al. use MoOx as a hole transport layer for perovskite solar cells but only achieved low efficiency of 5.9%.16 Hou et al. used bilayer MoOx/PEDOT:PSS to improve the performance and stability of the solution processed perovskite solar cells.9 Chen et al. used thermally evaporated MoOx in the inverted structure of the perovskite solar cell achieving 13% efficiency and good stability.10 These results suggest that MoOx is a promising material as an alternative HTM.

Different techniques have been used for the preparation of MoOx thin films with an amorphous and crystalline structure. Films can be deposited using solution-based methods and vacuum deposition techniques. The solution methods include spray pyrolysis,17 sol-gel process,18 electrodeposition, and spin coating19 whereas the vacuum methods are thermal evaporation20,21 and electron beam evaporation (e-beam).22 Solution processing is usually easy and cheap; however, there is less control over the film properties, leading to poor reproducibility. Vacuum deposition however has good control over the film properties and reproducibility. Also, a wide range of experimental parameters are available which can tailor the structure, morphology, composition, and other properties of the films according to the required applications.

In this study, empirical simulation using the solar cell capacitance simulator software (SCAPS) was applied to investigate MoOx as an alternative HTM for perovskite solar cells. MoOx thin films deposited at different substrate...
temperatures by the electron beam evaporation technique were considered for the simulation. Helium Ion Microscopy (HIM) and Atomic Force Microscopy (AFM) were used to study the surface morphology of the films. The surface roughness was measured by AFM and confirmed using the stylus profilometer. The film thickness was measured by both stylus profilometer and ellipsometry. The chemical composition was investigated using X-Ray Photoelectron Spectroscopy (XPS), and the crystalline nature of the films was determined using Raman spectroscopy. The optical properties of the films were characterized using UV-Vis-NIR spectrometry. From these measurements, the thickness and bandgap energy ($E_g$) of the MoO$_x$ thin films were used as input parameters in the empirical simulation analysis to study the films as an alternative HTM layer in both the inverted and regular structures of the PSC.

II. EXPERIMENTAL

A. Thin film deposition

MoO$_x$ thin films were deposited on a glass substrate using the electron beam evaporation technique (PVD 75 Kurt J. Lesker) from MoO$_3$ pellets (99.9% purity). The films were deposited at different substrate temperatures (room temperature, 100 °C and 200 °C). For the films deposited at 100 °C and 200 °C, the substrate was preheated to the desired temperature and maintained at that temperature throughout the deposition. Before each deposition, the glass was cleaned thoroughly with acetone, ethanol, and then dried with nitrogen gas. The chamber was evacuated to a base pressure of less than 1 × 10$^{-6}$ Torr. As expected, the pressure slightly increased to 4 × 10$^{-5}$ Torr during the film deposition due to the vapour of the MoO$_3$. The films were deposited at a deposition rate of 1 Å/s as controlled by the quartz crystal monitor. All the films had a nominal thickness of 100 nm. During deposition the substrate was continuously rotating at 10 rpm in order to ensure uniform and homogenous coatings fully covering the substrate. After deposition, the samples were allowed to naturally cool in the vacuum chamber to less than 50 °C before they were taken out of the chamber for characterization.

