

# Nonfullerene Self-Assembled Monolayers As Electron-Selective Contacts for *n-i-p* Perovskite Solar Cells

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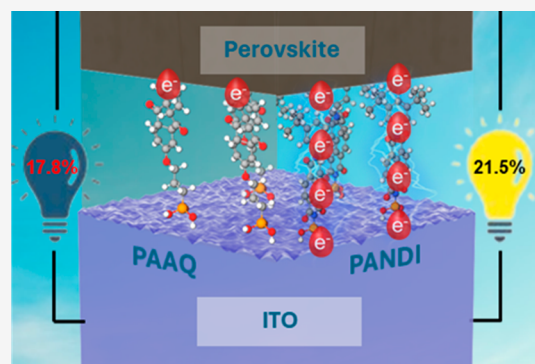


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**ABSTRACT:** Organic, nonfullerene semiconductors capable of self-assembly and composed of either anthraquinone (AQ) or naphthalenediimide (NDI) central fragments have been designed as electron-selective materials for *n-i-p* perovskite solar cells (PSCs). Both types of self-assembled monolayer (SAM) molecules contain phosphonic acid as an anchoring group, allowing covalent binding with indium tin oxide (ITO) surfaces. In particular, the NDI-based SAMs showed a more homogeneous anchoring on the ITO substrate and a stronger band bending at the ITO-SAM/perovskite interface than AQ-based SAMs. As a result, low-temperature-processed *n-i-p* PSCs with NDI SAMs as an electron-selective bottom contact showed a maximum power conversion efficiency (PCE) of 21.5%, representing the highest PCE among *n-i-p* PSCs with organic electron transporting layers (ETLs). In addition, our NDI-SAM-based devices demonstrate substantially improved long-term stability under operating temperature conditions when compared to devices using SnO<sub>2</sub> as the ETL.



Self-assembled monolayers (SAMs) have recently emerged as a viable replacement for traditional charge transport layers in perovskite solar cells (PSCs).<sup>1–6</sup> In particular, SAM molecules based on carbazole and phosphonic acid groups, usually referred to as *n*PACz, where *n* is the aliphatic chain length, have attracted great interest as hole-selective contacts.<sup>1,2,7–10</sup> The carbazole group has a strong electron-donating moiety and thus hole-selective properties when in contact with a semiconductor, while the phosphonic acid group can covalently bond on transparent conducting oxide (TCO) surfaces to form a monolayer.<sup>1,2,4,9,10</sup> Thanks to their fast hole extraction and the good electronic passivation of the SAMs/perovskite interface, *p-i-n* single-junction perovskite and monolithic perovskite/silicon tandem solar cells have achieved impressive certified power conversion efficiencies (PCEs) already above 26% and 33%, respectively.<sup>3,10–13</sup> While new SAMs are being intensely investigated toward efficient hole collection for a range of perovskite photoabsorbers with different bandgaps, reports toward the design of novel electron-selective SAM molecules remain rare.<sup>2,14,15</sup> Arguably, this is due to the relative scarcity of suitable electron acceptors that are available for the synthesis of electron-selecting materials, and the required efforts toward identification and application of nonfullerene derivatives as electron-selective

materials in PSCs.<sup>16,17</sup> Generally, the class of organic electron transport layers (ETLs) is dominated by fullerene derivatives, including C<sub>60</sub> and its functionalized analogues (e.g., PC<sub>61</sub>BM, PC<sub>71</sub>BM, ICBA, etc.).<sup>16–20</sup> Nonetheless, the application of fullerene-based ETLs for solution processing is limited by the poor solubility of unmodified fullerenes in organic solvents. Moreover, synthesizing and purifying modified fullerenes, particularly C<sub>70</sub> derivatives, are challenging, necessitating complex and costly multistage processes.<sup>16,18,21</sup>

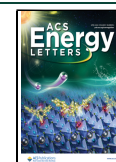
As a nonfullerene alternative, naphthalenediimide (NDI)-based small molecules offer favorable ETL properties, such as a high electron affinity, electron mobility, and photochemical stability, coupled with simple synthesis procedures, which motivates the design of new electron-selective NDI SAMs.<sup>22–24</sup> Initially, NDI SAMs such as *N,N'*-bis(1-*n*-hexylpyridinium-4-ylmethyl)-1,4,5,8-naphthalenetetracarboxy-

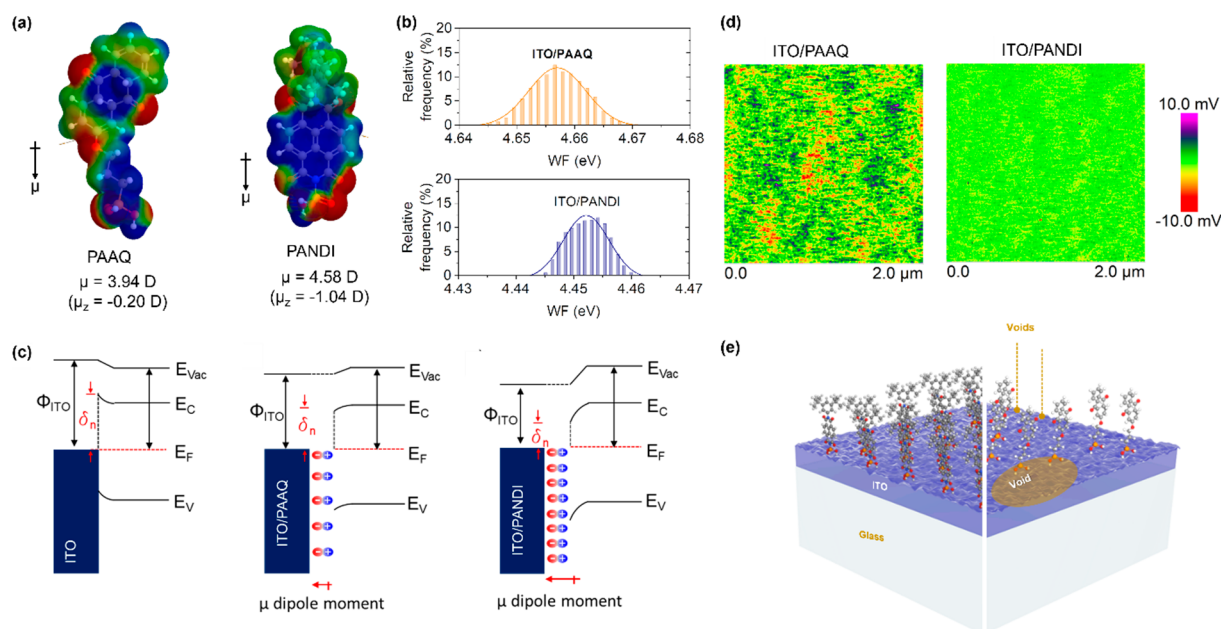
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**Figure 1.** (a) Electrical dipole moment of PAAQ and PANDI single molecules. (b) WF histogram of ITO/PAAQ and ITO/PANDI structures. (c) Band matching of pristine ITO/perovskite, ITO/PAAQ/perovskite, and ITO/PANDI/perovskite junctions, respectively. (d) Flattened KPFM mapping images of ITO/PAAQ and ITO/PANDI films. The scale of the image ranges was normalized from  $-10$  to  $10$  mV. (e) Illustration of SAM homogeneity on the ITO surface.

