

Cite this: *RSC Adv.*, 2017, 7, 7890

A simple synthesis method to prepare a molybdenum oxide hole-transporting layer for efficient polymer solar cells

Yanping Li,^{ab} Huangzhong Yu,^{*a} Xinxin Huang,^{ab} Zuping Wu^{ab} and Mingdong Chen^a

We report a simple synthetic method to prepare amorphous molybdenum oxide (p-MoO₃) using a favorably stable peroxomolybdic acid organosol as the precursor solution prepared by an ultrasonic reaction for the first time. The favorably smooth and dense surface morphology of the p-MoO₃ layers are obtained under 150 °C thermal treatment with good optical properties and a high work function (W_F) of 5.26 eV. During the annealing treatment two different oxidation states of Mo ions are observed with increasing the annealing temperature to 150 °C and 200 °C. The best performance of the P3HT:PC₇₁BM devices with p-MoO₃ anode buffer layers has been achieved under 150 °C treatment with a power conversion efficiency (PCE) of 4.02%, a V_{OC} of 0.59 V, a J_{SC} of 10.70 mA cm⁻², and a FF of 63.7%, superior to the corresponding PEDOT:PSS modified devices. Furthermore, the performance of the PTB7:PC₇₁BM devices with the annealed p-MoO₃ buffer layers has also been dramatically improved with the best performance parameters of a PCE of 8.46%, a V_{OC} of 0.73, a J_{SC} of 17.02 mA cm⁻², a FF of 68.1% for 150 °C. The improved performance of the devices originates from the following factors; (i) the favorable and compact surface morphology of the annealed p-MoO₃ films leading to a higher rectification ratio and lower leakage current. (ii) The formation of oxygen vacancies and the growing Mo⁵⁺ cation leading to the change of W_F under the annealing treatment. The highest W_F of 5.26 eV for 150 °C treatment influences the built-in electric field of the devices with the photocurrent being extracted efficiently at a short-circuit.

Received 9th January 2017
Accepted 20th January 2017

DOI: 10.1039/c7ra00303j

www.rsc.org/advances

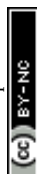
1. Introduction

In recent years the development of bulk heterojunction (BHJ) polymer solar cells (PSCs) continues to show progress,^{1–7} and PSCs have also been a hot research topic due to their advantages of low-cost manufacturing, light weight and good flexibility.^{8–11} A BHJ polymer solar cell generally consists of a transparent anode (typically indium tin oxide, ITO), a BHJ active layer (a mixture of donor polymer and fullerene acceptor), and a cathode (*e.g.* Al, Ca/Al or LiF/Al). Anode/cathode interfacial layers are usually used as charge selective contacts between the BHJ active layer and the electrodes. The development of appropriate electron (or hole) charge blocking and opposite transporting charge layers is paramount to the fabrication of more efficient solution processed PSC devices. The role of such layers is also to create asymmetrical interfaces to enable reduced recombination of photogenerated electrons and holes at the electrodes and collect charge effectively.¹²

Nowadays the aqueous poly(3,4-ethylenedioxythiophene)-poly(styrene sulfonate) (PEDOT:PSS) is typically used as the anode buffer layer in PSCs. However, PEDOT:PSS has the hygroscopic and acidic nature with an associated reduction in device stability.^{13,14} Therefore, developing a straightforward method to prepare more moderate, stable and effective anode interlayers is highly desirable for the low-cost, high throughput manufacturing of efficient PSC modules. In this regard, transition metal oxide semiconductors including NiO,^{15,16} MoO₃,^{17,18} V₂O₅ (ref. 19) and WO₃ (ref. 20) are emerging as a promising candidates of substitutes for PEDOT:PSS as the hole-transporting layer (HTL), which possess the demonstrable advantages of the wide range of energy level aligning capabilities, high transmittance in the visible region, and excellent ambient stability. Moreover, the developed solution-processed metal oxides semiconductors are compatible with low-cost all-solution high-throughput manufacturing of PSCs. Among these semiconductors, solution-processed MoO₃ thin films have been prepared from different precursors such as ammonium molybdate,^{21–23} MoO₂(acac)₂,^{24,25} Mo(CO)₃(EtCN)₃ (ref. 26) and dispersed MoO₃ nanoparticles^{27,28} and proved to be a promising candidate as efficient hole extraction layers. However, many weak points can be discerned in the above solution-based synthetic methods for MoO₃ films, which may ultimately decrease their hole-

^aSchool of Physics and Optoelectronics, South China University of Technology, Guangzhou 510640, China. E-mail: hzhyu@scut.edu.cn; Tel: +86-20-87110768

^bSchool of Materials Science & Engineering, South China University of Technology, Guangzhou 510640, China



transporting efficiency when they become part of an organic optoelectronic devices. For instance, the inadequately homogeneous surface of MoO₃ films that are usually formed, and the high temperature of thermal treatment along with additional process steps (such as oxygen plasma treatment) are generally needed after spin-coating the precursor solution. In addition, the sol-gel method involving dissolution of MoO₃ in hydrogen peroxide, and then dispersion with polyethylene glycol and 2-methoxyethanol, resulting in smooth MoO₃ films after the time-consuming reflux and high temperature (275 °C) thermal treatment.^{29,30}

In this paper, we report a simply synthetic method to prepare the favorably stable dark blue peroxomolybdic acid organosol by the ultrasonic reaction for the first time. Then the molybdenum oxide (marked as p-MoO₃) films with varying reduction degrees are prepared using the peroxomolybdic acid organosol as the precursor solution by spin-coating method. The optical characteristic, surface topography and electrical characteristics of p-MoO₃ films annealed in a broad temperature range are compared and discussed in depth. In order to validate the effect of our prepared p-MoO₃ films annealed under various temperature on the OPV devices, the OPV devices with the structures of ITO/HTL/P3HT (or PTB7):PC₇₁BM/LiF/Al are fabricated. Finally, the mechanism of the efficiency enhancement upon incorporation of p-MoO₃ films as anode buffer layers under various annealing temperatures is proposed.

2. Experimental section

Regioregular electron-donor poly(3-hexylthiophene) (P3HT), poly[[4,8-bis[(2-ethylhexyl)oxy]benzo[1,2-*b*:4,5-*b'*]dithiophene-2,6-diyl][3-fluoro-2-(2-ethylhexyl) carbonyl]thieno[3,4-*b*]thiophenediyl]] (PTB7) and acceptor [6,6]-phenyl C₇₁-butyric acid methyl ester (PC₇₁BM) were purchased from Luminescent Technology, Inc. The PEDOT:PSS (Clevios 4083) were purchased from H.C. Starck. The chlorobenzene, 1,2-dichlorobenzene and MoO₃ (99.98%) were purchased from Sigma-Aldrich. All the chemicals were used as received without further purification. The structures of the fabricated PSC devices are ITO/HTL/P3HT (or PTB7):PC₇₁BM/LiF/Al. The structures of the PSC devices, chemical structures and energy level diagram of materials used in the PSC devices are shown in Fig. 1.

