

SCIENTIFIC REPORTS

OPEN

Spiro-Phenylpyrazole/Fluorene as Hole-Transporting Material for Perovskite Solar Cells

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Spiro-OMeTAD with symmetric spiro-bifluorene unit has dominated the investigation of hole-transporting material (HTM) for efficient perovskite solar cells (PSCs) despite of its low intrinsic hole conductivity and instability. In this study, we designed and synthesized three asymmetric spiro-phenylpyrazole/fluorene base HTMs, namely: WY-1, WY-2 and WY-3. They exhibit excellent electrochemical properties and hole conductivities. Moreover, the PSC based on WY-1 exhibits the highest power conversion efficiency (PCE) of 14.2%, which is comparable to the control device employing spiro-OMeTAD as HTM (14.8%). These results pave the way to further optimization of both molecular design and device performance of the spiro-based HTMs.

Organic-inorganic lead halide perovskites have received a great deal of attention since the first report of perovskite solar cell (PSC) by Miyasaka *et al.*¹. These materials possess many attractive merits, including strong visible absorbability^{2,3}, high carrier mobility^{4,5} and long carrier diffusion length^{6,7}. The rational design of devices, effective modulation of interfacial contact and development of superior charge transporting materials have all contributed to the recent record-high power conversion efficiency (PCE) of 22.1%⁸ (<http://www.nrel.gov/>). To date, 2,2',7,7'-tetrakis(*N,N*-di-*p*-methoxyphenylamine)-9,9'-spirobifluorene (spiro-OMeTAD) has been utilized as the hole-transporting materials (HTMs) in fabrication of PSCs with PCEs of over 20%^{8–10}. However, spiro-OMeTAD suffers from poor hole mobility in the pristine form due to its poor crystallinity caused by the spiro structure^{11,12}. Generally speaking, there are two strategies to overcome such drawbacks^{13–16}. The first method is to replace spiro skeleton with other ensembles such as pyrene¹⁷, 4,4-*N,N'*-dicarbazole-1,1'-biphenyl (CBP)¹⁸, quinolizino acridine¹⁹, functional thiophene^{20,21}, 1,1,2,2-tetraphenylethene (TPE)^{22,23} and bifluorenylidene²⁴, *etc.* for tuning the HOMO energy, hole mobility and conductivity^{25–31}. The second method is to alter the spiro-bifluorene unit with the acridine/fluorene^{32,33} or xanthene/fluorene^{34,35} core to achieve the superior device performances.

Herein, we presented unprecedented HTMs based on spiro-phenylpyrazole/fluorene architecture; namely: WY-1, WY-2 and WY-3 (Fig. 1a). They exhibit excellent hole transporting properties as half of the molecule is constructed based on spiro-OMeTAD, while the orthogonal phenyl pyrazole fragments were fine-tuned by varying the π -conjugation and introducing substituents with distinctive steric and electronic properties²⁰. Remarkably, WY-1 with the simplest structure has achieved the highest PCE of 14.2% which is comparable to spiro-OMeTAD reference (14.8%), demonstrating the advantage of simple molecular structure in achieving high efficiency PSCs.

Results and Discussion

Synthetic procedures. The synthetic routes of HTMs are depicted in Fig. 1b. The pyrazole intermediates were first obtained by Claisen condensation using the functional ethyl 2-bromobenzoate and acetyl compounds, followed by hydrazine cyclization in refluxing ethanol. Next, treatment of the 2-bromophenyl pyrazoles with tetrahydro-2*H*-pyran (THP) afforded a mixture of two isomers, for which the THP is located at either one of the pyrazolic nitrogen atoms. This mixture was not separated, but directly used for the subsequent reaction with *n*-BuLi to afford the lithiated compounds. Upon treatment with 2,7-bis(bis(4-methoxyphenyl)amino)-9*H*-fluorene-9-one in tetrahydrofuran (THF), followed by hydrolysis in a mixture of acetic acid and concentrated hydrochloric acid at room temperature, the desired spiro compounds with a variety of phenyl pyrazole fragment were obtained in high yields. All obtained compounds WY-1, WY-2 and WY-3 showed good solubility in common

