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Influence of Schottky contact on the C-V and J-V characteristics of HTM-free perovskite solar cells

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Abstract The influence of the Schottky contact is studied for hole transport material (HTM) free $\text{CH}_3\text{NH}_3\text{PbI}_3$ perovskite solar cells (PSCs), by using drift-diffusion and small signal models. The basic current-voltage and capacitance-voltage characteristics are simulated in reasonable agreement with experimental data. The built-in potential of the finite $\text{CH}_3\text{NH}_3\text{PbI}_3$ layer is extracted from a Mott-Schottky capacitance analysis. Furthermore, hole collector conductors with work-functions of more than 5.5 eV are proposed as solutions for high efficiency HTM-free $\text{CH}_3\text{NH}_3\text{PbI}_3$ PSCs.

1 Introduction

Due to their potential for photo-induced carrier separation [1], various Hetero-Junction Solar Cells (HJSCs) have been experimentally [2, 3] and theoretically [4, 5] investigated. Selected doped functional materials are added on each side of the Light Harvesting Material (LHM) to select photo-induced carriers. The electron transport material (ETM) is used to extract photo-induced electrons and block holes, while the hole transport material (HTM) has a complementary function. HJSCs based on low-cost, easy processed [6–8] and highly absorbing [9, 10] semiconductor [11] halide perovskites have indeed led to high Photon-to-electron Conversion Efficiency (PCE) rising from 3.8% (2009) up to 22.1% (2016). Nowadays, these values are very close to the record value of silicon based solar cells (25.6%) [12].

As predicted from the detailed balance principle [13], if no defect-assisted recombination occurs in LHM and if the cell open circuit voltage (V_{OC}) equals to LHM's energy band gap (E_g) divided by elemental electron charge (q) the silicon based and the Perovskite based Solar Cells (PSC) should achieve maximum PCE of 44% and 37%, respectively. However, due to limited Internal PL quantum yield (iQY) and non-zero entropy, the maximum V_{OC} [14] is smaller than E_g :

$$qV_{OC} = E_g - T\Delta S - k_B T |\ln iQY| \quad (1)$$

where T is the absolute temperature and k_B is the Boltzmann constant. If optical losses are weak and the

contacts are almost ideal, an open circuit voltage (V_{OC}) of about 1.2 V is expected for $\text{CH}_3\text{NH}_3\text{PbI}_3$ [15]. In pace with the enhancement of stability [16, 17], the influence of defects was weakened down to an acceptable level [18, 19], while the band offsets between the LHM and ETM or HTM remain major factors impeding PCE [15, 20–23]. ZnO nanorods [24] or PCBM [25], ETM [26] are able to minimize the band offset at the conduction band minimum (CBM) and allow building almost ideal contact at ETM/LHM interface. However HTM very often present large band offsets at valence band maximum (VBM) and low carrier mobility [23, 27–29]. Alternatively, PSC without HTM layer was proposed as a solution towards high efficiency. After Etgar and coworkers early directly deposited gold on $\text{CH}_3\text{NH}_3\text{PbI}_3$ and demonstrated that the $\text{CH}_3\text{NH}_3\text{PbI}_3$ material can be simultaneously considered as a light harvester and a hole conductor, leading to PCE of 8% [30]. Then porous carbon film was used as contact for fully printable HTM-free PSCs with efficiency of 12.8% [17]. And PSCs with single-walled carbon nanotubes as hole collector achieved efficiency of 15% [31, 32]. In such case, solar cells benefit from fewer interfaces, and the optical and electrical losses in the HTM layer are eliminated as well. In Figure 1, the architecture of the HTM-free $\text{CH}_3\text{NH}_3\text{PbI}_3$ PSC is schematic represented by comparison to that of classical PSC Gold for example, is directly connected with $\text{CH}_3\text{NH}_3\text{PbI}_3$ as the hole collection electrode and a Schottky contact is formed [33].