2.1 Preparation of the molybdenum solution

The synthesis was based on the preparation of the peroxomolybdic acid solution with the treatment of the ultrasonic reaction, which was very crucial for the preparation of the peroxomolybdic acid solution. First, ethanediol and isopropanol were mixed to obtain 40 mL complex-solvent by the volume ratio of 1 : 9 (ethanediol to isopropanol), then 400 mg MoO₃ powder reacted with 40 mL complex-solvent, and the 30% aqueous H₂O₂ was added to the reacted system by the volume ratio of 7 : 93 (H₂O₂ to complex-solvent). The system was dissolved completely after ultrasonication for 30 min, forming the dark blue peroxomolybdic acid organosol (inserted in Fig. 2). The molybdenum concentration was 10 mg mL⁻¹ in mother liquor

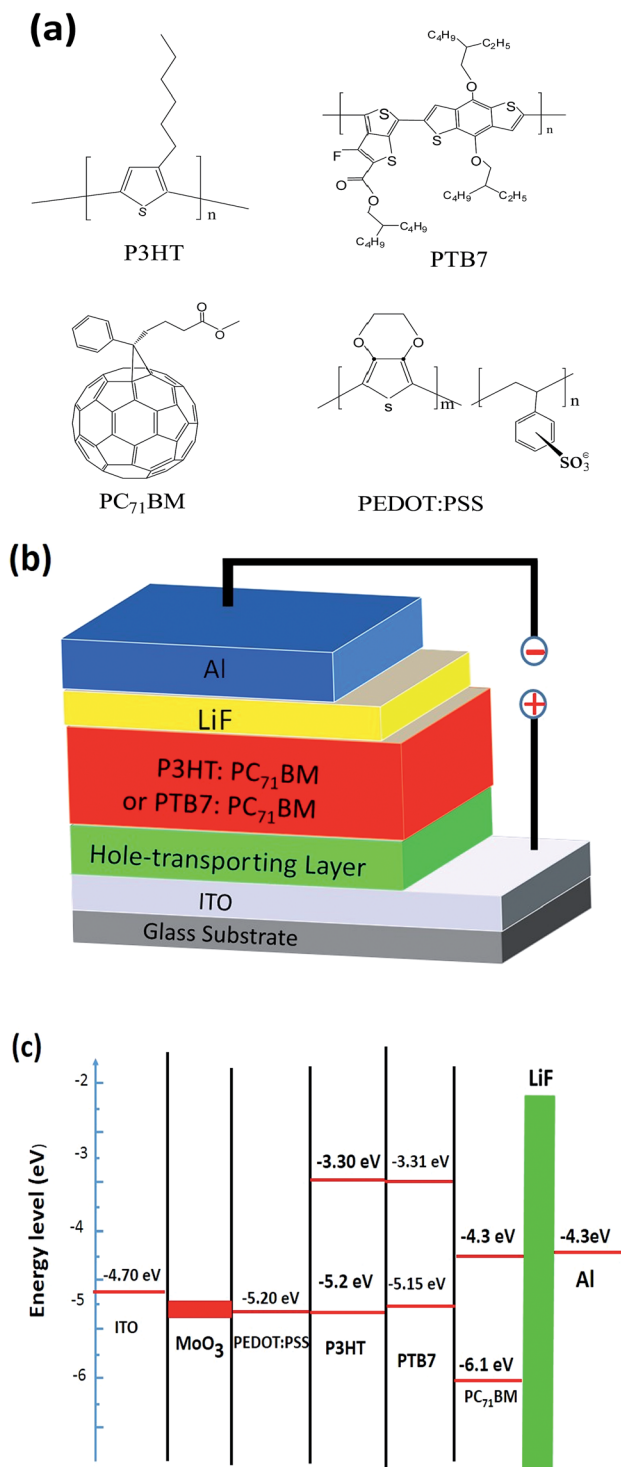


Fig. 1 (a) The molecular structures of P3HT, PTB7, PC₇₁BM, p-MoO₃ and PEDOT:PSS. (b) The typical device structure of P3HT:PC₇₁BM or PTB7:PC₇₁BM polymer solar cell. (c) Energy level diagram of the component materials used.^{2,31}

and had good dispersion and stability for several months. For each experimental run, the peroxomolybdic acid organosol was diluted with isopropanol to a certain molybdenum concentration.



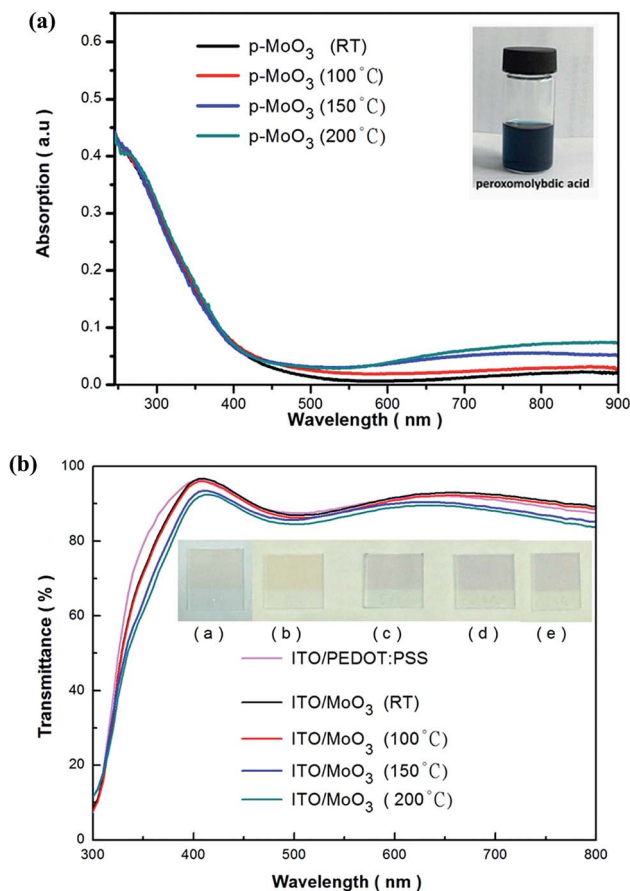


Fig. 2 (a) UV-vis absorption spectra of the p-MoO₃ thin films under various annealing temperature. (b) Transmission spectra of PEDOT:PSS thin film (a) and p-MoO₃ thin films treated with room temperature drying (b), 100 °C (c), 150 °C (d), 200 °C (e).

2.2 Fabrication of PSC devices

Prior to device fabrication, the solutions of light-absorbing layer were prepared. Both 1,2-dichlorobenzene solution containing P3HT and PC₇₁BM (1 : 0.8, w/w) and a mixed solvent of chlorobenzene and 1,8-diiodooctane (97 : 3, v/v) containing PTB7 : PC₇₁BM (1 : 1.5, w/w) were stirred in glovebox at 60 °C overnight. Upon cooling to room temperature, the solutions were filtered through a 0.2 µm polytetrafluoroethylene (PTFE) filter before using. The ITO substrates (CSG Holding Co. Ltd., Shenzhen, China) were cleaned by sonication using detergent, deionized water, acetone and isopropanol sequentially for 15 min. After an oven drying for a few hours, the ITO surface was treated with plasma treatment for 10 min. Then the layers of PEDOT:PSS (or peroxomolybdic acid organosol) were spin-coated onto the cleaned ITO to obtain the PEDOT:PSS films (or p-MoO₃ films) as an anode buffer layers at 4000 rpm for 40 s and baked at a certain annealing temperature in glovebox, and the thickness of PEDOT:PSS (or MoO₃) was about 40 nm (or 10 nm). The molybdenum concentration was diluted with isopropanol to 2 mg mL⁻¹ using mother liquor to regulate the solution viscosity. And the annealing time to construct device HTL in this study was for 20 min. Next, the active layers were

spin-coated in the glovebox (O₂ < 0.1 ppm, H₂O < 0.1 ppm) by employing the active layer solutions. The LiF and Al electrode were deposited continuously in a deposition chamber under a basic vacuum pressure lower than 3×10^{-4} Pa. The shadow mask was used during thermal evaporation to define the active area of 0.15 cm². All devices were unencapsulated and measured in air.

2.3 Characterization of thin film and PSC devices

All thickness measurements were analysed using alpha-step surface profiler. Structural assessments of the thin film samples under different annealing temperature *via* XRD was performed using a Bruker Discovery 08 with a Hi-Star area detector.

The UV-vis absorption and transmittance spectra of the p-MoO₃ thin films under various annealing temperature and PEDOT:PSS thin film were measured with a HP 8453 UV-vis spectrophotometer.