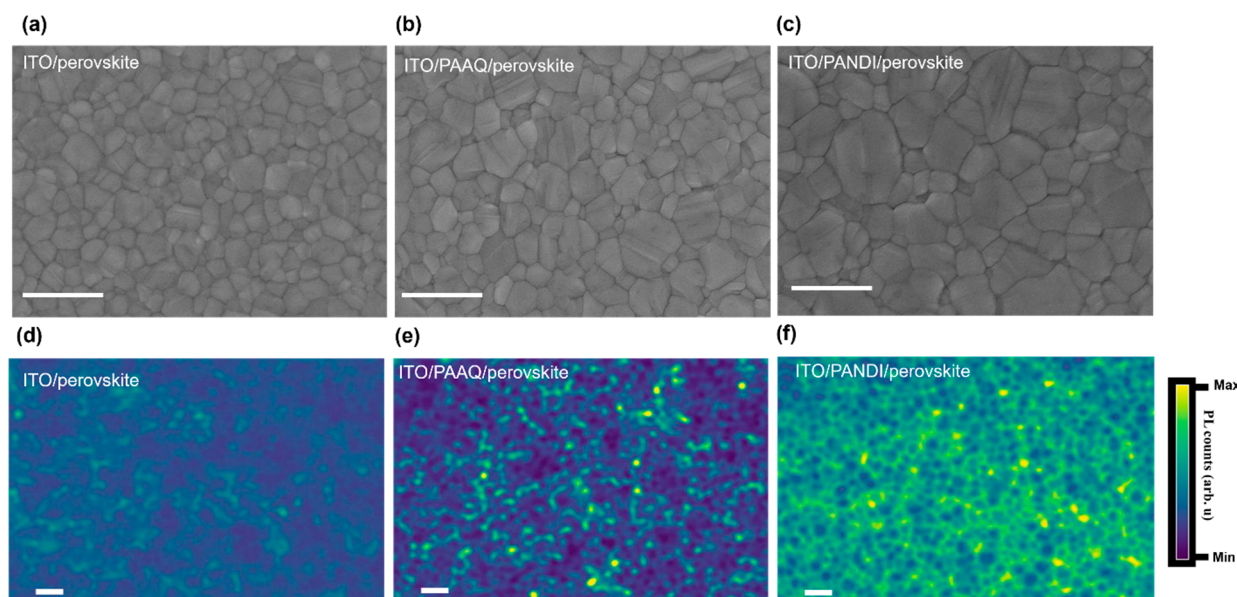
diimide and ammonium-bearing NDI polymers were applied as an interlayer between PC<sub>61</sub>BM and the metal electrode at the top contact to improve the performance of *p-i-n* PSCs.<sup>22,25,26</sup> Later, NDI SAMs, containing a carboxylic anchoring group, were applied directly onto indium tin oxide (ITO) substrates to lower their work function, enabling the collection of electrons in *n-i-p* PSCs.<sup>15</sup> These NDI-based devices demonstrated a PCE of 16% with a fill factor (FF) of 70%. Whereas these results show potential, they lag behind the best performances of regular *n-i-p* PSCs, relying typically on metal oxide ETLs (such as mesoporous TiO<sub>2</sub> or planar SnO<sub>2</sub>).<sup>27</sup> To narrow this discrepancy in device performance mandates further improvements in electron-selective NDI SAMs.

In the search for other options for electron-selective SAMs, anthraquinone (AQ) derivatives may also be considered. The AQ structure contains two electron-accepting carbonyl functional groups, which can increase the electron affinity and are easily tunable via molecule engineering and has been reported previously as an ETL in organic solar cells (OSCs).<sup>28,29</sup> In recent work, the AQ derivative 1-((3-(dimethylamino)propyl)amino)anthracene-9,10-dione was utilized in OSCs as an interface-modification layer of SnO<sub>2</sub>, improving the conductivity of the electron-selective contact stack and improving the energy level alignment with the organic photoactive layers.<sup>28</sup> The resulting devices showed a PCE of 18.1%, which is among the highest values reported for *n-i-p* OSCs.<sup>30</sup> Inspired by these results, we explore here how the AQ moiety can also be a promising candidate toward electron-selective SAMs in *n-i-p* PSCs.