To get an insight into the HTM-free PSC operation, direct current and small signal simulation analyses [5, 34–37]

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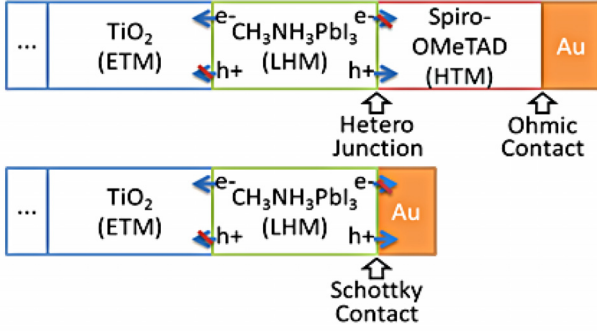


Fig. 1. Schematic representations of perovskite solar cells with (a) a hetero-junction or (b) a Schottky contact at hole collector side, respectively.

were performed including basic semiconductor models: the Poisson equation, the current continuity equation and a drift-diffusion model. The critical transport and recombination processes in solar cells can thus be quantitatively analyzed. Nevertheless, few numerical analyses were dedicated up to now to HTM-free PSCs. In our work, the basic current-voltage (J - V) and capacitance-voltage (C - V) characteristics of HTM-free $\text{CH}_3\text{NH}_3\text{PbI}_3$ based PSCs are studied with drift-diffusion and small signal models [38], which are integrated in Silvaco Atlas simulator [39].

2 Numerical modeling

The physical model is numerically simulated in Atlas by solving a set of coupled equations including Poisson's equation (2), continuity (3a) and (3b) and transport equations (4a) and (4b) for electrons and holes densities. These equations link together the electrostatic potential profile and the charge distributions, and describe the evolution of electron and hole densities under external bias and light illumination, including carrier transport, generation, and recombination processes. The bimolecular recombination model corresponds to the formula (5). The trap-assisted recombination model is described in formula (6), (7a) and (7b), while the photo-induced carrier generation processes are introduced through complex refractive index of materials. Simulations were carried out under equilibrium and small AC conditions, with and without AM1.5 sun illumination, in order to obtain J - V and C - V characteristics of HTM-free $\text{CH}_3\text{NH}_3\text{PbI}_3$ PSCs

$$\Delta\psi = -\frac{\rho}{\varepsilon} \quad (2)$$

where ψ is the potential, ρ is the charge density and ε is the dielectric constant.

$$\frac{\partial n}{\partial t} = \frac{1}{q} \text{div} \vec{J}_n + G_n - R_n \quad (3a)$$

$$\frac{\partial p}{\partial t} = -\frac{1}{q} \text{div} \vec{J}_p + G_p - R_p \quad (3b)$$

where $n(p)$ is the electron (hole) density, t is the time, J_n (J_p) is the electron (hole) current density, G and R are

the generation and recombination rates respectively. The footnote $n(p)$ is related to electron (hole).

$$\vec{J}_n = -q\mu_n n \nabla \phi_n \quad (4a)$$

$$\vec{J}_p = -q\mu_p p \nabla \phi_p \quad (4b)$$

where μ is mobility and ϕ is the quasi-Fermi level

$$R_{bi} = k_{bi} (np - n_i^2) \quad (5)$$

where k_{bi} is the bimolecular recombination coefficient and n_i the intrinsic electron density

$$R_{SRH} = \frac{pn - n_i^2}{\tau_n \left[p + n_i \cdot \exp\left(\frac{-\Delta E}{k_B T}\right) \right] + \tau_p \left[n + n_i \cdot \exp\left(\frac{\Delta E}{k_B T}\right) \right]} \quad (6)$$

$$\tau_n = \frac{1}{SIG_n \cdot v_n \cdot N_t} \quad (7a)$$

$$\tau_p = \frac{1}{SIG_p \cdot v_p \cdot N_t} \quad (7b)$$

The Shockley-Read-Hall (SRH) recombination mechanism is described by equation (6) τ is the charge carrier lifetime for trap-assisted process. The relationship between τ and trap density (N_t) (Eq. (7)) depends on the traps capture cross section (SIG) and the thermal velocity (v). ΔE is the absolute energy difference between the trap level and the intrinsic Fermi level (E_i) in the bulk. E_i is approximately located in the middle of energy band gap. If $\Delta E = 0$, the maximum of SRH recombination rate is obtained. In other words, the deep trap centers lead to the highest recombination rates and are harmful for photo-induced carrier extraction. We set $\tau_n = \tau_p$, $\mu_n = \mu_p$ to reduce the number of parameters in the present work.