XPS analysis was carried out on an XPS/ESCA (electron spectroscopy for chemical analysis) instrument (Axis Ultra DLD from Kratos Inc.) utilizing a monochromatized Al Kα X-ray source. Spectra were referenced to the residual hydrocarbon contamination at 284.6 ± 0.1 eV. The thin films (*ca.* 40 nm) were deposited on the ITO substrate for XPS analysis.

The work functions of the p-MoO₃ thin film samples under different annealing temperatures were investigated using Kelvin Probe measurements, carried out on a KP Technology SKP5050 in a glove-box filled in a conditioned chamber where the O₂ level is <0.1 ppm.

Scanning electron microscope (SEM) and Tapping-mode atomic force microscope (AFM) images using a NanoScope NS3A system (Digital Instruments) were obtained to observe the surface morphology of the annealed films.

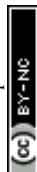
The current density-voltage (*J-V*) characteristics of the devices were performed using a computer controlled Keithley 2400 source-meter under illumination with calibrated AM1.5G (100 mW cm⁻²) at room temperature.

3. Result and discussion

3.1 Optical characteristics of MoO₃ thin films

Firstly the UV-vis measurement was carried on the spin-coated thin films with the same spin-coating conditions and different annealing temperatures. As clearly shown in Fig. 2(a), all the p-MoO₃ thin films are almost transparent within visible light range, and this will be of benefit for light harvesting. However, a growing small broad absorption ranging from 600 nm to 800 nm is observed for the annealed thin films under 150 °C. Increasing the treated temperature to 200 °C, this small broad absorption becomes more evident. This fascinating phenomenon may be attributed to the free electrons being trapped in oxygen vacancies in these p-MoO₃ thin films.³²

In accordance with the condition of device fabrication, we further investigated the transmission spectra of the thin films including PEDOT:PSS film to evaluate the transparency of these films in devices (Fig. 2(b)). These films are spin-coated on ITO



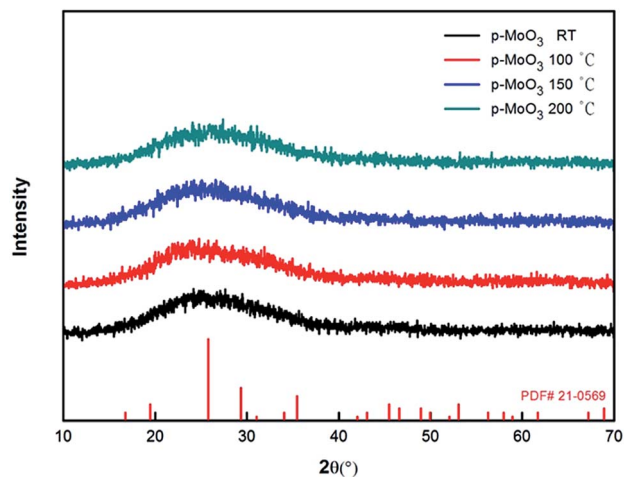


Fig. 3 XRD pattern of the resulting p-MoO₃ thin films annealed under various temperatures.

glasses. It is obvious that transmission values of all the films show good light transmittance except in the wavelength near 500 nm and 750 nm. This phenomenon is attributed to the transparency of ITO glass and small broad absorption mentioned above.^{33,34} Nevertheless, the transmission values are above 80% for all films regardless of the annealing temperature, implying that p-MoO₃ thin films are qualified to be good interfacial layer without significant absorption losses.

3.2 Morphology of the p-MoO₃ thin film

Then XRD measurements were performed on p-MoO₃ thin films deposited on ITO substrates and annealed in glovebox with different annealing temperatures. Fig. 3 shows the XRD patterns of p-MoO₃ thin films that are formed at various heat treated temperatures, the p-MoO₃ layers do not yield significant diffraction peaks as confirmed by XRD as shown in Fig. 3, which shows that these p-MoO₃ thin films have an amorphous structure. But these p-MoO₃ thin films with amorphous structure have been demonstrated to be effective to the device performance.^{24,25,35}

To understand the surface morphology of the annealed films, SEM and AFM were performed to observe the morphology characteristics of the p-MoO₃ films. Fig. 4 shows SEM images of the annealed p-MoO₃ thin films with different annealing temperatures. It is clear that the surface under the condition of room temperature drying is quite uneven and not uniform with some obvious small MoO₃ grains. However, it can also be noted that the surface become much flatter with the increase of heating temperature. We attribute this phenomenon to the evaporation of the solvent and progress of p-MoO₃ thin film formation. When the annealing temperature is increased to 150 °C, surface topography of the p-MoO₃ thin film is the most flat and uniform without any small grains. At this point, the heat treatment makes the film more compact, which prevents the current leakage of the solar cells, and is beneficial to the improvement of the devices. But further raising the annealing temperature to 200 °C, thin film appears critically gathered

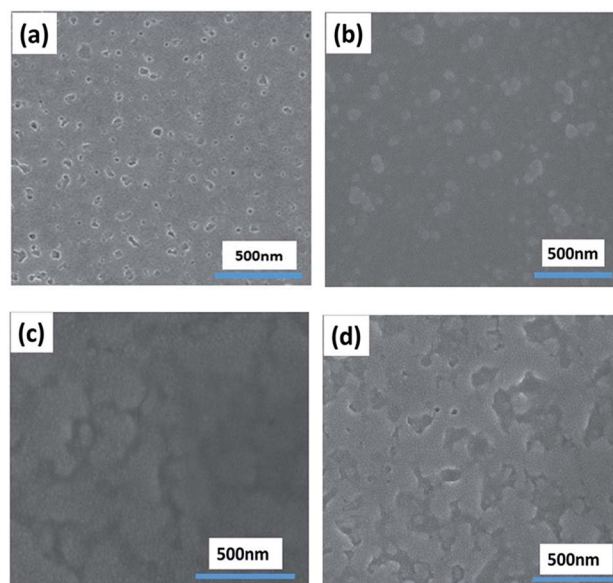


Fig. 4 SEM images of the thin films morphology of p-MoO₃ thin films morphology on ITO treated with room temperature drying (a), 100 °C (b), 150 °C (c), 200 °C (d).

aggregation phenomenon and less uniform. This can be due to enhanced surface diffusion of the species leading to smaller grains joining together to grow into larger grains.

Fig. 5 shows AFM images of the p-MoO₃ thin films with different annealing temperatures. The surface roughness (in Root Mean Square, RMS) of the p-MoO₃ thin film treated with different annealing temperatures is 2.36 nm, 2.12 nm, 1.46 nm and 1.51 nm for room temperature drying, 100 °C, 150 °C and 200 °C, respectively. The p-MoO₃ thin films treated with heat treatment of 150 °C possess the lowest surface roughness, which is quite well consistent with the result of SEM images. Low surface roughness of the p-MoO₃ thin film suggests that they can smoothen the ITO substrate surface effectively to avoid local shorts and at the same time be dense enough to prevent the current leakage, which is prerequisites to guarantee their implementation as effective buffer layer for organic electronics.