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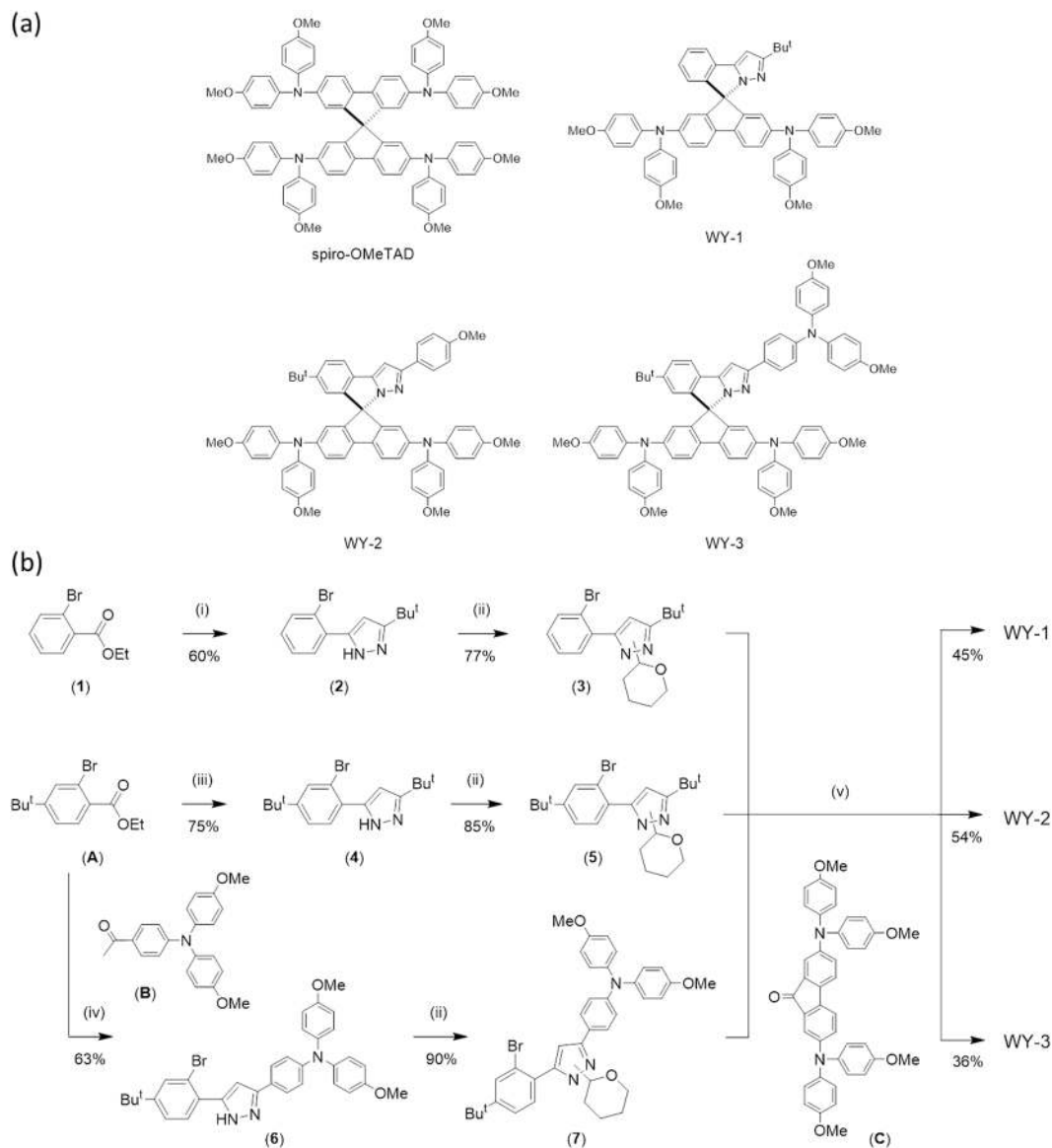


Figure 1. (a) Chemical structures of WY-1, WY-2 and WY-3 compared with Spiro-OMeTAD. (b) Synthetic routes for the studied HTMs.

organic solvents, such as THF, dichloromethane (CH_2Cl_2), toluene and chlorobenzene, ensuring excellent processibility for fabrication of PSCs using spin coating.