We first introduce our modification of NDI and AQ SAMs, functionalized with a phosphonic acid group to anchor directly onto ITO, toward energetic matching with the perovskite. We then investigate their use as an alternative to conventional metal oxide (i.e., TiO<sub>2</sub>, SnO<sub>2</sub>, and ZnO) ETLs in *n-i-p* PSCs. Contrasting with conventional processed metal oxide ETLs that usually require a higher temperature processing ( $\geq 150$

$^{\circ}\text{C}$ ), these new electron-selective SAMs are solution processable and only require moderate temperatures (typically up to  $100$   $^{\circ}\text{C}$ ), which can be of benefit for certain applications, such as those relying on temperature-sensitive substrates.<sup>31–34</sup> Moreover, conventional solution-processed metal oxide ETLs usually require additional surface modifications with organic or inorganic materials to minimize interfacial charge losses, adding complexity toward the scalability of PSCs.<sup>35</sup> Here, we systematically investigated the SAMs' chemical and physical properties, including the energetic alignment and the electron extraction properties, at the ITO-SAM/perovskite interface, the materials' thermal stability, the surface wettability of ITO/SAM films, and their influences on perovskite crystallization. Finally, we fabricated *n-i-p* devices obtaining a champion PCE of 21.5%, representing the highest PCEs among *n-i-p* PSCs with organic ETLs. We also tested the operational stability of our SAM-based devices at  $65$   $^{\circ}\text{C}$  with above 90% retention of their initial performance for 1000 h.

The synthesis routes of *N*-(2,5-di-*tert*-butylphenyl)-*N'*-(methyl)-1,4,5,8-naphthalene tetracarboxylic diimide phosphonic acid (PANDI) and {3-[(9,10-dioxo-9,10-dihydroanthracen-2-yl)oxy]propyl} phosphonic acid (PAAQ) are shown in Scheme S1. The reaction conditions and detailed synthetic procedures are discussed in the Supporting Information (SI). These results were confirmed by using nuclear magnetic resonance (NMR) analysis, as shown in Figures S1–S8. Cyclic voltammetry (CV) and thermogravimetric analysis (TGA) were conducted to understand our SAMs' electrochemical and thermal stability properties. The redox potential from CV determines the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) energy levels of PANDI- and PAAQ-based SAMs. Note that these values do not represent absolute solid-state electron affinities or ionization energies but can be used to make relative comparisons between the different compounds. The voltammograms of PANDI and PAAQ in solution show reversible



**Figure 2.** (a–c) Top view of SEM images of perovskite on the bare-ITO, ITO/PAAQ, and ITO/PANDI surfaces, respectively, with a scale bar of 1  $\mu\text{m}$ . (d–f) Normalized hyperspectral mapping of the perovskite surface on bare-ITO, ITO/PAAQ, and ITO/PANDI with a scale bar of 2  $\mu\text{m}$ .

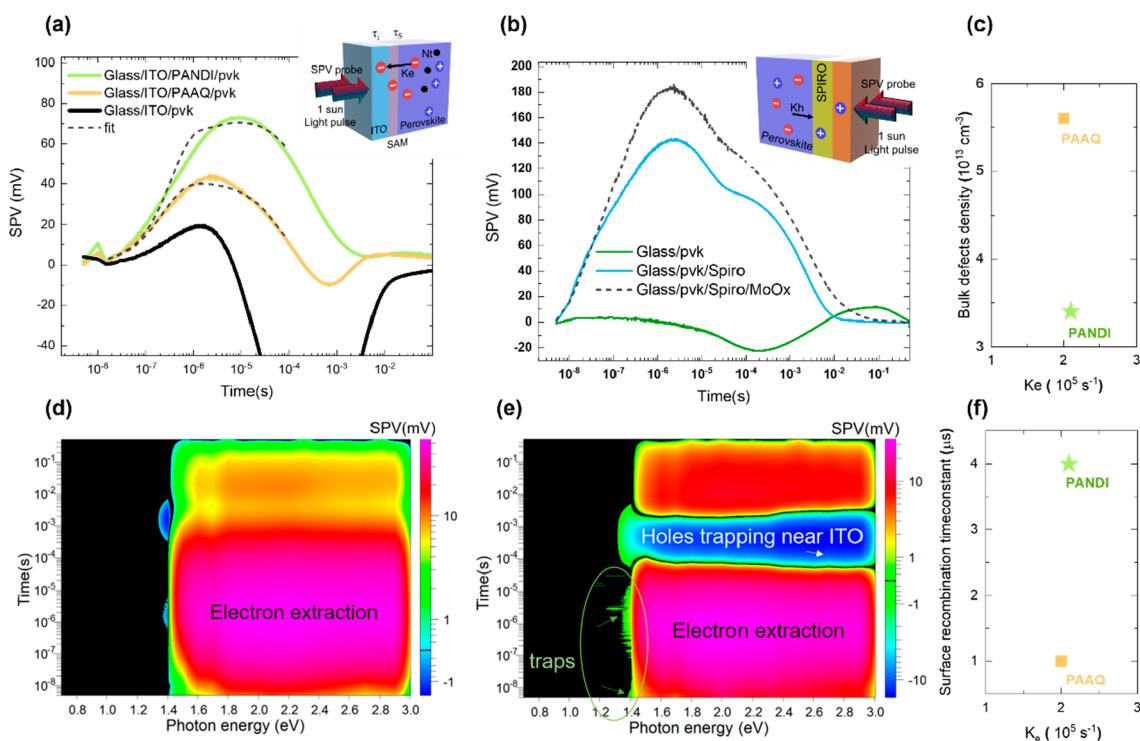
reduction waves during the scans and are typical for NDI and AQ derivatives (see Figure S9a). The reversible process indicates that both materials exhibit good electrochemical stability. The calculated energy level redox potential measurements result in  $E_{\text{LUMO}} = -3.81$  and  $-3.52$  eV while  $E_{\text{HOMO}} = -6.97$  and  $-6.79$  eV, respectively, for PANDI and PAAQ (summarized in Table S1). Both materials demonstrate good  $E_{\text{LUMO}}$  alignment with the conduction band minima (CBM) of the perovskite absorbers, while the  $E_{\text{HOMO}}$  energy levels are sufficiently high with reference to the valence band maxima (VBM) to block holes from the perovskite layer. In addition, both SAM molecules also provide good thermal and optical properties. For instance, TGA measurements prove that PANDI and PAAQ SAM molecules are sufficiently thermally stable with only 5% weight loss at 356 and 268  $^{\circ}\text{C}$ , respectively (Figure S9b). Furthermore, UV–vis transmittance results of SAM functionalization on the ITO surface show negligible optical losses compared to bare ITO and ITO/SnO<sub>2</sub> films (see Figure S10).