B. Thin film characterization

Various techniques were used to characterize the as-deposited MoO$_x$ thin films. The surface morphology of the films was characterized by Atomic Force Microscope (AFM) and Helium Ion Microscope (HIM). An NT-MDT Solver P47 scanning probe microscope (NT-MDT Co., Moscow, Russia) operated in the semi-contact mode with “Golden” Si cantilevers was used in the AFM measurement. The nominal tip diameter was 10 nm. High resolution micrographs were obtained using Zeiss Orion HIM at 25 kV with a 0.3 pA blank current. HIM is found to be the preferred technique for characterizing the as-deposited MoO$_x$ thin films without using conductive coating. Any charging caused by the insulating properties of the films can be compensated using a flood gun having a beam of electrons which compensate for the accumulated ion beam charge. The thickness of the films was measured by the stylus profilometer and confirmed by ellipsometry. X-ray photoelectron spectroscopy was performed using a Kratos Axis Supra with aluminium Al K$_\alpha$ X-ray radiation ($h\nu = 1486.7$ eV). Wide survey scans were acquired using the analyzer pass energy of 160 eV, and high resolution scans of the O s, C 1s, and Mo 3d regions were performed using a pass-energy of 20 eV to better discriminate the sub-structure of the spectral lines. In all cases, the binding energy scale was corrected by a rigid shift to align the peak of the C 1s core level to 284.8 eV, corresponding to adventitious carbon. Quantification was performed using the CasaXPS software version 2.3.17PR1.1, using the appropriate element sensitivity library for the Kratos instrument. Carbon present on the surface was assumed to be adventitious in nature, and as such a fraction of the measured oxygen signal was associated to the surface contamination, using the method of Payne et al.$^{23}$ Mo 3d and O 1s spectra were fitted with a series of Voigt line shapes, using area constraints across Mo and O chemical states to enforce the expected 3:1 and 5:2 intensity ratios for Mo (VI) and Mo (V). However, the Mo (V) line shape is known to be somewhat more complicated than a simple Voigt function due to multiple splitting.$^{24}$ We have found that our analysis using simple mathematical models is self-consistent. The mean ratio for the total O:Mo ratio determined from high resolution analysis yielded the same result as found using the total intensity of O:Mo from the survey spectrum.

The chemical structure and crystalline state of the MoO$_x$ films were studied using the Renishaw inVia Raman spectrometer. A Renishaw frequency doubled NdYAG laser excitation source of wavelength 532 nm was used. To avoid local heating of the samples, a low power of about 5 mW was applied to the samples. A Raman shift between the wavenumber 200 to 1200 cm$^{-1}$ was measured. The transmittance of the MoO$_x$ films on the glass substrate was measured using the Cary 5000 UV-Vis-NIR spectrophotometer with a 150 mm integrating sphere. The measurements were performed in the wavelength range 300 to 2500 nm at a near-normal angle of incidence. The measured transmittance values were subtracted from the base (zero) signal. A Teflon coating was used as a 100% reference. From these measurements, the weighted solar transmittance for A.M. 1.5 and optical bandgap of the films were obtained.

C. Simulation of perovskite solar cell

The SCAPS simulator was used for the simulation of the perovskite solar cell device using MoO$_x$ as HTM. SCAPS software is developed at Electronics and Information Systems (ELIS), University of Gent, which is modelled under an AM 1.5 light spectrum.$^{25}$ It is very well-known simulation software in thin film solar cells as well as solar cells that have a planar structure.$^{26}$ In this work, a planar structure consisting of regular (FTO/TiO$_2$/CH$_3$NH$_3$PbI$_3$/PCBM/MoO$_x$/Au) and inverted (FTO/MoO$_x$/CH$_3$NH$_3$PbI$_3$/PCBM/Au) configuration were used for the simulation of the perovskite solar cell. Input parameters used for the simulation of the device were obtained from experimental results of this work and literature as shown in Table I. Here, $N_A$ and $N_D$
denote the acceptor and donor densities, \( \varepsilon_r \) is the relative permittivity, \( \chi \) is the electron affinity, \( E_g \) is the bandgap energy, \( \mu_n \) and \( \mu_p \) are mobility of electron and hole, and \( N_t \) is the defect density.\(^{27,28}\) The other parameters not mentioned in the Table are the effective density of charge at the conduction band (\( N_C \)) and at the valence band (\( N_V \)), which have values of \( 2 \times 10^{18} \) and \( 1.8 \times 10^{19} \), respectively.\(^{28}\)