3 Basic properties of HTM-free PSC

A basic modeling of HTM-free Perovskite Solar Cells (PSCs) studied experimentally by Etgar's group [40], relies on $\text{TiO}_2/\text{CH}_3\text{NH}_3\text{PbI}_3/\text{Au}$ architecture with a computed static band alignment shown in Figure 2. Heavily n-type doped ETM anatase (TiO_2) [41–43] and hole collector gold are added on each sides of lightly n or p-doped $\text{CH}_3\text{NH}_3\text{PbI}_3$ [1, 44, 45]. Under thermal equilibrium and short circuit condition, the Fermi-level (E_f) remains constant as reference through the device, and the band offsets at each interface yield two potential barriers. It is clear that the major potential drop $\Delta\psi$ is located in the $\text{CH}_3\text{NH}_3\text{PbI}_3$ layer. Therefore, the major part of the electrical current originates from carrier drifting rather than carrier diffusion. The main properties of the materials used for the simulation are summarized in Table 1, including χ , E_g , the doping level (N), the effective masses of electron and hole (m_e^* & m_h^*) and the relative dielectric constant (ε_r). The thickness of TiO_2 and $\text{CH}_3\text{NH}_3\text{PbI}_3$ layers are both equal to 300 nm. The work function (WF) of gold is 5.1 eV [33, 46]. An Ohmic contact is considered

Table 1. Main properties of the materials.

	X (eV)	E_g (eV)	N (cm^{-3})	$m_e^* & m_h^*$ (m_0)	ϵ_r (ϵ_0)
HOIP	3.9	1.6	$n = 4e16^*$ or $p = 8e16^*$	0.23&0.29 [48]	70 [49]
TiO ₂	4.1	3.3	1e19 [41]	5.6&5.6 [50]	31 [51]

Notes: HOIP means CH₃NH₃PbI₃; * indicates fitting parameter.

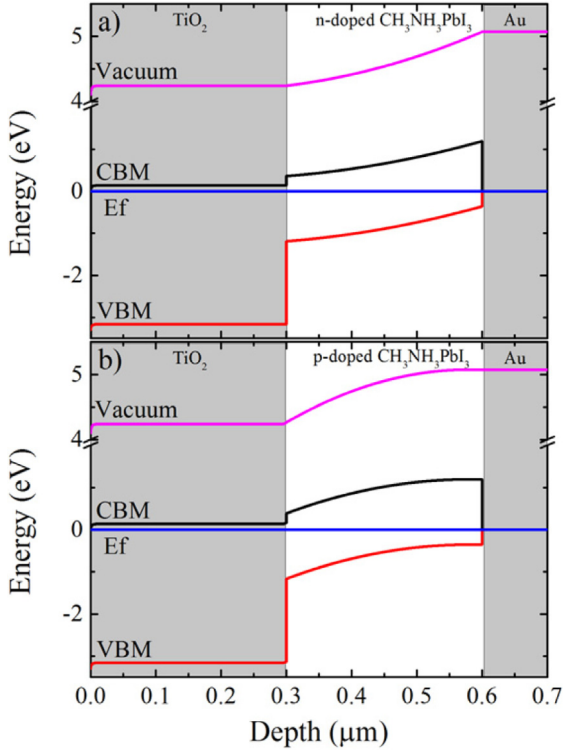


Fig. 2. Static band diagram of (a) n-doped and (b) p-doped CH₃NH₃PbI₃ based HTM-free PSCs.

at the bottom of the TiO₂ layer on the other side. The bimolecular recombination coefficient k_{bi} of CH₃NH₃PbI₃ is 10^{-9} cm³/s [47]. Due to lack of precise trap characterization, ΔE is set to zero and N_t is 10^{10} cm⁻³, while τ and μ are tuned to match the experimental data.

3.1 Capacitance characteristics

In order to obtain efficient energy conversion in solar cells with low mobility LHM, a high build in potential (V_{bi}) is necessary to prevent significant losses due to carrier recombination processes competing with charge extraction processes [52]. In our case, a Schottky contact [53] is formed at interface CH₃NH₃PbI₃/Au. Therefore, V_{bi} can be extracted from a Mott-Schottky capacitance analysis. The device architecture in our work is shown together with the circuit in Figure 3. The capacitance expression

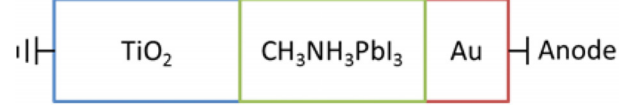


Fig. 3. Circuit used for Mott-Schottky capacitance analysis.