To gain a better understanding of how SAMs anchoring affects the work function (WF) of the ITO substrate, we also conducted calculations on the electric dipole moments of each SAM molecule in the gas phase (Figure 1a). Accordingly, we found that the net dipole moment of PAAQ at 3.94 D is smaller than that of PANDI (4.58 D). However, it is crucial to acknowledge that the net dipole moments of SAM molecules may not linearly follow the WF values, as the orientation and packing of the SAMs with respect to the ITO surface is still not fully understood.<sup>36,37</sup> Nevertheless, by assuming the perpendicular orientation of the SAM molecules to the substrates, the  $z$ -projection of the dipole moment becomes particularly relevant, given its higher contribution compared to the  $x$ - and  $y$ -projections. Upon calculation, we found that the  $z$ -projection dipole moment values for PAAQ and PANDI are  $-0.20$  and  $-1.04$  D, respectively (Figure S11 and Table S2). It is worth noting that while assuming a perpendicular orientation of PAAQ and PANDI to the surface, the presence of long alkyl chains connecting phosphonic acid and

anthraquinone in PAAQ may cause slight tilting of the SAMs. The negative sign indicates that the negative pole is directed toward the phosphonic acid anchoring group. The higher value of the dipole moment will enhance the WF of ITO to align better with the CBM of perovskite.

We then obtained the WF of the ITO/SAM samples from the contact potential difference (CPD; the detailed calculation is given in the SI) that was obtained by scanning Kelvin probe force microscopy (KPFM), as shown in Figure 1b and Figure S12. The WF of ITO/PANDI ( $\sim 4.45$  eV) is found to be lower than that of ITO/PAAQ ( $\sim 4.66$  eV), indicating a more favorable energetic alignment for electron extraction. Therefore, the larger WF difference of ITO/PANDI ( $\sim 0.27$  eV) than ITO/PAAQ ( $\sim 0.10$  eV), in reference to bare-ITO, enhances the band bending at the ITO/PANDI/perovskite interface, thereby improving its electron selectivity (see also the schematic illustration in Figure 1c).<sup>38,39</sup> The larger shift of the WF of PANDI can be associated with its larger interface dipole compared to PAAQ, as discussed above.<sup>40–42</sup> Apart from WF modifications, the SAM coatings do not significantly change the surface roughness of ITO, as shown by AFM results in Figure S13.

We also analyzed the chemical composition on the surface to understand the anchoring and coverage of SAM molecules on ITO by performing X-ray photoelectron spectroscopy (XPS; see Figure S14 and Table S3). Quantifying the surface coverage was also possible by calculating the presence of carbon (C 1s) and indium (In 3d<sub>5/2</sub>) in different positions across the sample area. This approach can determine the concentration of hydrogen bonding between SAMs and phosphonic acid attached to the indium-based substrate. The ratio P/In shows a high lateral variation in the surface region of PAAQ compared to PANDI on the ITO substrate (Tables S4 and S5). This analysis, therefore, provides a comprehensive picture of the electronic property dependence on the surface homogeneity. In addition, Figure 1d shows a surface mapping of CPD values of SAMs on ITO to further experimentally verify the uniformity of the SAM coverage. The more



**Figure 3.** (a, b) Charge extraction dynamics generated from trSPV measurements for ETL and HTL stack. (c) Bulk defect density. trSPV spectra from (d) PANDI and (e) PAAQ. (f) Surface recombination time constant defining how fast free carriers recombine near the perovskite surface.

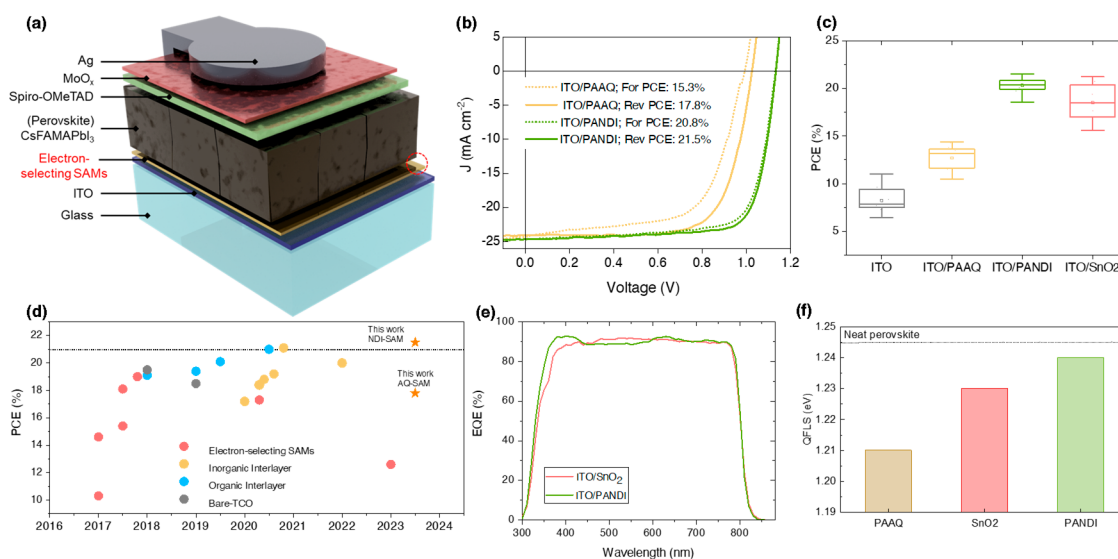
homogeneous pattern is proportional to the absence of WF fluctuations in a specific area. This shows that the ITO/PANDI interface has a narrower WF distribution than the ITO/PAAQ case, indicating a better PANDI coverage on ITO as illustrated in Figure 1e, in accordance with the XPS results.