III. RESULTS AND DISCUSSIONS

A. Structural properties

The Raman spectra of the MoO\(_x\) thin films deposited at different substrate temperatures is shown in Fig. 1. Three sharp Raman peaks obtained at 665, 820, and 995 cm\(^{-1}\) belong to the orthorhombic \( \alpha \)-phase of molybdenum oxide.\(^{35}\) Such a phase is normally obtained after high temperature annealing (450°C). The film deposited at room temperature has shown peaks of \( \alpha \)-phase with very weak Raman intensity which is the characteristics of amorphous films.\(^{36}\) The peak intensity, however, increases with increasing substrate temperature and this shows that the films deposited at higher substrate temperatures are dominated by the crystalline properties.\(^{36}\) The Raman band at 995 cm\(^{-1}\) is assigned to the terminal oxygen (Mo=O) stretching mode, whereas the 820 cm\(^{-1}\) to the doubly connected bridge-oxygen (Mo=O=Mo) stretching mode and the 665 cm\(^{-1}\) to the triply connected bridge-oxygen (Mo\(_3\)-O) vibration.\(^{37}\) The peak at 820 cm\(^{-1}\) has high sharpness for the films deposited at 100°C and 200°C which indicates that the corresponding vibrational modes are due to the highly ordered structure. The corner-sharing chains of MoO\(_6\) octahedra may be visualized in the \( \alpha \)-phase which shares the edge with two similar chains to form the MoO\(_3\) stoichiometry. Each octahedron of MoO\(_3\) has one unshared oxygen atom, two oxygen atoms are common to two octahedra, and three oxygens are in part-shared edge and common to the three octahedra,\(^{36}\) which referred as the \( \alpha \)-phase.\(^{38}\)

To observe the effect of substrate temperature on the chemical characteristics of MoO\(_x\) thin films, XPS analysis was conducted. Figure 2 shows a series of survey and high resolution (Mo 3d, O 1s and C 1s) spectra of the MoO\(_x\) thin films deposited at different substrate temperatures. The films produced at room temperature consist of pure MoO\(_3\), as they exhibit the oxygen to molybdenum ratio of 3:1. This implies that a pure stoichiometric film is deposited from the electron
beam evaporator at room temperature. High resolution scans of the Mo 3d and O 1s core level confirm this finding, where the Mo 3d doublet can be fitted with a single doublet pair with the expected 3:2 intensity ratio between the 3d_{5/2} and 3d_{3/2} states. By contrast, the films deposited at elevated temperatures (100°C and 200°C) show a significant reduction of Mo(V) state, with films deposited at 100°C containing 75% Mo (V) and films deposited at 200°C containing 76% Mo (V) as shown in Fig. 3. This analysis points to an overall oxygen to molybdenum ratio of 2.7:1 for the sample deposited at 200°C.

The high resolution Mo 3d and O 1s scans in Fig. 3 are fitted based on an assumed 5:2 and 3:1 oxygen weighting of the Mo (V) and Mo (VI) components, respectively. The spectral weight of oxygen was adjusted for the presence of C-Ox species in adventitious carbon, as well as ascribing high-binding energy spectral weight to adsorbed water. As explained by Bulpett et al., the conduction band of MoO3 consists of empty 4d and 5s states whereas the valence band consists of oxygen 2p state. At higher substrate temperature, the transition from oxygen 2p to an empty Mo^{6+} 4d level will give rise to the incorporation of lower valency Mo^{5+} ion in the lattice.

B. Morphological properties

Figure 4 shows the surface morphology of the MoOx thin films deposited at room temperature, 100°C and at 200°C characterized using HIM [Figs. 4(a)–4(c)] and AFM [Figs. 4(d)–4(f)]. Both films are homogenous with no cracks or pits and have smooth surface morphology covering the substrate uniformly as observed from the HIM micrographs in Figs. 4(a)–4(c). There is usually an increase in grain size with increasing deposition temperature, which is not obvious in the HIM and AFM images of this work. However, the surface roughness of the films slightly decreases with increasing deposition temperature as observed from the AFM images in Figs. 4(d)–4(f) and Table II. These surface roughness values were confirmed by the stylus profilometer and AFM.