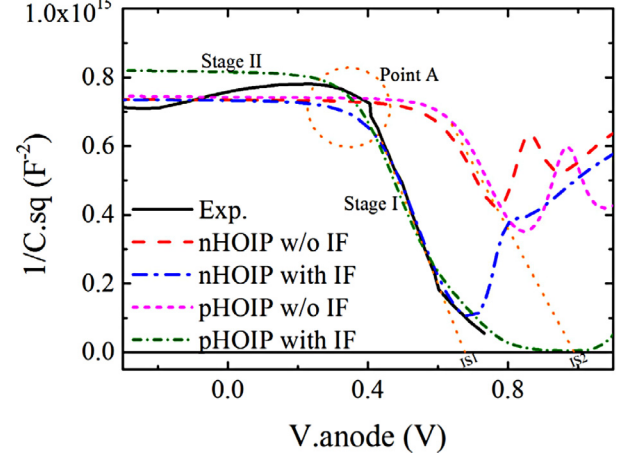


Fig. 4. The computed and experimental (solid line) C - V characteristics in dark of HTM-free PSC. N-doped HOIP with (without) an n-doped interfacial layer (IF) at Au/HOIP interface is indicated as dash dot (dash) line. P-doped HOIP with (without) a p-doped IF at TiO₂/HOIP interface is indicated as short dash dot (short dash) line. The V_{bi} values extracted from the IS1 and IS2 intersections are equal to 0.6 and 0.9 V, respectively.

is given by:

$$\frac{1}{C^2} = \frac{2}{qA^2\epsilon_0} \left(\frac{1}{N\epsilon_r} \right) \left(V_{bi} - V_{bias} + \frac{k_B T}{q} \right) \quad (8)$$

where C is the junction capacitance, A is the junction area, ϵ is the vacuum permittivity, N is the activated dopant density in semiconductor and ϵ_r is the relative permittivity. In our work, a small signal analysis [38] is used to simulate C - V characteristics of HTM-free CH₃NH₃PbI₃ PSC. The signal frequency is set at 1 kHz for simulation, as in practical measurement. The theoretical characteristics is presented in Figure 4 and compared to available experimental data [40]. In order to fit the experimental data, an effective interfacial layer (IF) was introduced into the architecture for each type of CH₃NH₃PbI₃. For n-doped CH₃NH₃PbI₃, the IF of 8.5 nm is heavily n-doped and located between Au and CH₃NH₃PbI₃. For p-doped CH₃NH₃PbI₃, the IF of 3.4 nm is heavily p-doped and located between TiO₂ and CH₃NH₃PbI₃. The doping level of each IF is equal to $2e19$ cm⁻³. The influence of IF is further discussed at the end of the section.

As the bias reversely increases, the extension of the depletion region starts at the CH₃NH₃PbI₃/Au interface, then goes through the CH₃NH₃PbI₃ and finally into the TiO₂. Because of the different N and ϵ_r values in both CH₃NH₃PbI₃ and TiO₂, C - V curves under reverse bias are bent into two stages. Similar phenomena were observed

for III-V semiconductors [54,55]. The roughly constant capacitance at stage II is due to the heavy doping level in TiO_2 , in comparison with the smaller slope related to the small doping level in $\text{CH}_3\text{NH}_3\text{PbI}_3$. The point A in Figure 4 corresponds to the transition point of the depletion region from stage I to stage II. According to expression (8), the V_{bi} of a finite $\text{CH}_3\text{NH}_3\text{PbI}_3$ layer is extracted from the intersection (IS) as pointed out in Figure 4. The fluctuations of experimental data at stage II can be explained by non-uniform doping in the TiO_2 layer. Because of the huge effective surface area of nano-porous TiO_2 [56–58] and the rough surface of $\text{CH}_3\text{NH}_3\text{PbI}_3$ layer [58], it is risky to extract N , ϵ_r or thickness (d) of $\text{CH}_3\text{NH}_3\text{PbI}_3$ from the expression (8) and classic parallel plate capacitance expression (9), directly. In the model the effective area of capacitance interface (A_{eff}) is around two times as large as the active area of practical gold electrode

$$C = \frac{\epsilon_r \epsilon_0 A_{eff}}{d}. \quad (9)$$

If more uniform growth of material layers for HTM-free PSCs is achieved in the future, it will be possible to extract more quantitative information from C - V measurements, related to N , ϵ_r and the thickness of the $\text{CH}_3\text{NH}_3\text{PbI}_3$ layer.