The coated PANDI molecules onto ITO significantly increase ITO's surface hydrophobicity to a contact angle of around  $78.4^\circ$  versus PAAQ-coated ITO and bare ITO, with values of around  $42.2$  and  $28.5^\circ$ , respectively (Figure S15). It is noted that all of the samples were washed with chlorobenzene or methanol to remove unbound SAM molecules from the ITO substrate. This reveals that only SAMs attached onto ITO will affect the surface hydrophobicity that can be directly associated with our SAMs' coverage on ITO.<sup>43</sup> The higher surface hydrophobicity of the ITO/SAM substrate will affect the crystallization and morphology of the perovskite film.<sup>39,44</sup> The more hydrophobic (i.e., less wetting) surface of the ITO/PANDI case can enhance the surface mobility of the perovskite inks, resulting in larger-grain-size perovskite crystals, as indicated from top-view scanning electron microscopy (SEM) images in Figure 2a–c.<sup>39,44,45</sup> In addition, the functional head methyl group at PANDI can interact well with the perovskite layer, resulting in fewer pinholes or voids formed at the ITO/PANDI/perovskite interface (see the cross-sectional SEM image in Figure S16).<sup>3,4</sup> In contrast, the poor contact can result in some voids at the ITO/perovskite and ITO/PAAQ/perovskite bottom interface.

Next, we investigate the perovskite quality film via steady-state photoluminescence (PL) and time-resolved PL decay measurements. The steady-state PL intensity of the perovskite film coated on ITO/PANDI surfaces achieves a higher PL intensity than the ITO- and ITO/PAAQ-based samples (Figure S17a). This improvement is consistent with the

increased carrier lifetime of the ITO/PANDI/perovskite sample ( $\tau_{\text{average}}$  of  $3.1 \mu\text{s}$ ) compared to the ITO/PAAQ/perovskite sample ( $\tau_{\text{average}}$  of  $2.2 \mu\text{s}$ ), extracted from time-resolved PL measurements (Figure S17b). A larger perovskite crystal grain size implies a decrease of the relative amount of grain boundaries, directly reducing nonradiative recombination.<sup>39,44</sup> To visualize how the SAMs influence the perovskite film quality, confocal microscopy and local PL mapping of perovskite films was performed. Figure 2d–f shows hyperspectral PL microscopy images of the corresponding microscopy images above (Figure 2a–c) and corresponds to wavelengths emitted at 798 nm. We investigated the local distribution of PL on a microscale at the perovskite film surface.<sup>46</sup> This local PL distribution is correlated to the perovskite film quality, which is depicted through a color gradation from black to light yellow. For the ITO/perovskite sample, a lower PL emission spectrum is observed, indicated by a darker region, while the ITO/PAAQ/perovskite sample exhibits a higher PL emission but with a large spatial variation. Interestingly, the ITO/PANDI/perovskite sample has fewer local spatial variations with a stronger emission, indicating a higher perovskite film quality.<sup>46,47</sup>

To demonstrate the charge extraction capabilities at the ETL/perovskite and perovskite/HTL interfaces, we utilized the transient surface photovoltage (trSPV) with a 5 ns laser pulse and 1 sun equivalent fluence, as shown in Figure 3a,b. Generally, the trSPV signal is directly proportional to the concentration of extracted carriers, thereby providing a maximum signal amplitude and signal rise time, which offer qualitative information on the effectiveness of charge transport. Both SAMs exhibit positive signals, indicating efficient electron extraction (Figure 3a). Among them, PANDI demonstrated a larger amplitude and a faster rise time. The initial spike may be



**Figure 4.** (a) Device scheme of employed *n-i-p* PSCs with electron-selective SAM. (b) Representative *J-V* characteristics of PAAQ and PANDI under the forward and reverse scan directions. (c) PCE statistics for bare-ITO-, ITO/PAAQ-, ITO/PANDI-, and ITO/SnO<sub>2</sub>-based PSCs, respectively. (d) Summary of device efficiency of electron-selective SAMs and ETL-free *n-i-p* devices from the literature. (e) EQE spectra of ITO/SnO<sub>2</sub>- and ITO/PANDI-based devices. (f) QFLS values of perovskite film with different ETLs.

attributed to the transfer of a small fraction of hot carriers to the SAM. Notably, the ITO/perovskite interface exhibited a negative signal after 4  $\mu$ s, likely associated with free hole trapping or accumulation near the ITO interface. A similar behavior was observed for PAAQ, most probably induced by direct contact between ITO/perovskite due to the nonuniform coverage of PAAQ SAMs onto ITO, causing the trSPV signal to represent the superposition of electron extraction (6 ns to 4  $\mu$ s) and hole trapping (4  $\mu$ s to 10 ms). In contrast to electron extraction, the hole extraction by spiro-OMeTAD displayed a larger amplitude and a faster rise time (Figure 3b), indicating that the ETL/perovskite interface is a charge extraction bottleneck in the devices. It is noted that an additional MoO<sub>x</sub> layer can slightly enhance the hole extraction capabilities at the perovskite/spiro-OMeTAD/MoO<sub>x</sub> top contact stack.