Thermal properties. Thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC) measurements were recorded to study their fundamental properties (Table 1 and Figure S1). As can be seen, the decomposition temperature (T_d) of WY-1, WY-2 and WY-3 were recorded to be 401 °C, 422 °C and 427 °C, respectively, which are only slightly lower than that of spiro-OMeTAD (452 °C)³⁵, but are high enough to withstand the harsh condition required for the cell operation. Furthermore, the glass transition temperature (T_g) increased from WY-1 (122 °C), WY-2 (136 °C) to WY-3 (145 °C) due to the higher molecular weight and skeletal rigidity. Nevertheless, all T_g are comparable to that of spiro-OMeTAD (125 °C)³⁵, showing good morphological stability demanded for PSCs.

Optical and electrochemical properties. Figure 2 showed the UV-vis absorption of WY-1, WY-2 and WY-3 in both solution and solid states, together with those of spiro-OMeTAD for comparison, while the corresponding numerical data are depicted in Table 1. The lowest-energy absorption peak in solution occurred at 386 nm for all new HTMs and spiro-OMeTAD, which can be attributed to $\pi-\pi^*$ electronic transition at the diphenylamine substituted fluorene moiety²⁷. Their optical energy gap ($E_{g,\text{opt}}$), estimated from absorption onset, was calculated to be ca. 2.98 eV, which is identical to that of spiro-OMeTAD (3.00 eV) documented in literature³⁶. Going from solution to solid state, their absorption profiles are essentially unchanged, indicating no obvious $\pi-\pi$ stacking interaction in the solid state due to the orthogonal spiro geometry.

HTMs	λ_{abs}^a (nm)	$E_{\text{g,opt}}^b$ (eV)	E_{ox}^c (V)	HOMO ^d (eV)	LUMO ^e (eV)	T_d (°C)	T_g (°C)
Spiro-OMeTAD	303, 365, 386	3.00	0.002	-5.10	-2.10	452 [*]	125 [*]
WY-1	290, 386	2.98	0.065	-5.16	-2.18	401	122
WY-2	269, 386	2.98	0.065	-5.16	-2.18	422	136
WY-3	306, 346, 386	2.98	0.057	-5.15	-2.17	427	145

Table 1. Electrochemical, photophysical and thermal properties of Spiro-OMeTAD, WY-1, WY-2, and WY-3. ^aMeasured in THF with a concentration of 10^{-5} M. ^bCalculated from the absorption onset. ^cThe oxidative onsets (vs FcH/FcH⁺) measured by cyclic voltammetry in CH₂Cl₂. ^dHOMO = $-(E_{\text{ox}} + 5.1)$ eV and LUMO = HOMO + $E_{\text{g,opt}}$.

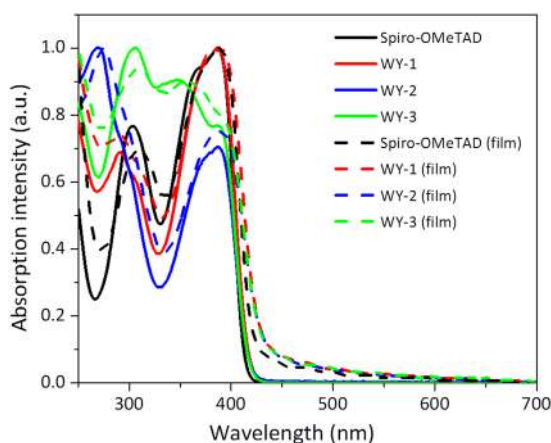


Figure 2. UV-vis absorption spectra of four HTMs measured in solutions (solid line) and thin films (dash line).