3.2 Photovoltaic characteristics

Due probably to the different growth procedures employed by the experimental groups, some deviations are found for the absorption coefficients values of $\text{CH}_3\text{NH}_3\text{PbI}_3$ in reference [59]. For that reason, the absorption coefficient curve used in our simulation was rather obtained by fitting the experimental IPCE spectrum data. In Figure 5, the simulated J - V characteristics under 1 sun of 1.5 AM illumination are presented along with experimental data. A good matching to the experimental J - V curve is achieved based on n -type $\text{CH}_3\text{NH}_3\text{PbI}_3$ when its $\tau = 80$ ns and $\mu = 0.2$ cm^2/Vs . These empirical values are consistent with commonly measured values for the $\text{CH}_3\text{NH}_3\text{PbI}_3$ material. The paper will only discuss n -doped $\text{CH}_3\text{NH}_3\text{PbI}_3$ based PSCs in the following sections, because a better agreement with experimental C - V and J - V characteristics is obtained in this case

From the comparison of the experimental and computed C - V characteristics (Fig. 4), it is necessary to assume that a heavily n -doped IF exists at the $\text{CH}_3\text{NH}_3\text{PbI}_3/\text{Au}$ contact. Alternative hypotheses with layers containing acceptors or surface states, were explored but without success. Indirect evidences of the existence of such an IF can be found in the report of Liu's group [33]. Using ultraviolet photoemission spectroscopy (UPS), these authors indeed showed that during the deposition of the gold contact, the Fermi level undergoes a progressive shift. Noteworthy, the presence of metal nano particles [60] or charged ions [61–63] at the interface was discussed by other groups. This Fermi level shift is simulated in the present work by introducing an effective and

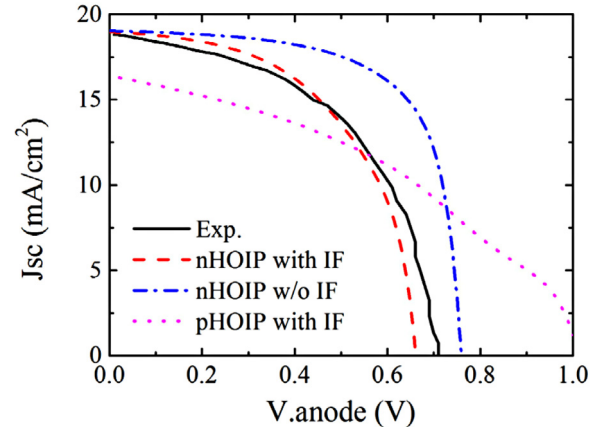


Fig. 5. J - V characteristics under 1 sun illumination of n -doped $\text{CH}_3\text{NH}_3\text{PbI}_3$ based HTM-free PSCs with (dash line) and without (dash dot line) interfacial layer (IF). The τ and μ of n -doped $\text{CH}_3\text{NH}_3\text{PbI}_3$ are equal to 80 ns and 0.2 cm^2/Vs , respectively. And p -type $\text{CH}_3\text{NH}_3\text{PbI}_3$ is pictured by dot line as example with τ of 18 ns and μ of 0.2 cm^2/Vs .