We then simulated charge transport, using a model based on rate equations (eq 1), to provide a quantitative understanding of the charge extraction and the defect activation energies. Our model accounts for nonradiative recombination channels in the bulk and at the surfaces, characterized by the defect concentration in the bulk ( $N_t$ ) and time constants responsible for surface and interface recombination ( $\tau_s$  and  $\tau_i$ , respectively). The trSPV signal is given by eq S7. The dynamics of the electron concentration decay is given by the equation

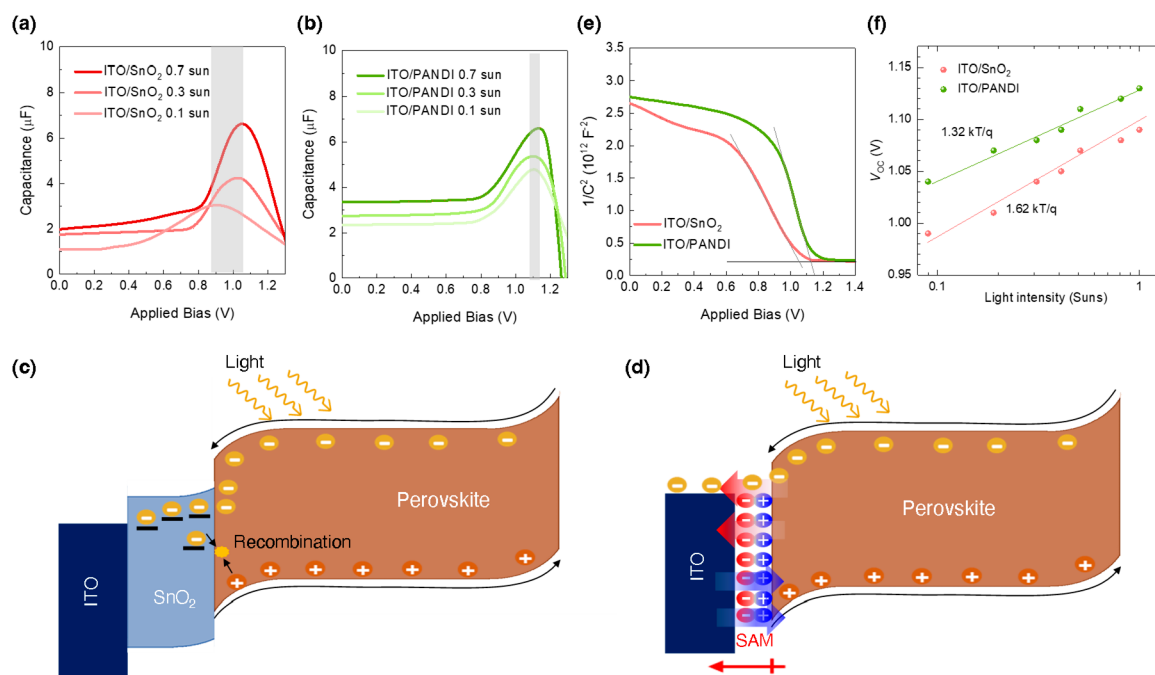
$$\frac{dn}{dt} = -K_{e\_ETL}n + K_{eb\_ETL}n_{ETL} - C_b(np) - n\sigma_{et}v_e(N_t - n_t) - \frac{n}{\tau_s} \quad (1)$$

where  $K_{e\_ETL}$  and  $K_{eb\_ETL}$  are electron rate constants for the extraction/injection of electrons from/to the perovskite, respectively.  $n$  and  $p$  are the photoinduced electron and hole densities, respectively.  $C_b$ ,  $\sigma_e$ , and  $v_e$  are the radiative recombination constant, electron capture cross-section of the charge trap, and the thermal velocity of electrons, respectively, taking values from previous work.<sup>8</sup> We then determined the

charge transport constants by fitting our model to the experimental data, using  $K_{e\_ETL}$ ,  $N_t$ ,  $\tau_s$ , and  $\tau_i$  as fitting parameters (dashed line in Figure 3a,b), shown in Table S6. The charge extraction rate constant showed a comparable value for both SAMs (Figure 3c).

To investigate the defect activation energies in the perovskite, we measured trSPV with a tunable wavelength laser with photon energies in the range of 0.7–3 eV, as shown in Figure 3d,e. The perovskite deposited on PAAQ exhibited clear signals below the bandgap, indicating charge accumulation due to the presence of a deep defect. In addition, the ITO/PAAQ/perovskite film also displayed a signature of hole trapping, demonstrated by the blue region in the SPV dynamics. In contrast, the ITO/PANDI/perovskite film displayed negligible signals below the bandgap and for hole trapping. As a result, perovskites deposited on PANDI showed a marked lower defect density ( $3.4 \times 10^{13} \text{ cm}^{-3}$ ) compared to the PAAQ ( $5.6 \times 10^{13} \text{ cm}^{-3}$ ), highlighting the beneficial role of PANDI in improving the electronic quality of the perovskite film (see Table S7).<sup>8</sup> Additionally, PANDI significantly suppressed surface and interface recombination by a factor of 4 (Figure 3f).<sup>48,49</sup> Furthermore, the capabilities of SAM in passivating surface and bulk defects, particularly through crystallization modification, resulted in better performance of the solar cell devices, as discussed below. The reported time constants are in agreement with trPL data in Figure S17b and previous reports for high-quality perovskite thin films.<sup>50–52</sup>

We then evaluated the performance of the PAAQ and PANDI SAMs on ITO as electron-selective materials at the bottom contact of *n-i-p* PSCs, with spiro-OMeTAD as HTL in the top contact (see Figure 4a). Following our previous work, we applied a 2D perovskite passivation treatment on top of perovskite films to improve the overall device performance and stability.<sup>9</sup> As shown in Figure S18, the absence of an electron-selective SAM layer on ITO leads to a high surface recombination at the bottom contact, limiting the device performance to a PCE of only 11.9%.<sup>8,53</sup> In contrast, insertion of the PAAQ and PANDI SAMs between the ITO/perovskite interface—besides enhancing the electron selectivity of the



**Figure 5.** Photoassisted capacitance–voltage of different devices: (a) SnO<sub>2</sub> and (b) PANDI. (c, d) Illustration of trap charge at interface of pristine SnO<sub>2</sub> and SAM based devices, respectively. (e) Mott–Schottky plot and (f)  $V_{OC}$  with respect to light intensity of devices based on SnO<sub>2</sub> and PANDI ETLs.