3.3 Components and electrical characterization of p-MoO₃ thin films

X-ray photoelectron spectroscopy (XPS) shown in Fig. 6 was operated to obtain the evidence for the composition of the p-MoO₃ thin films treated with various annealing temperature. The binding energy (BE) obtained in the XPS analysis was corrected for specimen charge by referencing the C 1s peak to 284.6 eV. Fig. 6(a) compares the full scan spectra of the annealed p-MoO₃ thin films, the characteristic peaks of the O 1s, Mo 3s, Mo 3p and Mo 3d levels are shown in the full spectra of the p-MoO₃ thin films. Moreover, no other peaks of the impurity and ITO substitute are found, which suggests that the synthetic precursor can be spread uniformly and completely decomposes to p-MoO₃ by annealing. In Fig. 6(b), the Mo 3d doublet presents a spin-orbit splitting between the Mo 3d_{3/2}



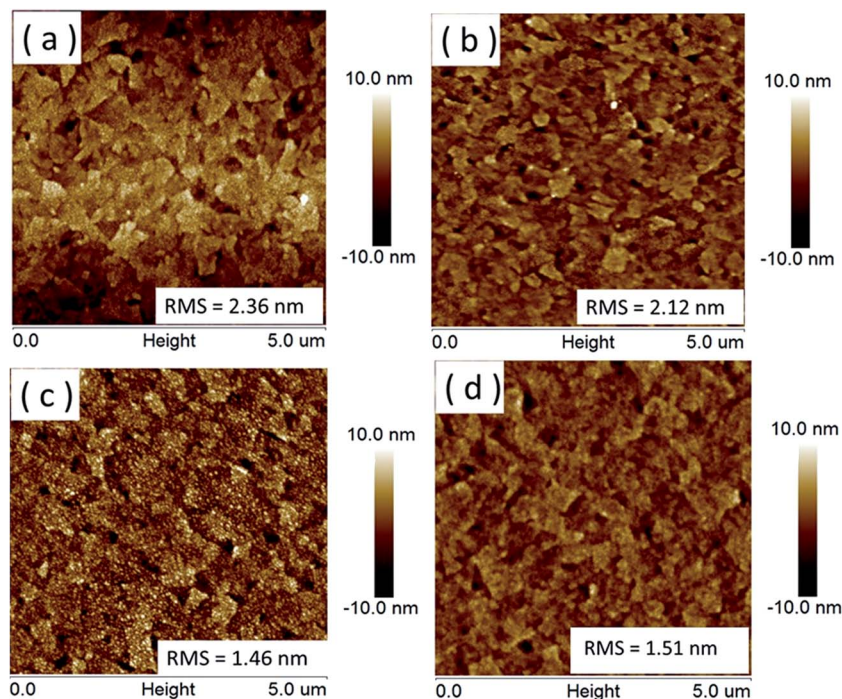


Fig. 5 AFM morphology images of p-MoO₃ thin films on ITO treated with room temperature drying (a), 100 °C (b), 150 °C (c), 200 °C (d).

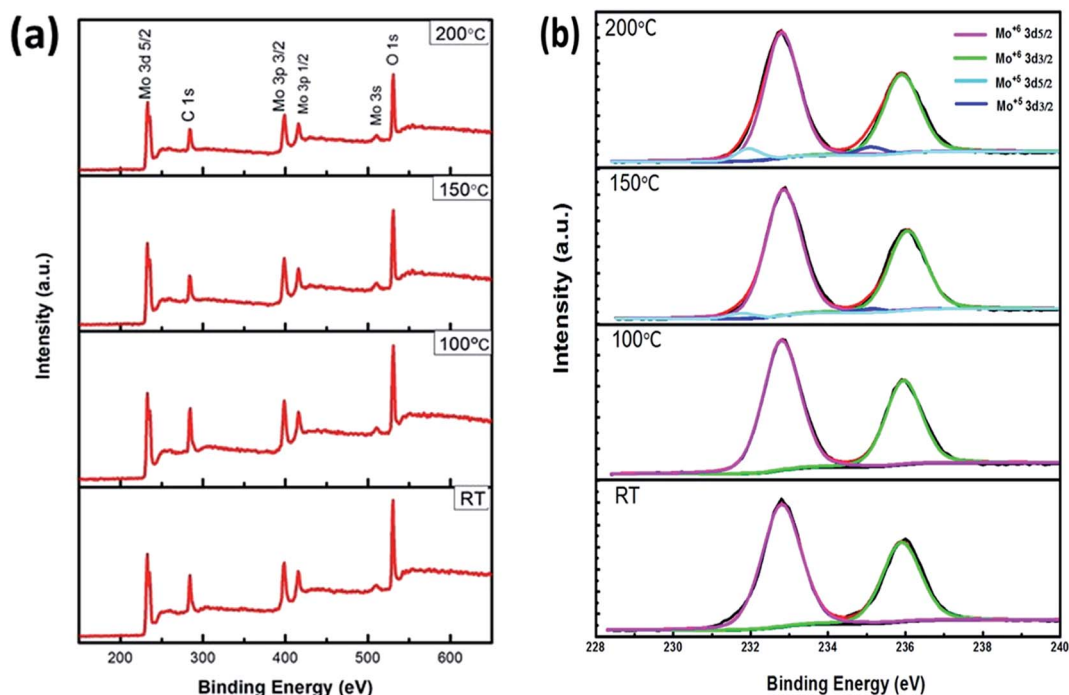
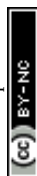


Fig. 6 X-ray photoelectron spectroscopy of p-MoO₃ thin films. (a) Full scanned spectra, (b) Mo 3d core level of thin films treated with various annealing temperature.

and 3d5/2 bands. The Mo 3d3/2 and 3d5/2 peaks of the p-MoO₃ films treated under RT and 100 °C located at 232.8 and 235.9 eV, respectively. The lines of doublet are Gaussian with the d intensity ratio of 2 : 3, separated by 3.1 eV. It is apparent that

the position of the Mo (3d) lines is expected for Mo⁶⁺ ions as in p-MoO₃.^{30,36,37}

However, two different oxidation states of Mo ions are observed with further increasing the annealing temperature to



150 °C and 200 °C, the positions of newly formed Mo⁵⁺ peaks in the 150 °C and 200 °C annealed films fitted by two 3d5/2 and 3d3/2 doublets in the form of a Gaussian function located 231.7 eV and 234.9 eV, respectively, which is identified as Mo⁵⁺.^{38,39} While the positions of Mo⁶⁺ peaks annealed films has not changed with further improve the annealing temperature. The calculated Mo⁵⁺/Mo⁶⁺ ratios for the 150 °C and 200 °C annealed films are 0.030 and 0.071, respectively. By using the Mo⁵⁺/Mo⁶⁺ obtained from the Mo 3d core level spectra, the estimated Mo:O stoichiometries of p-MoO₃ films annealed in the 150 °C and 200 °C are 2.99 and 2.96, respectively. It can be ascribed that the concentration of the Mo⁵⁺ component in the p-MoO₃ films is greatly connected with thin film processing conditions, which coincides with the fact that preparation of p-MoO₃ in the oxygen deficient environment (in glovebox) and annealing temperatures led to present Mo⁵⁺ states, corresponding to the electron transfer from O 2p to Mo 4d and generation of oxygen vacancies.^{39–42}

To further evaluate the energetics of the solution-processed p-MoO₃ thin films under annealing conditions, Kelvin probe (KP) analysis was performed to obtain the work function (W_F) of annealed films in glovebox referring to literature.²⁶ Kelvin probe analysis of the PEDOT:PSS coated on ITO as a reference gives a W_F of 5.20 eV, which is in agreement with the literature.² The W_F of annealed p-MoO₃ thin films shows a gradual shift towards the higher values from 4.90 eV to 5.26 eV with increasing annealing temperatures from RT to 150 °C, reaching the highest W_F value of 5.26 eV with 150 °C. We attribute this phenomenon to the desorption of adsorbates and evaporation of the solvent.^{43,44} With further increasing the annealing temperature to 200 °C, the W_F value of the film decreases slightly to 5.22 eV. The main reason may be linked with the distinctly reduced cation oxidation shown in Fig. 6(b) and occupied oxygen vacancy defect states within the band that played the part of donor levels. The W_F values are shown in Table 1.