Cyclic voltammetry (CV) was next performed to determine the relative energy of the highest occupied molecular orbital (HOMO) (Table 1). As shown in Figure S2, all samples display reversible multi-oxidative behaviours. Particularly, the numbers of reversible CV wave are proportional to the number of diphenylamino substituents, confirming excellent electrochemical stability. In addition, their oxidative onsets versus FcH/FcH⁺ were found to be slightly more positive than spiro-OMeTAD, giving the more stabilized HOMO ($-5.15 \sim -5.16$ eV) compared with spiro-OMeTAD (-5.10 eV), and were expected to have higher V_{OC} values for the PSCs. The stabilized HOMOs may be attributed to the reduced π -conjugation and greater electron withdrawing effect exerted by the orthogonal phenyl pyrazole. This is confirmed by theoretical calculation based on density functional theory (DFT) method using B3LYP hybrid functional and 6-31 g* basis set (Figure S3), for which the electron density of HOMO and LUMO for WY-1, WY-2 and WY-3 are all residing on both the diphenylamino substituents and adjacent fluorene unit. In sharp contrast, the frontier orbitals of spiro-OMeTAD are delocalized over both of the bifluorene cores. Such different electronic density distributions result in reduced HOMO levels for WY-1 to WY-3 which is in accordance with the experimental tendency.

Mobility measurement. The hole mobilities were studied by measuring the electrochemical impedance measurements on a symmetrical, hole-only device comprising of Au/HTM/Au architecture (Fig. 3a) under different voltage bias over 1 MHz to 100 Hz in the dark. A typical Nyquist plot of Au/WY-1/Au measured at -0.4 V bias was shown in Fig. 3b; data points were fitted using an equivalent circuit provided in the insert of Fig. 3b. The equivalent circuit contains a series resistance (R_0), a charge transfer resistance (R_1) and a constant phase element (CPE_1) which associates with charge accumulation in the depletion layer near the interface of metal electrode (gold) and semiconductor (HTM)^{37, 38}. Because the HTM is an organic semiconductor itself; therefore, R_1 and CPE_1 decreases when forward bias increases³⁹. The hole mobility could be estimated using the equation (1):

$$\mu = \frac{eL^2}{k_b T R_1 \text{CPE}_1} \quad (1)$$

where L is the width of the depletion^{40, 41} zone and is approximately equal to the film thickness at such resistive and thin domain, k_b is Boltzmann constant, and T is the absolute temperature. CPE_1 and R_1 were obtained by curve fitting. The relationship of hole mobility with different applied bias is shown in Fig. 3c. It can be seen that WY-1 has identical mobility to spiro-OMeTAD except at high bias over 0.5 V, while the mobilities of WY-2 and WY-3 are both lower than WY-1 and spiro-OMeTAD. The lower mobility for WY-2 and WY-3 may originate from their greater size and higher rigidity³³, both are expected to jeopardize the hole transporting property.