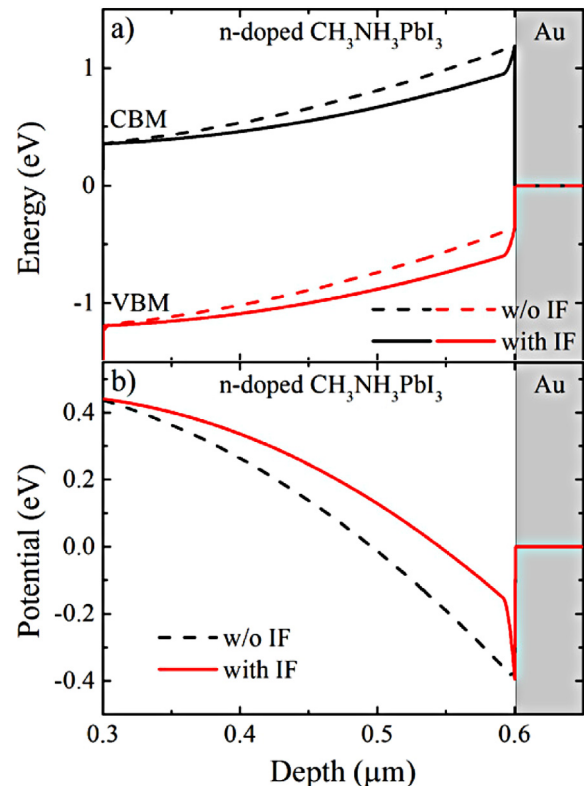


Fig. 6. (a) Static band alignment and (b) potential profile with and without interfacial layer (IF).

heavily n -doped $\text{CH}_3\text{NH}_3\text{PbI}_3$ IF. The high density of positive ionized charges in the IF leads to a reduction of the V_{bi} from 0.9 to 0.6 V, in good agreement with the experimental value (Fig. 4).

The static band alignment and potential profile with and without IF are represented in Figures 6a and 6b, to have an insight into the device operation. Even though

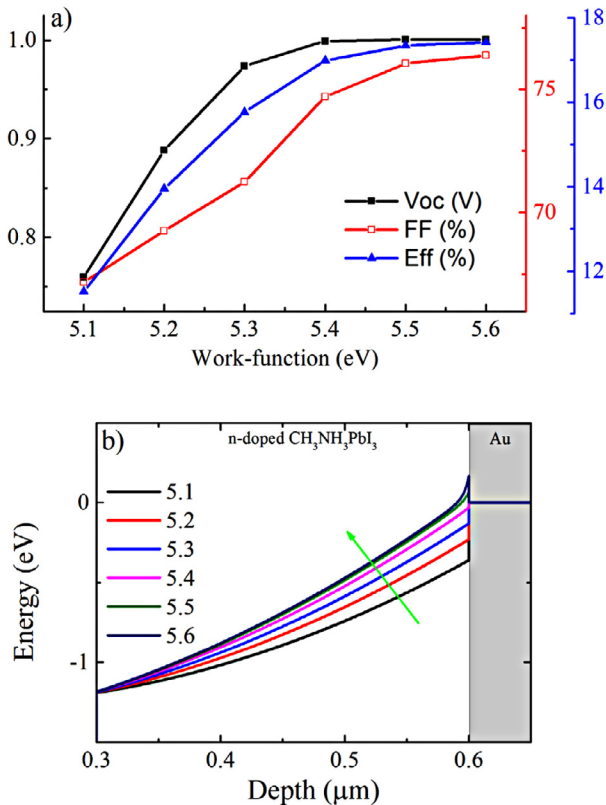


Fig. 7. (a) Open circuit voltage (V_{OC}), fill factor (FF) and efficiency (Eff) of HTM-free CH₃NH₃PbI₃ PSC as a function of the work-function (WF) of hole collector conductor. (b) Valence band variation as a function of WF.

the band offset at the CH₃NH₃PbI₃ surface is pinned by the gold contact, the effective potential drop across the CH₃NH₃PbI₃ layer is lowered by the presence of the IF. As a consequence, the losses due to carrier recombination processes increase and the efficiency of PSC decreases from 11% to 8%, as shown in Figure 5. A thick IF has clearly a detrimental effect on the photovoltaic efficiency.

4 Influence of work-function

The efficiencies of the PSC can be increased by improving the intrinsic properties of the perovskite (τ and μ). We propose in this work to explore an additional possibility for HTM-free PSC. It is indeed possible to enlarge V_{bi} by increasing the WF of the metal used for the Schottky contact. In Figure 7a, V_{OC} , fill factor and efficiency are presented as a function of WF. These parameters are improved until saturation is reached for a WF value of 5.6 eV, while the short circuit current (J_{SC}) is almost constant and equal to 19 mA/cm². The efficiency of HTM-free PSC can be enhanced up to 17% (Fig. 7a) by this way, and as stated before further enhancements could also be expected by improving the intrinsic properties of the perovskite.