3.4 Effect of the p-MoO₃ thin film on the performance of PSCs

To investigate the effect of the p-MoO₃ buffer layer on the photovoltaic performance of PSCs, firstly we selected the P3HT:PC₇₁BM blended composite as the light-absorbing layer and constructed some solar cells. Device structures of P3HT:PC₇₁BM solar cells fabricated with p-MoO₃ thin films treated with various annealing temperatures are presented as follow:

Table 1 W_F values of anode interface coated with PEDOT:PSS and MoO₃ by Kelvin probe evolution

Anode interface	T (°C)	W_F (eV)
PEDOT:PSS	150	5.20
p-MoO ₃	RT	4.90
p-MoO ₃	100	5.10
p-MoO ₃	150	5.26
p-MoO ₃	200	5.22

- (A) ITO/PEDOT:PSS (150 °C)/P3HT:PC₇₁BM/LiF/Al.
- (B) ITO/p-MoO₃ (RT)/P3HT:PC₇₁BM/LiF/Al.
- (C) ITO/p-MoO₃ (100 °C)/P3HT:PC₇₁BM/LiF/Al.
- (D) ITO/p-MoO₃/P3HT:PC₇₁BM/LiF/Al.
- (E) ITO/p-MoO₃ (200 °C)/P3HT:PC₇₁BM/LiF/Al.

Devices A is the control sample with a PEDOT:PSS buffer layer, devices B, C, D and E are samples with the annealed p-MoO₃ layer. The device performance parameters are summarized in Table 2.

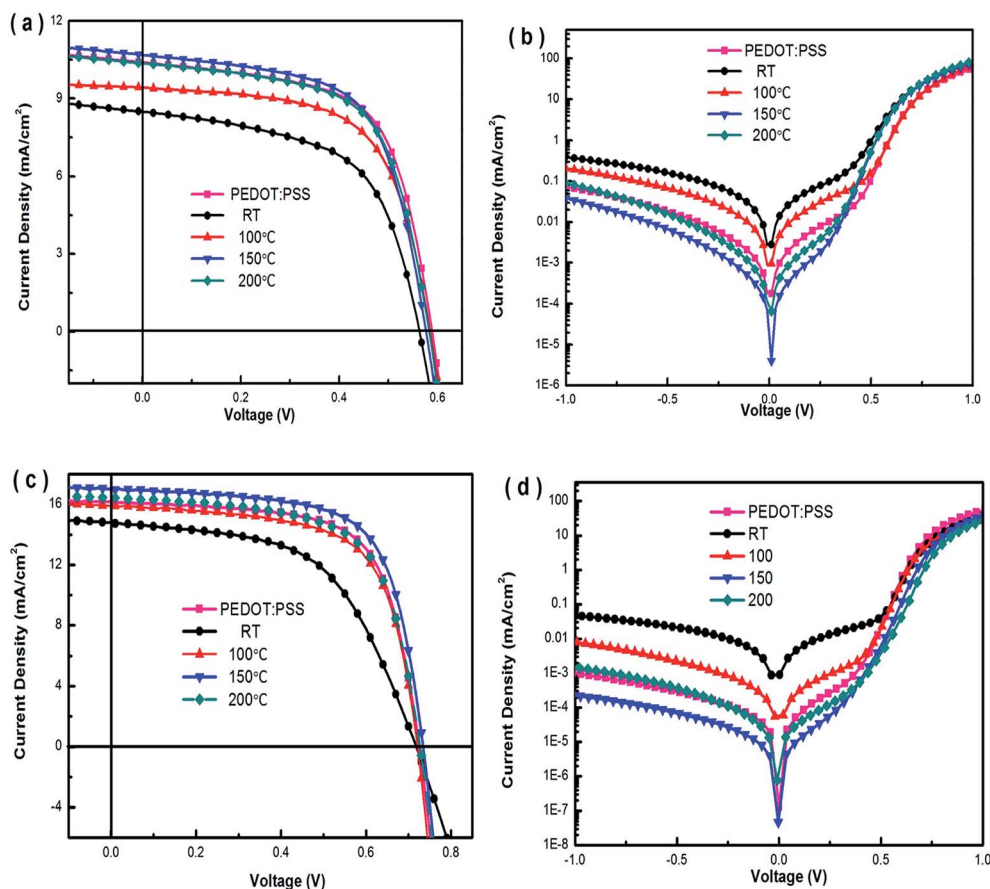
J - V curves of devices A–E under the illumination of AM1.5G, 100 mW cm^{−2} are depicted in Fig. 7(a). For device A with PEDOT:PSS buffer layer, it shows a PCE of 3.91% with a short-circuit current density (J_{SC}) of 10.50 mA cm^{−2}, an open-circuit voltage (V_{OC}) of 0.59 V, and a fill factor (FF) of 63.1%. In contrast, the four parameters (V_{OC} , J_{SC} , FF, and PCE) of devices with the p-MoO₃ buffer layer exhibit different degrees of growth under the annealing temperature increased from RT to 150 °C. The device with the p-MoO₃ anode buffer layer annealed at 150 °C shows the best PCE of 4.02, with a V_{OC} of 0.59 V, a J_{SC} of 10.70 mA cm^{−2}, and a FF of 63.7%, which is better than that using the PEDOT:PSS buffer layer (PCE = 3.91%). Further increasing the annealing temperature to 200 °C, the device performance is decreased apparently with a PCE of 3.78%, a V_{OC} of 0.58 V, a J_{SC} of 10.36 mA cm^{−2}, and a FF of 62.9%.

A closer look at the J - V curves are shown in Fig. 7(b), we can find that the both reverse current density and leakage current of the devices are reduced with the increase of heat treatment p-MoO₃ buffer layer, and the devices for 150 °C possess the lowest reverse current density and leakage current. This is because the 150 °C heat treatment makes the film more compact, which has fewer defects, reduces charge recombination, and is beneficial to the improvement of the devices. Moreover, the rectification ratio of the devices calculated based on the J - V curves at ± 1.0 V is also calculated. For devices A, B, C, D and E, the rectification ratio is 2.2×10^3 , 2.0×10^2 , 5.1×10^2 , 3.1×10^3 and 1.1×10^3 , respectively. Clearly the rectification ratio values of devices D and E with the p-MoO₃ buffer layer annealed under 150 °C and 200 °C are simultaneously higher than that of device B and C with the p-MoO₃ buffer layer under temperature drying and annealed under 100 °C. Series resistance R_s and shunt resistance R_{sh} are clear as shown in Table 2. R_{sh} of P3HT:PC₇₁BM based devices with p-MoO₃ buffer layer annealed at 150 °C is 1145.46 Ω cm², which is much greater than the devices with p-MoO₃ buffer layer without annealing (639.03 Ω cm²), annealed at 100 °C (743.96 Ω cm²) and at 200 °C (921.46 Ω cm²). This is presumably due to the increase of compact density of the film surface (see Fig. 4(c) and 5(c)), leading to the decrease of charge recombination and leakage current, which is also in agreement with the Fig. 7(b). Moreover, the R_s of the devices decrease with the increase of heat treatment p-MoO₃ buffer layer, and the value of 8.08 Ω cm² for the device with p-MoO₃ layer annealed 150 °C is the lowest among those devices with p-MoO₃ layer other heat treatment temperature. Increase of heat treatment temperature usually causes the reduction of molybdenum oxide, which leads to rise in conductivity of molybdenum oxide. In general, reduced transition metal oxides (TMOs) not only shows an increase in conductivity, but also causes stronger



Table 2 Performance parameters of P3HT:PC₇₁BM solar cells fabricated with annealed p-MoO₃ thin films

Device	<i>T</i> (°C)	<i>J</i> _{SC} (mA cm ⁻²)	<i>V</i> _{OC} (V)	FF (%)	<i>R</i> _s (Ω cm ²)	<i>R</i> _{sh} (Ω cm ²)	PCE (%)
A	150	10.50	0.59	63.1	8.15	968.57	3.91
B	RT	8.49	0.57	56.9	10.50	639.03	2.75
C	100	9.42	0.58	62.0	8.49	743.94	3.38
D	150	10.70	0.59	63.7	8.08	1145.46	4.02
E	200	10.36	0.58	62.9	8.20	921.46	3.78

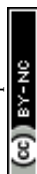
Fig. 7 *J*-*V* characteristic curve of different devices fabricated with annealed p-MoO₃ thin films under illumination and in the dark. (a, b) P3HT:PC₇₁BM solar cells under illumination and in the dark, (c, d) PTB7:PC₇₁BM solar cells under illumination and in the dark.

optical absorption.⁴⁵ We attribute this interesting phenomenon to a result of the forming process of p-MoO₃ and a slightly non-stoichiometric composition with some oxygen deficiency leading to reduced oxides.