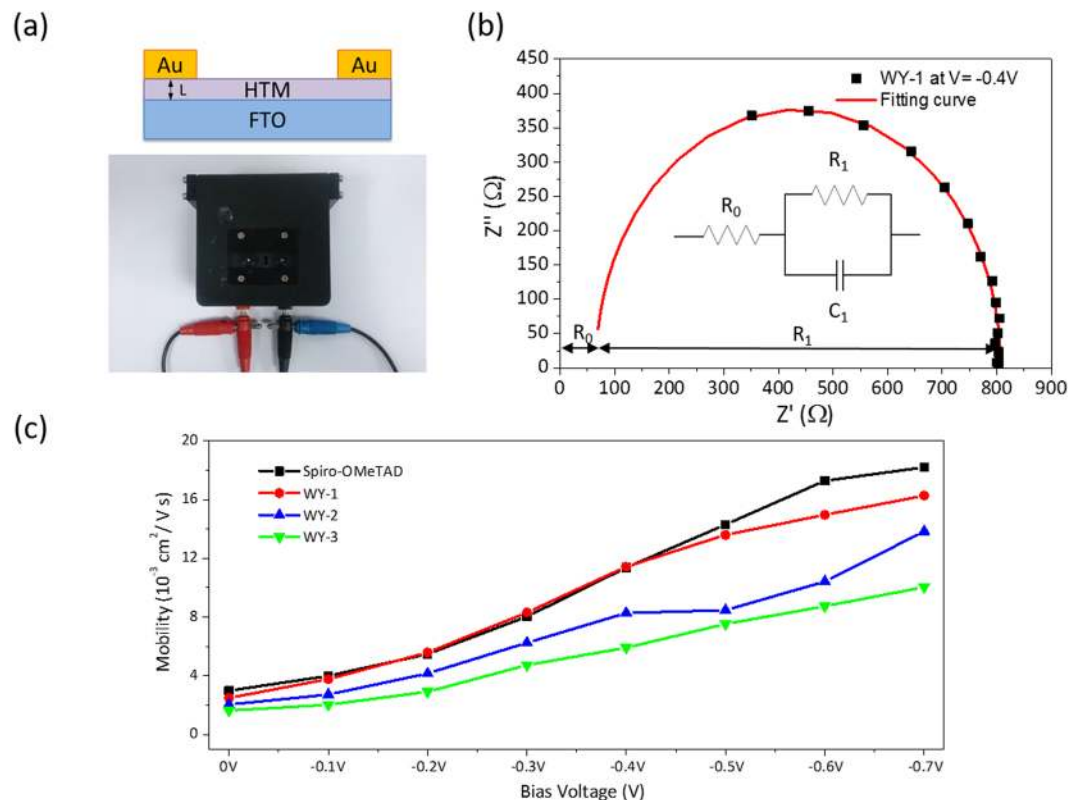


Figure 3. (a) Structural diagram of the hole-only cell Au/HTM/Au for electrochemical impedance measurement. (b) The typical Nyquist plot of Au/WY-1/Au measured at -0.4 V and its fitting curve using an equivalent circuit (inset). (c) The calculated hole mobility of HTMs vs. applied bias.

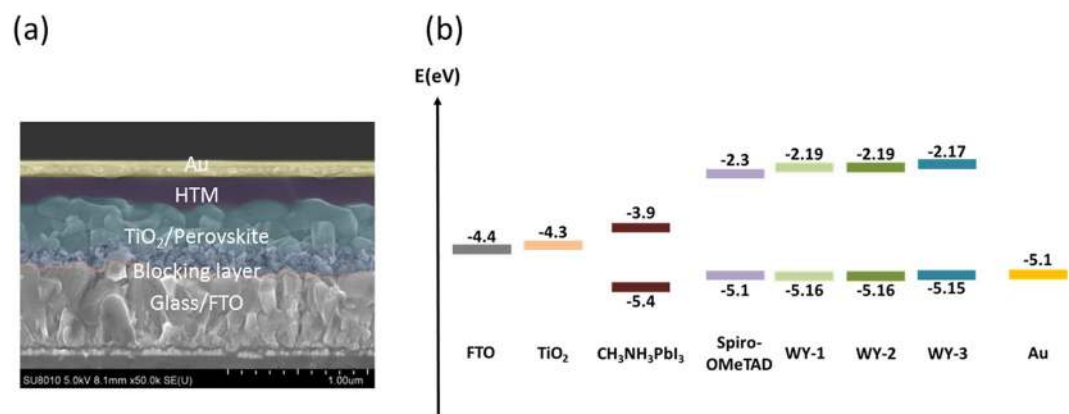


Figure 4. (a) Cross-section view of representative PSC and (b) energy level diagram of the materials used in the studied PSCs (right).