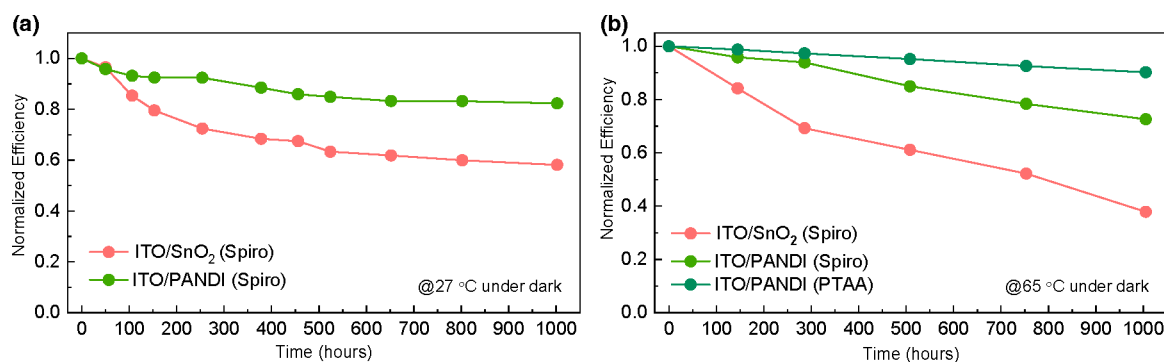
bottom contact—could suppress this type of recombination by repelling holes due to the strong internal electric field at the interface.<sup>19,39,54</sup> As a result, PAAQ-based devices show an increase in performance with a maximum PCE of 17.8% ( $V_{OC}$  of 1.03 V,  $J_{SC}$  of 24.1 mA cm<sup>-2</sup>, FF of 72%; Figure 4b). Despite this improvement over the SAM-free reference device, PAAQ-based devices feature limited  $V_{OC}$  and FF values, which could be due to the low surface coverage of PAAQ on ITO, resulting in unfavorable electronic properties and a lesser perovskite quality, as discussed above.

The PANDI SAM shows a more homogeneous coverage on ITO, with a higher surface dipole, enhancing the band bending and suppressing surface recombination effectively.<sup>14,41</sup> As a result, PANDI-based devices exhibit a higher device performance with a maximum PCE of 21.5% ( $V_{OC}$  of 1.13 V,  $J_{SC}$  of 24.7 mA cm<sup>-2</sup>, FF of 77%) under a reverse scan with a negligible hysteresis. This result is among the highest reported for electron-selective SAMs and ETL-free based *n-i-p* PSCs (see Table S8 and Figure 4d). The statistical distribution of our electron-selective SAMs from 25 devices is shown in Figure 4c and Figure S18, which, with its narrower PCE distribution, shows the reproducibility of our PANDI-based devices. We also fabricated *n-i-p* PSCs with pristine SnO<sub>2</sub> as the ETL for comparison, resulting in a PCE of ~21.2% (comparable to that in the literature; Figure S19).<sup>32,33,55,56</sup> In addition, the application of PANDI SAMs was further explored for flexible substrates, giving a PCE of 17.7% (Figure S20a,b). This outcome underscores the use of SAMs in accommodating substrates that require lower-temperature processing. The successful implementation of PANDI SAMs on ITO-PET highlights its potential for improving the performance of emerging technologies reliant on a flexible and low-temperature substrate (Table S9).<sup>57,58</sup>

Figure 5e shows the external quantum efficiency (EQE) of the PANDI- and SnO<sub>2</sub>-based devices. Interestingly, the EQE of the PANDI-based device exhibits a slightly higher overall  $J_{SC}$

than SnO<sub>2</sub>-based devices because of a higher absorption in the blue part of the spectrum, likely due to the lower optical reflectance losses of ITO/PANDI structures, as mentioned above. We also calculated the quasi-Fermi-level splitting (QFLS) from absolute PL measurements by using a calibrated 532 nm excitation laser at ~1 sun condition to understand the correlation to the  $V_{OC}$  improvement of our devices (Figure 4f). Notably, ITO/PANDI/perovskite samples show the highest QFLS value of around 1.24 eV compared to the ITO/SnO<sub>2</sub>/perovskite (1.23 eV)- and ITO/PAAQ/perovskite-based (1.21 eV) samples. The higher QFLS value on the PANDI indicates effective suppression of nonradiative recombination at the electron-selective buried interface.<sup>59</sup> This highlights how our PANDI-SAM yields lower optical reflectance and energetic losses, simultaneously, compared to the conventionally used SnO<sub>2</sub> ETL.

To evaluate charge extraction (accumulation/trapping) at the device level, we measured photoassisted capacitance–voltage ( $C-V$ ).<sup>60,61</sup> Figure 5a,b shows the  $C-V$  characteristics for PANDI and pristine SnO<sub>2</sub> ETL based devices under different light intensities. Interestingly, the change in peak potential ( $V_{peak}$ ) under reverse bias as the function of different light intensities is correlated with interfacial charge accumulation.<sup>60,61</sup> The SnO<sub>2</sub>-based device shows a significant  $V_{peak}$  shift due to interfacial charge trapping, as illustrated in Figure 5c. In contrast, due to a higher internal electric field from the dipole moment of SAM, the PANDI-based device exhibits a marginal  $V_{peak}$  shifting, indicating negligible charge accumulation (Figure 5d).<sup>14,37,39,41,60,62–64</sup> The reduced charge accumulation also resulted in a negligible photocurrent hysteresis in PANDI-based devices. Further, the enhanced built-in potential ( $V_{bi}$ ) values of PANDI-based devices extracted from Mott–Schottky plots correspond to the shallower WF of ITO/PANDI and a higher electric dipole of the SAM (Figure 5e). As a result, the higher  $V_{bi}$  can be