The improved PCE of devices with annealed p-MoO₃ hole transporting layers (HTL) may origin from the following factors; (i) the favorable surface morphology of the annealed p-MoO₃ HTL. As shown in SEM images (Fig. 4) and AFM images (Fig. 5), the p-MoO₃ HTL spin-coated on ITO substrates without the annealing is quite not smooth and dense compared to substrates, leading to the poor device performance. By increasing the annealing temperature, the p-MoO₃ surface is growing smoother and compacter with the optimum surface and smallest roughness (RMS) of 1.46 nm for 150 °C. This

decreased roughness accounts for the different dark *J*-*V* characteristics (Fig. 7(b)) of the devices with p-MoO₃ layer annealed various temperatures, in which the devices exhibit increasingly excellent diode characteristics with higher rectification ratio and lower leakage current under reverse bias with improved annealing temperature. It means that the annealed p-MoO₃ layer is capable of lowering the leakage current caused by the disagreeable roughness of ITO substrate. Further improving the annealing temperature leads to the slight aggregation and rising back of roughness. Therefore, the contact between the p-MoO₃ layer and the active layer becomes less favorable to the hole extraction.

(ii) The formation of oxygen vacancies and the growing Mo⁵⁺ cation leading to the change of *W*_F under the annealing. UV-vis



absorption spectra analysis (Fig. 2(a)) indicates the formation of oxygen vacancies of the annealed p-MoO₃ layer started with the annealing temperature 150 °C and the amount of oxygen vacancies further increased with improved temperature to 200 °C, which is all consistent with the XPS analysis as presented in Fig. 6. During annealing treatment, small amount of oxygen vacancies generated owing to the process of dehydration and desorption is capable of causing effective n-type doping for 150 °C, which is greatly conducive to the improved performance of the devices D with a PCE of 4.02%, the further increasing amount of oxygen vacancies with improved temperature to 200 °C leads to a slightly declined PCE of 3.78%. Furthermore, the oxygen ion vacancies of the substoichiometric p-MoO₃ are occupied by two electrons and acted as donor centres with the donor level at 0.27 eV below the conduction band, the charge transfer from oxygen vacancies to Mo ions creates reduced oxidation states of Mo⁵⁺. It is reported that a general correlation between cation oxidation state and an oxide's W_F is that metal oxides in their more oxidized forms tend to have higher work functions than their reduced forms.⁴³ Therefore, the increasing change of the W_F from 4.90 eV to 5.22 eV under the annealing from RT to 200 °C couples with a fluctuation of the highest W_F 5.26 eV for 150 °C, which influences the built-in electric field of the devices and the photocurrent being extracted efficiently at short-circuit.

In order to further testify the possibility of general usage of the p-MoO₃ layer as a hole-transporting layer under the mild thermal-annealing in polymer solar cells, we also prepare the devices with the structures of PTB7:PC₇₁BM solar cells fabricated with PEDOT:PSS and p-MoO₃ thin films treated with various annealing temperatures as following:

(F) ITO/PEDOT:PSS (150 °C)/PTB7:PC₇₁BM/LiF/Al.

(G) ITO/p-MoO₃ (RT)/PTB7:PC₇₁BM/LiF/Al.

(H) ITO/p-MoO₃ (100 °C)/PTB7:PC₇₁BM/LiF/Al.

(I) ITO/p-MoO₃ (150 °C)/PTB7:PC₇₁BM/LiF/Al.

(J) ITO/p-MoO₃ (200 °C)/PTB7:PC₇₁BM/LiF/Al.

In Fig. 7(c), the J - V characteristics of the PTB7:PC₇₁BM devices under AM1.5G illumination are presented. It is evident that the annealing to p-MoO₃ layer induces large improvement in J_{SC} from 14.78 mA cm⁻² to 16.43 mA cm⁻² and FF from 54.9% to 65.1% with the annealing temperature from RT to 200 °C. The best J_{SC} , FF and PCE of the p-MoO₃-based devices is 17.02 mA cm⁻², 68.1% and 8.46% for 150 °C, respectively, which apparently surpass the PEDOT-based devices with a J_{SC} of 16.17 mA cm⁻², FF of 67.2% and PCE of 7.93%. The performance parameters are summarized in Table 3.

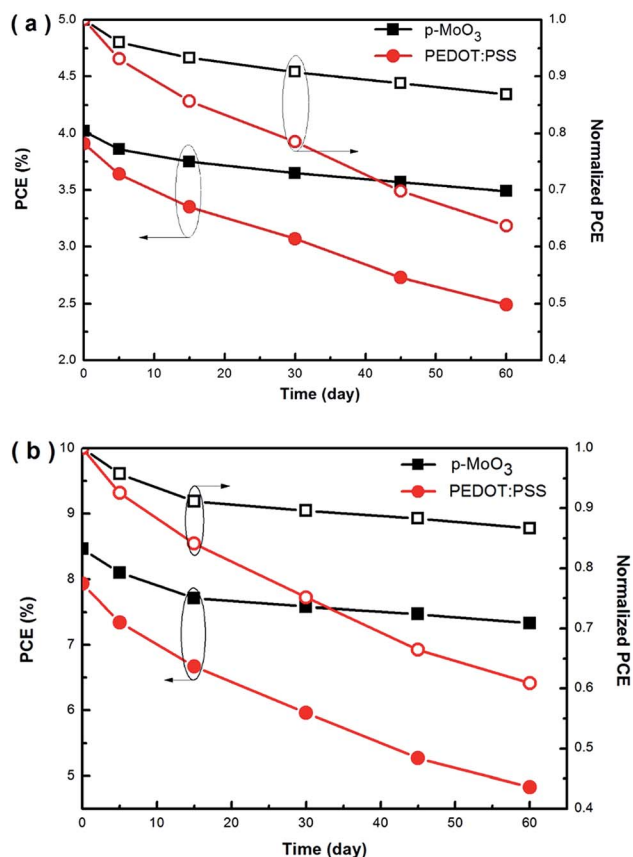


Fig. 8 PCEs and normalized PCEs as functions of aging time of P3HT:PC₇₁BM (a) and PTB7:PC₇₁BM (b) devices incorporating p-MoO₃ thin films.

As a consequence, the devices with the p-MoO₃ films annealed at 150 °C exhibit the largest increase in performance with the best PCE of 8.46%, further confirming that our prepared p-MoO₃ thin films annealed at moderate temperature are fit for the PTB7:PC₇₁BM devices due to the favorable surface morphology and electrical properties as discussed above. Meanwhile, according to the leakage dominated region (<0.5 V) as shown in Fig. 7(d), the dark current of the PTB7:PC₇₁BM devices with the annealed p-MoO₃ thin films present the trend toward miniaturization with the minimum at 150 °C, which is much less than that of the device F with PEDOT:PSS. This also indicates that p-MoO₃ thin film is more efficient in hole-collection and electron-blocking when served as a HTL in PTB7:PC₇₁BM BHJ devices.