C. Optical properties

Figure 5(a) shows the spectral transmittance of the 100 nm MoOx films deposited at different substrate temperatures. The films are highly transparent in the visible and near-infrared wavelength region and their transmittance sharply dropped in the ultraviolet wavelength. With increasing...
deposition temperature, the absorption edge shifts towards the longer wavelength. From the Raman spectra, the film deposited at room temperature has amorphous behaviour and this property can also be observed from the optical spectra in Fig. 5(a) having low interference fringes. The strong coloration of the films deposited at high temperature is due to oxygen ion vacancies, which promotes the reduction of oxygen atoms in the oxide structure which is confirmed from the XPS data.\textsuperscript{40} The weighted optical transmittance of all the films in the solar wavelength is more than 80% as shown in Table II with slightly higher value observed from the amorphous film. This amorphous film has slightly higher transmittance in the infrared compared to the crystalline MoO\textsubscript{x} films. The interference fringes would also have effect on the variation in the weighted solar transmittance.

The bandgap energy ($E_g$) of the MoO\textsubscript{x} films (100 nm) was calculated using the relation\textsuperscript{40}

$$ (a h\nu)^2 = A(h\nu - E_g), \quad (1) $$

where $a$ is the absorption coefficient, $A$ is the band edge parameter, $h$ is the Plank constant, and $\nu$ is the frequency of light. The plot produced from this relation is known as the Tauc plot and gives a linear behaviour in the high energy region. The $E_g$ of the films is then calculated by fitting the linear region of the plot to zero. Figure 5(b) shows the $(a h\nu)^2$ vs photon energy ($h\nu$). The estimated $E_g$ value of MoO\textsubscript{x} deposited at room temperature is 3.75 eV (see Table II). This value decreases to 3.45 eV when the film was deposited at 200°C. The decrease in bandgap energy is attributed to the oxygen vacancies which enable to capture electrons and act as a donor centres.\textsuperscript{41} The deficiency of oxygen is confirmed by the XPS result which shows that the atomic percentage of

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|c|}
\hline
Substrate temperature ($^\circ$C) & Transmittance (T\%) & O:Mo ratio & RMS surface roughness (nm) & Bandgap Energy $E_g$ (eV) \\
\hline
RT & 84 & 2.96 & 2.0 & 3.75 \\
100 & 82 & 2.76 & 1.59 & 3.50 \\
200 & 81 & 2.69 & 1.50 & 3.45 \\
\hline
\end{tabular}
\caption{Transmittance, atomic ratio of O:Mo, surface roughness, and bandgap energy of MoO$_x$ thin films deposited at different substrate temperatures.}
\end{table}

Fig. 4. Micrograph of MoO$_x$ thin films deposited at room temperature (a), (d), 100°C (b), (e), and 200°C (c), (f). (a), (b), (c) are HIM images having a scale bar of 1 $\mu$m and (d), (e), (f) are AFM images scanned over 2 $\mu$m x 2 $\mu$m.

Fig. 5. (a) Transmittance spectra and (b) $(a h\nu)^2$ vs $h\nu$ plot of 100 nm thick MoO$_x$ thin films deposited at various substrate temperatures.
oxygen is reduced with the increase in the substrate temperature (see Table II). The bandgap energy of MoOx obtained in this work (3.45–3.75 eV) are found to be within the values of e-beam deposited molybdenum oxide films reported in the literature (3.1–3.7 eV). These experimentally obtained Eg values of MoOx for a 100 nm thick film are used in the SCAPS simulation to predict the efficiency of the regular and inverted structures of the PSC. In the simulation, the MoOx film thickness was varied between 50 and 250 nm that included the experiment film thickness in order to observe the effect of film thickness on the various parameters of the solar cells.