Performance of Perovskite solar cells. The procedures of perovskite solar cell (PSC) fabrication are detailed in the experimental section. The PSC features a multi-layered structure of FTO/TiO₂ compact layer/mesoporous TiO₂ scaffold/perovskite/HTM/Au (Fig. 4a). The perovskite layer (CH₃NH₃PbI₃) was sequentially deposited, following the method developed by Burschka *et al.*⁴² Fig. 4b reveals that the energy levels of new HTMs are suitable for making efficient PSCs. HTMs were spin-coated with typical additives such as bis(trifluoromethane) sulfonimide lithium salt (Li-TFSI) and 4-tert-butylpyridine (tBP) (detailed information can be found in the experimental section).

The photovoltaic performances of PSCs made with all studied HTMs and benchmark spiro-OMeTAD are presented in Fig. 5. Figure 5a–d summarizes photovoltaic performance of multiple devices from different batches using the same recipe and under 100 mW cm^{-2} AM1.5 G solar illumination. It can be seen that WY-1 exhibits competitive PCE vs. spiro-OMeTAD, while WY-2 and WY-3 show lowered short-circuit current density (J_{SC}) and

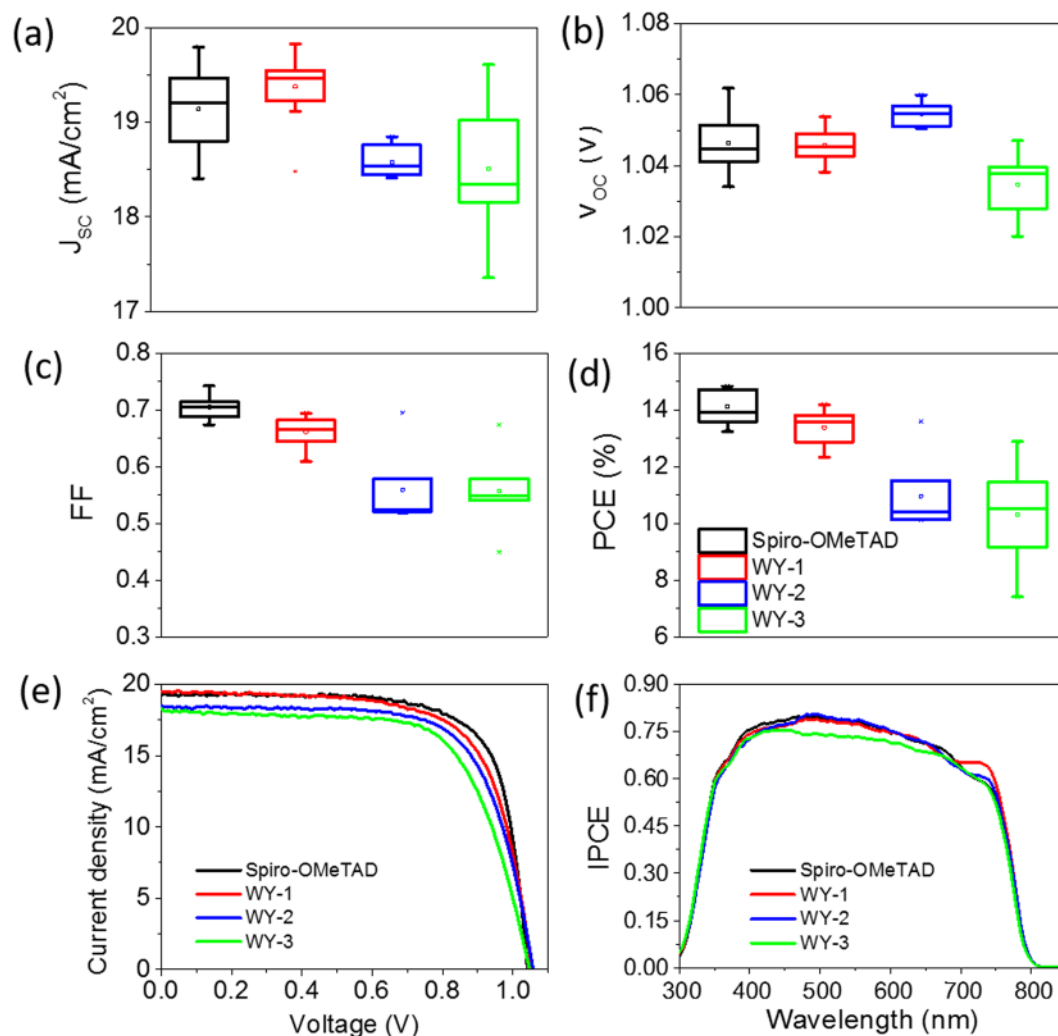


Figure 5. (a–d) Averaged J_{sc} , V_{oc} , FF and PCE of PSCs employed with different HTMs. (e) I–V curves of best-performing PSCs employing the studied HTMs, measured under AM 1.5 G illumination, 100 mW cm^{-2} . (f) IPCE diagrams of the corresponding devices; the total currents derived from the integrated curves are: $18.05 \text{ mA}/\text{cm}^2$, $18.16 \text{ mA}/\text{cm}^2$, $18.08 \text{ mA}/\text{cm}^2$ and $17.32 \text{ mA}/\text{cm}^2$ for the spiro-OMeTAD, WY-1, WY-2, and WY-3-based device, respectively.

fill factor (FF), resulting in slightly inferior PCE. Figure 5e depicts the I–V curves of the best devices based on new HTMs and spiro-OMeTAD, while corresponding I–V parameters are listed in Table 2. The PSC using WY-1 achieves a J_{sc} of $19.48 \text{ mA}/\text{cm}^2$, an open-circuit voltage (V_{oc}) of 1.05 V , a fill factor (FF) of 0.69 , and a PCE of 14.20% , respectively. Under the same fabrication conditions, the benchmark spiro-OMeTAD-based device exhibits J_{sc} of $19.20 \text{ mA}/\text{cm}^2$, V_{oc} of 1.04 