Furthermore, as shown in Figure 7b, the change of VBM is very small except for the part close to the

CH₃NH₃PbI₃/Au interface, as the WF of the metal increases up to 5.6 eV, larger than the VBM of CH₃NH₃PbI₃ (5.5 eV). As a result, the overall V_{bi} in CH₃NH₃PbI₃ layer becomes almost saturated. When the carrier recombination rate is small enough, the J_{SC} mainly depends on absorption properties and is almost independent of V_{bi} . Palladium [46, 64] or Selenium [65] are examples of hole collector conductors with WF larger than 5.5 eV.

5 Conclusion

In summary, a detailed investigation of C - V and J - V characteristics of HTM-free CH₃NH₃PbI₃ PSC has been proposed, based on the drift-diffusion model and small signal analysis. The simulation results are in good agreement with experimental data. An effective heavily doped interfacial layer was introduced at the interface to fit the C - V characteristics. It is also shown in this work that, an increase of WF of the hole collector conductor, is expected to enhance the PSC efficiency.

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References

1. C.-S. Jiang et al., Nat. Commun. **6**, 8397 (2015)
2. M. Grätzel, J. Photochem. Photobiol. C Photochem. Rev. **4**, 145 (2003)
3. W. Zhang, G.E. Eperon, H.J. Snaith, Nat. Energy **2016**, 16048 (2016)
4. Q. Wang et al., J. Phys. Chem. B **110**, 25210 (2006)
5. J. Bisquert, L. Bertoluzzi, I. Mora-Sero, G. Garcia-Belmonte, J. Phys. Chem. C **118**, 18983 (2014)
6. W. Nie et al., Science **347**, 522 (2015)
7. M.M. Lee, J. Teuscher, T. Miyasaka, T.N. Murakami, H.J. Snaith, Science **338**, 643 (2012)
8. J. Burschka et al., Nature **499**, 316 (2013)
9. J. Even, L. Pedesseau, J.-M. Jancu, C. Katan, J. Phys. Chem. Lett. **4**, 2999 (2013)
10. J.M. Ball et al., Energy Env. Sci **8**, 602 (2015)
11. J. Even et al., J. Phys. Chem. C **119**, 10161 (2015)
12. NREL Efficiency Chart Rev. (2016) Available at: http://www.nrel.gov/ncpv/images/efficiency_chart.jpg
13. W. Shockley, H.J. Queisser, J. Appl. Phys. **32**, 510 (1961)
14. C.M. Sutter-Fella et al., Nano Lett. **16**, 800 (2016)
15. P. Gao, M. Grätzel, M.K. Nazeeruddin, Energy Environ. Sci. **7**, 2448 (2014)
16. W. Nie et al., Nat. Commun. **7**, 11574 (2016)
17. A. Mei et al., Science **345**, 295 (2014)
18. W. Qiu et al., Energy Env. Sci **9**, 484 (2016)
19. T.M. Brenner, D.A. Egger, L. Kronik, G. Hodes, D. Cahen, Nat. Rev. Mater. **1**, 15007 (2016)
20. T. Minemoto, M. Murata, Sol. Energy Mater. Sol. Cells **133**, 8 (2015)
21. E.J. Juarez-Perez et al., J. Phys. Chem. Lett. **5**, 680 (2014)
22. W. Li et al., Energy Env. Sci. **9**, 490 (2016)
23. P. Schulz et al., Energy Environ. Sci. **7**, 1377 (2014)