D. Simulation of perovskite solar cell with MoOx as HTM

Empirical simulation analysis using the SCAPS simulation software was performed to study the performance of perovskite solar cell with MoOx as an HTM layer. The schematic diagram of the inverted and regular structures used in the simulation is shown in Figs. 6(a) and 6(b), respectively. A typical perovskite solar cell configuration adopts a solid planar heterojunction p-i-n structure where the perovskite material in the regular structure is sandwiched between Electron Transport Layer (ETL) (TiO2) and HTM (MoOx). In the regular structure, the HTM is applied on the top of the perovskite material; thus, deposition of MoOx at only room temperature is suitable as the perovskite material is sensitive to elevated temperature (more than 100 °C). In the inverted structure, the HTM (MoOx) is applied on the top of fluorine doped tin oxide (FTO) and PCBM is typically used as ETL and allows high substrate temperature deposition of the MoOx thin films. FTO coated glass is used as a transparent conductive oxide, and Au serves as a back metal contact. Thus, using the experimentally obtained bandgap energy (3.75 eV) and film thickness (100 nm) of MoOx, the solar cell efficiency for both the regular and inverted structures are found to be 18.21% to 17.85%, respectively.

An optimum thickness of the HTM with full surface coverage is very important for the device performance. Figure 7 shows the effect of MoOx thickness on the open circuit voltage (Voc), short-circuit current density (Jsc), recombination current density (Jrec), fill factor (FF), and efficiency for both the regular [Fig. 7(a)] and inverted [Fig. 7(b)] structures. As observed in the figure, the Jsc slightly decreases with the increase in HTM (MoOx) thickness whereas the Voc remains unchanged. This decrease in Jsc can be linked to the increase in recombination current as shown in the Fig. 7(a). The total recombination current density (Jrec) increases with the increase of HTM thickness which causes a reduction in Jsc and overall efficiency of the solar cell. The same trend is followed in both regular and inverted structures. This increase in recombination current density is due to longer distance that photo-generated carriers should transport to be driven out, whereas their life-time and mobility are kept constant. So a minimum of MoOx layer thickness that gives full layer coverage to minimize recombination in the HTM layer is desirable. Also, in the inverted structure [Fig. 7(b)] where the light enters through the MoOx side, the FF decreased with the increase of MoOx thickness due the increase in recombination current density and an increase in light absorption. From the HIM images in Fig. 4, the 100 nm thick MoOx films deposited at different substrate temperatures have fully covered the substrate. By varying the MoOx layer thickness from 50 to 250 nm for the highest bandgap energy of MoOx obtained in Table II, the efficiency of the PSC only slightly decreases from 18.25% to 18.15% for the regular structure [Fig. 7(a)] and from 17.88% to 17.70% for the

![FIG. 6. Schematic diagram of (a) inverted structure and (b) regular structure of a perovskite solar cell used in SCAPS simulation.](image)

![FIG. 7. Open circuit voltage (Voc), short-circuit current density(Jsc), current recombination (Jrec), fill factor (FF), and efficiency of PSC as a function of MoOx layer thickness in (a) regular and (b) inverted PSC structure. The room temperature deposited bandgap energy (3.75 eV) of MoOx films is considered.](image)
The best cells with 50 nm thicknesses have shown high $V_{oc}$ and $J_{sc}$ values of 1.03 V and 22.3 mA/cm$^2$ for the regular structure, and 0.97 V and 22.7 mA/cm$^2$ for the inverted structure with a fill factor of 78.7 and 82.5, respectively.

The quantum efficiency (QE) is one of the important parameters that describe the quality of light absorption, charge transfer, and collection of a solar cell. As shown in Fig. 8, the QE for the (a) regular and (b) inverted structures ($E_g = 3.75$ eV) decreases with the increase of HTM thickness in the wavelength region below 400 nm and this is more prominent in the regular structure of the solar cell. The decrease in QE at the lower wavelength can be due to the recombination in the MoO$_x$ layer and/or the light absorption of MoO$_x$ layer. The carriers which are generated by the high energy photons (>bandgap energy of MoO$_x$) do not contribute to the quantum efficiency because they cannot reach the back contact of the inverted structure PSC due to their limited life time and mobility leading to enhanced recombination in the MoO$_x$ layer. Smaller bandgap of MoO$_x$ means the photons at wavelength less than 400 nm can be absorbed by the MoO$_x$ layer, which in turn reduces the effective light absorption of the perovskite light absorber and thus the QE. The higher the MoO$_x$ thickness, the higher is the light absorption by the HTM of the inverted structure and the lower the QE.