Table 3 Performance parameters of PTB7:PC₇₁BM solar cells fabricated with annealed p-MoO₃ thin films

Device	T (°C)	J_{SC} (mA cm ⁻²)	V_{OC} (V)	FF (%)	R_s (Ω cm ²)	R_{sh} (Ω cm ²)	PCE (%)
F	150	16.17	0.73	67.2	5.05	1174.06	7.93
G	RT	14.78	0.72	54.9	13.35	705.19	5.85
H	100	15.90	0.72	66.2	5.21	827.53	7.57
I	150	17.02	0.73	68.1	4.49	1275.09	8.46
J	200	16.43	0.73	65.1	4.82	976.58	7.81



Apart from device efficiency, device stability is another important aspect affecting the realization of commercial polymer solar. In this study, we find that the devices with p-MoO₃ thin film exhibit enhanced stability. All of these devices constructed as described in Section 2.2 are not encapsulated and stored in the glovebox filled with nitrogen atmosphere (O₂ < 0.1 ppm, H₂O < 0.1 ppm). We choose the devices with p-MoO₃ and PEDOT:PSS thin films annealed at optimal annealing temperature 150 °C as tested objects. Fig. 8 displays the PCEs and normalized PCEs as a function of aging time, of P3HT:PC₇₁BM and PTB7:PC₇₁BM devices incorporated p-MoO₃ thin films. For P3HT:PC₇₁BM devices as shown in Fig. 8(a), the PCEs decrease dramatically for the devices fabricated with PEDOT:PSS and reach from 3.91% to 2.40% after 60 days, only keeping about 60% of their initial efficiencies. In comparison, the p-MoO₃-based P3HT:PC₇₁BM devices get PCEs of 3.40% after storage for 60 days, remaining about 85% of their initial efficiencies (from 4.02% to 3.40%). On the other hand, Fig. 8(b) presents the stability of PTB7:PC₇₁BM devices with different HTL. The PEDOT-based devices degrade rapidly after 60 days with about 40% decay of PCEs (from 7.93% to 4.83%), whereas the devices incorporating the p-MoO₃ HTLs get only 13.3% decay of PCEs (from 8.46% to 7.33%) after 60 days, revealing the high stability of the p-MoO₃ interfaces. Clearly, it is indicated that the devices incorporating the p-MoO₃ thin films exhibit higher stability.

4. Conclusions

The favorably stable dark blue peroxomolybdic acid organosol is simply synthesized by the treatment of the ultrasonic reaction. The prepared p-MoO₃ films were characterized from the spectroscopic measurements, SEM, AFM, XPS analysis, and Kelvin probe analysis. It is found that the favorably smooth and dense surface morphology of the p-MoO₃ layers are obtained under 150 °C with the good optical and electrical properties. Furthermore, the performance of the P3HT:PC₇₁BM and PTB7:PC₇₁BM devices with p-MoO₃ thin films is found to depend strongly on the annealing and show best performance for 150 °C. The improved performance of devices origin from the factors; (i) the favorable surface morphology of the annealed p-MoO₃ films leading to higher rectification ratio and lower leakage current. (ii) The formation of oxygen vacancies and the growing Mo⁵⁺ cation leading to the change of W_F under the annealing. Furthermore, the devices containing p-MoO₃ thin film exhibited enhanced environmental stability. Overall, the simple and stable peroxomolybdic acid solution that we have developed is a viable candidate for replacing the PEDOT:PSS anode modifying layers in many organic solar cells including other electronic devices that require an ultrasoft and high W_F solution-processed hole selective contact.

Acknowledgements

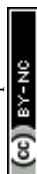
This work was financially supported by the National Natural Science Foundation of China (Grant No. 61474046 and 61176061).

References

- 1 G. Yu, J. Hummelen, F. Wudl and A. J. Heeger, Polymer photovoltaic cells: Enhanced efficiencies *via* a network of internal donor-acceptor heterojunctions, *Science*, 1995, **270**, 1789–1791.
- 2 Z. He, C. Zhong, S. Su, M. Xu, H. Wu and Y. Cao, Enhanced power-conversion efficiency in polymer solar cells using an inverted device structure, *Nat. Photonics*, 2012, **6**, 591–595.
- 3 J. You, L. Dou, K. Yoshimura, T. Kato, K. Ohya, T. Moriarty, K. Emery, C. Chen, J. Gao, G. Li and Y. Yang, A polymer tandem solar cell with 10.6% power conversion efficiency, *Nat. Commun.*, 2013, **4**, 1446.
- 4 Y. Liu, J. Zhao, Z. Li, C. Mu, W. Ma, H. Hu, K. Jiang, H. Lin, H. Ade and H. Yan, Aggregation and morphology control enables multiple cases of high-efficiency polymer solar cells, *Nat. Commun.*, 2014, **5**, 5293.
- 5 Z. He, B. Xiao, F. Liu, H. Wu, Y. Yang, S. Xiao, C. Wang, T. P. Russell and Y. Cao, Single-junction polymer solar cells with high efficiency and photovoltage, *Nat. Photonics*, 2015, **9**, 174–179.
- 6 H. Zhou, Y. Zhang, C. Mai, S. D. Collins, G. C. Bazan, T. Nguyen and A. J. Heeger, Polymer Homo-Tandem Solar Cells with Best Efficiency of 11.3%, *Adv. Mater.*, 2015, **27**, 1767–1773.
- 7 A. R. M. Yusoff, D. Kim, H. P. Kim, F. K. Shneider, W. J. Silvab and J. Janga, A high efficiency solution processed polymer inverted triple-junction solar cell exhibiting a power conversion efficiency of 11.83%, *Energy Environ. Sci.*, 2015, **8**, 303–316.
- 8 J. J. Zhao, B. Z. Xia, K. Lu, D. Deng, L. Yuan, J. Q. Zhang, L. Y. Zhu, X. W. Zhu, H. Li and Z. X. Wei, Dialkoxyphenyldithiophene-based small molecules with enhanced absorption for solution processed organic solar cells, *RSC Adv.*, 2016, **6**, 60595–60601.
- 9 M. C. Scharber and N. S. Sariciftci, Efficiency of bulk-heterojunction organic solar cells, *Prog. Polym. Sci.*, 2013, **38**, 1929–1940.
- 10 X. Guo, N. Zhou, S. J. Lou, J. Smith, D. B. Tice, J. W. Hennek, R. P. Ortiz, J. T. L. Navarrete, S. Li, J. Strzalka, L. X. Chen, R. P. H. Chang, A. Facchetti and T. J. Marks, Polymer solar cells with enhanced fill factors, *Nat. Photonics*, 2013, **7**, 825–833.
- 11 A. J. Heeger, 25th anniversary article: bulk heterojunction solar cells: understanding the mechanism of operation, *Adv. Mater.*, 2014, **26**, 10–28.
- 12 R. Steim, F. R. Kogler and C. J. Brabec, Interface materials for organic solar cells, *J. Mater. Chem.*, 2010, **20**, 2499–2512.
- 13 J. J. Lee, S. H. Lee, F. S. Kim and J. H. Kim, Simultaneous enhancement of the efficiency and stability of organic solar cells using PEDOT:PSS grafted with a PEGME buffer layer, *Org. Electron.*, 2015, **26**, 191–199.
- 14 Y. Sun, J. H. Seo, C. J. Takacs, J. Seifter and A. J. Heeger, Inverted polymer solar cells integrated with a low-temperature-annealed sol-gel-derived ZnO film as an electron transport layer, *Adv. Mater.*, 2011, **23**, 1679–1683.