V , FF of 0.74 and PCE of 14.84% . From the incident photon to current efficiency (IPCE) provided in Fig. 5f, the higher J_{sc} obtained in WY-1-based device is derived from its higher IPCE at 700 to 750 nm , which may be due to the superior optical transmittance of WY-1 at around 700 nm (Figure S4), allowing better IPCE over that of spiro-OMeTAD. But owing to the lowered hole mobility vs. spiro-OMeTAD, WY-1 device suffers from notable ohmic losses in FF due to the slightly higher resistivity. In addition, WY-2 and WY-3 based PSCs exhibit very similar photovoltaic performance with J_{sc} of 18.50 and 18.27 mA cm^{-2} , V_{oc} of 1.06 and 1.05 V , FF of 0.70 and 0.67 and PCE of 13.60 and 12.89% , respectively. The inferior performance of WY-2 and WY-3 than WY-1 may be attributed to the larger molecular volume vs. WY-1. Hence, the greater steric hindrance in WY-2 and WY-3 renders poor intermolecular π -stacking interaction (i.e. carrier transport) in the deposited thin film and thus lowers the J_{sc} and FF of the PSC.

Conclusions

In summary, we reported three HTMs with asymmetric spiro-phenylpyrazole/fluorene architecture, which renders the effective modification on conventional symmetric spiro-OMeTAD. These new HTMs possess suitable HOMO level which could be beneficial to the high V_{oc} . Moreover, they exhibit adequate thermal, optical and electrochemical properties but a slightly different hole-transporting ability, while WY-1 shows the best hole mobility relative to that of WY-2 and WY-3. As a result, WY-1 based device exhibits the highest PCE of

Type of BL		J _{sc} (mA/cm ²)	V _{oc} (mV)	FF	PCE (%)	R _s (Ω)	HI
Best performing	Spiro-OMeTAD	19.20	1.041	0.742	14.84	32.47	0.04
	WY-1	19.48	1.053	0.691	14.19	53.52	0.07
	WY-2	18.50	1.057	0.696	13.60	63.10	0.08
	WY-3	18.27	1.047	0.674	12.89	85.45	0.05
Average*	Spiro-OMeTAD	19.14 ± 0.46	1.046 ± 0.007	0.705 ± 0.022	14.13 ± 0.57	47.96 ± 11.86	0.09 ± 0.07
	WY-1	19.38 ± 0.33	1.046 ± 0.005	0.662 ± 0.025	13.36 ± 0.58	51.64 ± 13.09	0.09 ± 0.04
	WY-2	18.58 ± 0.15	1.055 ± 0.003	0.560 ± 0.059	10.96 ± 1.17	89.89 ± 12.57	0.39 ± 0.13
	WY-3	18.50 ± 0.65	1.035 ± 0.003	0.556 ± 0.061	10.32 ± 1.72	100.10 ± 34.37	0.35 ± 0.16