As discussed before, the experimental results have confirmed that the increase in substrate temperature causes an increase in the crystallinity, a decrease in the bandgap energy along with the overall oxygen to molybdenum ratio and surface roughness of the MoO$_x$ thin film. Thus, the effect of MoO$_x$ bandgap energy on the PSC performance for the inverted structure has been investigated using SCAPS simulation. As shown in Fig. 9(a) for the 100 nm layer of MoO$_x$, the $J_{sc}$ increases from 22.3 mA/cm$^2$ to 22.6 mA/cm$^2$, whereas the $V_{oc}$ remains constant with increasing MoO$_x$ bandgap. The recombination current density decreases with the increase of bandgap energy. The increase in $J_{sc}$ governs an overall increase in the efficiency of the simulated PSC device to 17.85% as shown in Fig. 9(a). From the above analysis,
deposition of MoO\textsubscript{x} films at higher substrate temperature causes a reduction in the bandgap energy which reduces the simulated \( J_{sc} \). Variation in QE with different MoO\textsubscript{x} bandgap energies (100 nm layer of MoO\textsubscript{x}) is shown in Fig. 9(b). An increase in QE with the increase of the MoO\textsubscript{x} bandgap energy is observed at the lower wavelength region. However, with increasing deposition temperature, an increase in both crystallinity and defect density of the film and a reduction in surface roughness were observed. These properties which are useful for enhancing the performance of the solar cell were not considering in the SCAPS simulation.

As shown in the simulated results, the variation in bandgap energy of MoO\textsubscript{x} as HTM only slightly affects the efficiency of the PSC. However, as observed in the experiment, deposition of MoO\textsubscript{x} thin films at higher substrate temperature (e.g., 200 °C) gives rise to oxygen deficiency and enhanced crystallinity of the films which would be expected to improve the performance and stability of the PSC.\textsuperscript{43} From the calculation, a thinner MoO\textsubscript{x} film of about 50 nm thick that can completely cover the surface of the substrate is found to be beneficial for enhanced QE. Using these conditions, the performance of the PSC was determined as shown in Fig. 10 with optimum efficiency of 17.82% and high QE.

IV. CONCLUSION

SCAPS simulation software was used to determine MoO\textsubscript{x} as an alternative HTM in perovskite solar cells. The HTM layer thickness and bandgap energy obtained experimentally were used to study the device performance. Empirical simulation results indicate that a larger HTM thickness gives a lower efficiency due to higher recombination current (\( J_{rec} \)), whereas the bandgap energy has a slight effect on the efficiency of the PSC. Using the properties of electron beam evaporated MoO\textsubscript{x} film deposited at room temperature, the modelling shows an efficiency of 18.25% for the regular structure and 17.88% for the inverted structure. The MoO\textsubscript{x} deposited at higher substrate temperature (200 °C) is suitable for the inverted structure of PSC with promising efficiency of 17.82%. The overall simulated results suggest that the MoO\textsubscript{x} thin film is a potential hole transport inorganic material in the perovskite solar cell and can replace the organic hole transport materials with improved performance, stability, and lower cost. Experimental results of the MoO\textsubscript{x} films indicated a reduction in bandgap energy, oxygen composition, and surface roughness but enhanced crystallinity with increasing substrate temperature. Device fabrication using MoO\textsubscript{x} as HTM is needed to examine the effect of these parameters with the change of substrate temperature on the performance of perovskite solar cells.

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FIG. 10. Quantum efficiency (QE) of PSC using MoO\textsubscript{x} as HTM in the inverted structure of PSC. The inset shows the optimized efficiency of the PSC in the inverted structure using a film thickness of 50 nm.