24. J. Dong, J. Shi, D. Li, Y. Luo, Q. Meng, *Appl. Phys. Lett.* **107**, 073507 (2015)
25. O. Malinkiewicz et al., *Nat. Photonics* **8**, 128 (2014)
26. K.-W. Tsai, C.-C. Chueh, S.T. Williams, T.-C. Wen, A.K.Y. Jen, *J. Mater. Chem. A* **3**, 9128 (2015)
27. M. Saliba et al., *Nat. Energy* **1**, 15017 (2016)
28. C. Chappaz-Gillot et al., *Sol. Energy Mater. Sol. Cells* **120**, 163 (2014)
29. W. Chen et al., *Science* **350**, 944 (2015)
30. L. Etgar et al., *J. Am. Chem. Soc.* **134**, 17396 (2012)
31. S.N. Habisreutinger et al., *J. Phys. Chem. Lett.* **5**, 4207 (2014)
32. K. Aitola et al., *Energy Env. Sci.* **9**, 461 (2016)
33. X. Liu et al., *Phys. Chem. Chem. Phys.* **17**, 896 (2014)
34. V. Gonzalez-Pedro et al., *Nano Lett.* **14**, 888 (2014)
35. Y.T. Set, B. Li, F.J. Lim, E. Birgersson, J. Luther, *Appl. Phys. Lett.* **107**, 173301 (2015)
36. X. Sun, R. Asadpour, W. Nie, A.D. Mohite, M.A. Alam, *IEEE J. Photovolt.* **5**, 1389 (2015)
37. Y.T. Set, E. Birgersson, J. Luther, *Phys. Rev. Appl.* **5**, 054002 (2016)
38. S.E. Laux, *IEEE Trans. Electron Devices* **32**, 2028 (1985)
39. Silvaco Inc., ATLAS user's manual (2012), <http://silvaco.com>
40. W.A. Laban, L. Etgar, *Energy Environ. Sci.* **6**, 3249 (2013)
41. G. Liu, W. Jaegermann, J. He, V. Sundström, L. Sun, *J. Phys. Chem. B* **106**, 5814 (2002)
42. H. Tang, K. Prasad, R. Sanjinès, P.E. Schmid, F. Lévy, *J. Appl. Phys.* **75**, 2042 (1994)
43. L. Forro et al., *J. Appl. Phys.* **75**, 633 (1994)
44. E.M. Miller et al., *Phys. Chem. Chem. Phys.* **16**, 22122 (2014)
45. Q. Wang et al., *Appl. Phys. Lett.* **105**, 163508 (2014)
46. H.B. Michaelson, *J. Appl. Phys.* **48**, 4729 (1977)
47. A. Paulke et al., *Appl. Phys. Lett.* **108**, 113505 (2016)
48. G. Giorgi, J.-I. Fujisawa, H. Segawa, K. Yamashita, *J. Phys. Chem. Lett.* **4**, 4213 (2013)
49. Q. Lin, A. Armin, R.C.R. Nagiri, P.L. Burn, P. Meredith, *Nat. Photon.* **9**, 106 (2014)
50. S. Rühle, D. Cahen, *J. Phys. Chem. B* **108**, 17946 (2004)
51. S. Roberts, *Phys. Rev.* **76**, 1215 (1949)
52. A. Pivrikas et al., *Phys. Rev. Lett.* **94**, 176806 (2005)
53. C. Wang et al., *J. Vac. Sci. Technol. B* **33**, 032401 (2015)
54. A. Morii, H. Okagawa, K. Hara, J. Yoshino, H. Kukimoto, *Jpn J. Appl. Phys.* **31**, L1161 (1992)
55. H. Kroemer, W.-Y. Chien, J.S.H. Jr., D.D. Edwall, *Appl. Phys. Lett.* **36**, 295 (1980)
56. S. Nakade et al., *Electrochem. Commun.* **5**, 804 (2003)
57. P.M. Sommeling et al., *J. Phys. Chem. B* **110**, 19191 (2006)
58. S. Gamliel, A. Dymshits, S. Aharon, E. Terkieltaub, L. Etgar, *J. Phys. Chem. C* **119**, 19722 (2015)
59. N.-G. Park, *Nano Converg.* **3**, 1 (2016)
60. W. Zhang et al., *Nano Lett.* **13**, 4505 (2013)
61. Y. Yuan et al., *Adv. Energy Mater.* **6**, 1501803 (2016)
62. J.S. Yun et al., *Adv. Energy Mater.* **6**, 1600330 (2016)
63. H. Yu, H. Lu, F. Xie, S. Zhou, N. Zhao, *Adv. Funct. Mater.* **26**, 1411 (2016)
64. L. Baojun, L. Enke, Z. Fujia, *Solid-State Electron.* **41**, 917 (1997)
65. A.M. Patil, V.S. Kumbhar, N.R. Chodankar, A.C. Lokhande, C.D. Lokhande, *J. Colloid Interface Sci.* **469**, 257 (2016)

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