**Figure 6.** Evolution of normalized PCEs for nonencapsulated devices based on SnO<sub>2</sub>-ETL and PANDI SAM at (a) room temperature storage and (b) 65 °C under nitrogen and dark conditions.

associated with more efficient charge extraction/transport at the ITO/SAM/perovskite interface.<sup>31,65</sup>

On the other hand, Figure 5f shows the  $V_{OC}$  with respect to light intensity to determine the diode ideality factor of the PSCs. The diode ideality factor was calculated following  $n_{id} = \frac{q}{k_B T} \times \frac{dV_{OC}}{d \ln \theta}$ , where  $k_B T/q$  is the thermal voltage and can be extracted from the slope of the  $V_{OC}$  as a function of the light intensity.<sup>66</sup> The slope for SnO<sub>2</sub> devices shows a higher value ( $\sim 1.62 k_B T/q$ ) than that of PANDI-based ( $\sim 1.32 k_B T/q$ ) devices. Thus, the larger slope value, the greater the probability of trap-assisted recombination. The reduced ideality factor of PSCs with SAM compared to those with SnO<sub>2</sub> indicated suppressed trap-assisted bimolecular recombination at the electron-selective interface.<sup>67,68</sup>

Finally, we also evaluated the long-term stability of the *n-i-p* devices. First, we stored the devices based on SnO<sub>2</sub>-ETL and PANDI SAM inside a N<sub>2</sub>-filled glovebox at room temperature for 1000 h and measured the devices regularly (Figure 6a). The SnO<sub>2</sub>-based *n-i-p* devices could only retain around 58% of their initial PCEs after 1000 h of storage (in accordance with previous reports), while the PANDI SAM-based devices showed at 84% a much superior retention.<sup>69,70</sup> Next, we also evaluated the thermal stability of our devices kept on a hot plate at 65 °C in a N<sub>2</sub>-filled glovebox in the dark for more than 1000 h and compared PANDI SAM and SnO<sub>2</sub>-based devices in Figure 6b. This clearly showed that PANDI SAM-based devices demonstrated a better thermal stability, with around 72% retention of their initial PCEs, compared to SnO<sub>2</sub>-based devices with only 38% retention under identical conditions. The functional head group of PANDI consists of two methyl groups exhibiting a strong interaction to the PbI<sub>6</sub> framework in the perovskite, increasing the energy barrier for halide migration to the ETL bottom interface.<sup>71</sup> It should be noted that the degradation of *n-i-p* devices can also be initiated due to use of doped spiro-OMeTAD as an HTL under elevated temperature.<sup>2,17,72–76</sup> For this reason, we also evaluated our PANDI SAM-based devices employing poly[bis(4-phenyl)-(2,4,6-trimethylphenyl)amine] (PTAA), a more thermally stable HTL, with devices showing above 90% retention of their performance under identical conditions. In addition, the ITO-SnO<sub>2</sub>/perovskite interface also clearly further induced degradation due to redox reactions and charge accumulation during the stability tests, whereas these effects can be minimized with the use of organic SAM/perovskite interfaces.<sup>8,45,61,77–79</sup>

In summary, PANDI and PAAQ SAMs were successfully synthesized and investigated as electron-selective bottom contacts for *n-i-p* PSC. Both SAM molecules show a sufficient thermal stability and relatively suitable energetic alignment with the perovskite absorber and ITO contact. In particular, PANDI-based SAMs demonstrate a higher surface homogeneity on the ITO surface than PAAQ SAMs. We found that the increasing surface homogeneity on the ITO/PANDI can effectively suppress nonradiative interfacial recombination through the field-effect passivation, as indicated by longer charge carrier lifetimes and higher QFLS values. We also performed trSPV measurements and charge transport modeling, showing sufficient electron extraction and a relatively lower defect density at the ITO/PANDI/perovskite interface. Notably, the ITO/PANDI interface offers multiple benefits of lower optical reflectance and interfacial energetic losses, simultaneously, compared to the conventionally used SnO<sub>2</sub> ETL. As a result, ITO/PANDI-based *n-i-p* PSCs demonstrate a champion PCE of around 21.5%, representing the highest PCE among organic ETLs based on reported *n-i-p* PSCs, so far. Additionally, our PANDI SAMs also show the capability to be applied to flexible substrates. Finally, the PANDI-based device also showed improved operational long-term thermal stability, confirming that the PANDI SAM has a potential future to be utilized as ETL materials.

## ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsenenergylett.4c00306>.

Detailed experimental and characterization method, synthesis route, molecular NMR results, cyclic voltammetry and thermogravimetric measurements, SAM UV-vis spectra, Kelvin probe measurements, ITO/SAM topographic atomic microscopy, XPS analysis, surface water contact angle, cross-sectional SEM, steady-state PL and time-resolved PL, summary of photovoltaic parameters of various organic ETLs and ETL-free for *n-i-p* devices, device statistics,  $J-V$  performance for SnO<sub>2</sub> and flexible devices, and transient surface photovoltage fitting method (PDF)

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## Author Contributions

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## Author Contributions

D.S.U. and R.A. fabricated and did characterizations of perovskite samples and devices. L.M.S. and V.M. synthesized the SAMs and performed NMR measurements. L.M.S. performed TGA and CV measurements. D.S.U. performed SEM, hyperspectral photoluminescence, and UV–vis measurements. A.P. conducted the SKPM and AFM measurements and did analysis. P.D. performed XPS and did data analysis. A.M. conducted the trSPV measurements and analysis. R.A. and T.M. conceived the idea and wrote the original manuscript. R.A., D.S.U., L.M.S., T.M., A.M., E.A., and S.D.W. contributed

to reviewing and writing to the manuscript. R.A., T.M., V.G., and S.D.W. supervised the project. S.D.W. secured the funding. All authors contributed to the manuscript.

## Notes

The authors declare no competing financial interest.

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