- 15 L. Zhang, L. Favereau, Y. Farre, A. Maufroy, Y. Pellegrin, E. Blart, M. Hissler, D. Jacquemin, F. Odobel and L. Hammarstrom, Molecular-structure control of electron transfer dynamics of push-pull porphyrins as sensitizers for NiO based dye sensitized solar cells, *RSC Adv.*, 2016, **6**, 77184–77194.
- 16 J. Zhang, J. Wang, Y. Fu, B. Zhanga and Z. Xie, Efficient and stable polymer solar cells with annealing-free solution-processible NiO nanoparticles as anode buffer layers, *J. Mater. Chem. C*, 2014, **2**, 8295–8302.
- 17 D. Zhao, S. T. Tan, L. Ke and D. L. Kwong, Optimization of an inverted organic solar cell, *Sol. Energy Mater. Sol. Cells*, 2010, **94**, 985–991.
- 18 Y. Sun, C. J. Takacs, S. R. Cowan, J. H. Seo, X. Gong, A. Roy and A. J. Heeger, Efficient, Air-Stable Bulk Heterojunction Polymer Solar Cells Using MoO_x as the Anode Interfacial Layer, *Adv. Mater.*, 2011, **23**, 2226–2230.
- 19 K. Shen, R. Yang, D. Wang and K. Ho, Stable CdTe solar cell with V₂O₅ as a back contact buffer layer, *Sol. Energy Mater. Sol. Cells*, 2016, **144**, 500–508.
- 20 T. Stubhan, N. Li, N. A. Luechinger, S. C. Halim, G. J. Matt and C. J. Brabec, High fill factor polymer solar cells incorporating a low temperature solution processed WO₃ hole extraction layer, *Adv. Energy Mater.*, 2012, **2**, 1433–1438.
- 21 X. B. Chu, M. Guan, Y. Zhang, Y. Y. Li, X. F. Liu and Y. P. Zeng, ITO-free and air stable organic light-emitting diodes using MoO₃:PTCDA modified Al as semitransparent anode, *RSC Adv.*, 2013, **3**, 9509–9513.
- 22 K. H. Wong, K. Ananthanarayanan, J. Luther and P. Balaya, Origin of hole selectivity and the role of defects in low-temperature solution-processed molybdenum oxide interfacial layer for organic solar cells, *J. Phys. Chem. C*, 2012, **116**, 16346–16351.
- 23 S. Murase and Y. Yang, Solution processed MoO₃ interfacial layer for organic photovoltaics prepared by a facile synthesis method, *Adv. Mater.*, 2012, **24**, 2459–2462.
- 24 K. Zilberberg, H. Gharbi, A. Behrendt, S. Trost and T. Riedl, Low-temperature, solution-processed MoO_x for efficient and stable organic solar cells, *ACS Appl. Mater. Interfaces*, 2012, **4**, 1164–1168.
- 25 Z. Tan, D. Qian, W. Zhang, L. Li, Y. Ding, Q. Xu, F. Wang and Y. Li, Efficient and stable polymer solar cells with solution-processed molybdenum oxide interfacial layer, *J. Mater. Chem. A*, 2013, **1**, 657–664.
- 26 S. R. Hammond, J. Meyer, N. E. Widjonarko, P. F. Ndione, A. K. Sigdel, A. Garcia, A. Miedaner, M. T. Lloyd, A. Kahn, D. S. Ginley, J. J. Berry and D. C. Olson, Low-temperature, solution-processed molybdenum oxide hole-collection layer for organic photovoltaics, *J. Mater. Chem.*, 2012, **22**, 3249–3254.
- 27 J. Meyer, R. Khalandovsky and P. Görrn, MoO₃ Films Spin-Coated from a Nanoparticle Suspension for Efficient Hole-Injection in Organic Electronics, *Adv. Mater.*, 2011, **23**, 70–73.
- 28 M. Xu, L. Cui, X. Zhu and L. S. Liao, Aqueous solution-processed MoO₃ as an effective interfacial layer in polymer/fullerene based organic solar cells, *Org. Electron.*, 2013, **14**, 657–664.
- 29 S. Lin, C. Wang, K. Kao, Y. Chen and C. Liu, Electrochromic properties of MoO₃ thin films derived by a sol-gel process, *J. Sol-Gel Sci. Technol.*, 2010, **53**, 51–58.
- 30 C. Girotto, E. Voroshazi, D. Cheyns, P. Heremans and B. P. Rand, Solution-processed MoO₃ thin films as a hole-injection layer for organic solar cells, *ACS Appl. Mater. Interfaces*, 2011, **3**, 3244–3247.
- 31 F. Cheng, G. Fang, X. Fan and Y. Li, Enhancing the performance of P3HT:ICBA based polymer solar cells using LiF as electron collecting buffer layer and UV-ozone treated MoO₃ as hole collecting buffer layer, *Sol. Energy Mater. Sol. Cells*, 2013, **110**, 63–68.
- 32 T. Yang, M. Wang, Y. Cao, F. Huang, L. Huang, J. Peng, X. Gong, S. Z. D. Cheng and Y. Cao, Polymer Solar Cells with a Low-Temperature-Annealed Sol-Gel-Derived MoO_x Film as a Hole Extraction Layer, *Adv. Energy Mater.*, 2012, **2**, 523–527.
- 33 R. A. Afre, Y. Hayashi and T. Soga, Enhanced photovoltaic device performance upon modification of indium tin oxide coated glass by liquid nitrogen treatment, *J. Phys. D: Appl. Phys.*, 2009, **42**, 042002.
- 34 L. Cattin, F. Dahou, Y. Lare, M. Morsli, R. Tricot, S. Houari, A. Mokrani, K. Jondo, A. Khelil, K. Napo and J. C. Bernède, MoO₃ surface passivation of the transparent anode in organic solar cells using ultrathin films, *J. Appl. Phys.*, 2009, **105**, 034507.
- 35 X. Yu, T. J. Marks and A. Facchetti, Metal oxides for optoelectronic applications, *Nat. Mater.*, 2016, **15**, 383–396.
- 36 F. Xie, W. C. H. Choy, C. Wang, X. Li, S. Zhang and J. Hou, Low Temperature Solution Processed Hydrogen Molybdenum and Vanadium Bronzes for an Efficient Hole Transport Layer in Organic Electronics, *Adv. Mater.*, 2013, **25**, 2051–2055.
- 37 S. Kato, R. Ishikawa, Y. Kubo, H. Shirai and K. Ueno, Efficient Organic Photovoltaic Cells Using Hole-Transporting MoO₃ Buffer Layers Converted from Solution-Processed MoS₂ Films, *Jpn. J. Appl. Phys.*, 2011, **50**, 071604.
- 38 M. A. Moraes, B. C. Trasferetti, F. P. Rouxinol, R. Landers, S. F. Durrant, J. Scarmínio and A. Urbano, Molybdenum oxide thin films obtained by the hot-filament metal oxide deposition technique, *Chem. Mater.*, 2004, **16**, 513–520.
- 39 M. Anwar, C. A. Hogarth and R. Bulpett, Effect of substrate temperature and film thickness on the surface structure of some thin amorphous films of MoO₃ studied by X-ray photoelectron spectroscopy (ESCA), *J. Mater. Sci.*, 1989, **24**, 3087–3090.
- 40 J. J. Jasieniak, J. Seifter, J. Jo, T. Mates and A. J. Heeger, A Solution-Processed MoO_x Anode Interlayer for Use within Organic Photovoltaic Devices, *Adv. Funct. Mater.*, 2012, **22**, 2594–2605.
- 41 Y. Lee, J. Yi, G. F. Gao, H. Koerner, K. Park, J. Wang, K. Luo, R. A. Vaia and J. W. P. Hsu, Low-Temperature Solution-Processed Molybdenum Oxide Nanoparticle Hole Transport Layers for Organic Photovoltaic Devices, *Adv. Energy Mater.*, 2012, **2**, 1193–1197.



- 42 K. Kanaia, K. Koizumib, S. Ouchic, Y. Tsukamotoc, K. Sakanouec, Y. Ouchib and K. Sekib, Electronic structure of anode interface with molybdenum oxide buffer layer, *Org. Electron.*, 2010, **11**, 188–194.
- 43 M. T. Greiner, L. Chai, M. G. Helander, W. Tang and Z. Lu, Transition Metal Oxide Work Functions: The Influence of Cation Oxidation State and Oxygen Vacancies, *Adv. Funct. Mater.*, 2012, **22**, 4557–4568.
- 44 Irfan, H. Ding, Y. Gao, C. Small, D. Y. Kim, J. Subbiah and F. So, Energy level evolution of air and oxygen exposed molybdenum trioxide films, *Appl. Phys. Lett.*, 2010, **96**, 243307.
- 45 J. Meyer, S. Hamwi, M. Kröger, W. Kowalsky, T. Riedl and A. Kahn, Transition metal oxides for organic electronics: energetics, device physics and applications, *Adv. Mater.*, 2012, **24**, 5408–5427.