Table 2. Photovoltaic parameters of PSCs with different hole transporting layer. Photomasks (0.2 × 0.5 cm²) made of thin metal sheet were applied before all measurements. The hysteresis index (HI) is defined as $(P_{\max(\text{forward scan})}/P_{\max(\text{backward scan})}) - 1$, which P_{\max} is the maximum power in I-V measurement. *These values were obtained from 6–8 devices.

14.20% with a J_{sc} of 19.48 mA/cm², a V_{oc} of 1.05 V and a FF of 0.69 which are very close to the benchmark HTM spiro-OMeTAD (14.84%). On the other hand, higher series resistances originate from lowered hole mobility are found in these WY-based PSCs, rendering lowered J_{sc} and FF. Currently, modification on molecular structure and optimization on concentration of additives are undergoing in our laboratory to solve this issue. We believe that further modulation on such spiro-phenylpyrazole/fluorene structure will push the device performance to an even higher level.

Methods

General Information and Materials. All reactions were performed under a nitrogen atmosphere and solvents were distilled from appropriate drying agents prior to use. Commercially available reagents were used without purification unless otherwise stated. The starting materials (A)⁴³, (B)⁴⁴ and (C)³³ were prepared according to the literature methods.

Synthetic procedures. Experimental conditions: (i) 3,3-dimethylbutan-2-one, NaH, THF, reflux; then N₂H₄·H₂O, EtOH, reflux; (ii) 3,4-dihydro-2H-pyran, CH₂Cl₂, reflux; (iii) 1-(4-methoxyphenyl)ethanone, NaH, THF, reflux; then N₂H₄·H₂O, EtOH, reflux; (iv) NaH, THF, reflux; then N₂H₄·H₂O, EtOH, reflux; (v) *n*-BuLi, THF, −78 °C to RT; then HCl, AcOH, reflux.

3-Tert-butyl-5-(2-bromophenyl)-1H-pyrazole (2). NaH (478 mg, 19 mmol) was suspended in 16 mL of dry THF under nitrogen, and was added dropwise 3,3-dimethylbutan-2-one (1.27 g, 12.6 mmol) dissolved in 8 mL dry THF at 0 °C. The mixture was stirred for 1 h at RT, and ethyl 2-bromobenzoate (**1**, 4.38 g, 18.7 mmol) dissolved in 16 mL dry THF was added dropwise at 0 °C. The mixture was first refluxed for 4 h, and it was poured into a mixture of water and ethyl acetate and neutralized with 2 M HCl_(aq). The organic layer was washed with brine and water, and dried over anhydrous Na₂SO₄. The crude product was obtained by concentrating the solution to dryness. Then, this product was dissolved in EtOH (65 mL), and reacted with hydrazine hydrate (3 mL, 63 mmol) in presence of *p*-toluenesulfonic acid monohydrate (228 mg, 1.2 mmol) at reflux for 12 h. After then, the solution was evaporated to dryness and the residue dissolved in ethyl acetate. The organic layer was washed with NaHCO₃ solution, brine and water in sequence, and dried over anhydrous Na₂SO₄. The pure product can be obtained by column chromatography eluting with a mixture of hexane: EA = 4:1. Overall yield: 60%. ¹H NMR (400 MHz, CDCl₃): δ 7.63 (d, *J* = 8.0 Hz, 1 H), 7.59 (d, *J* = 7.8 Hz, 1 H), 7.31 (t, *J* = 7.5 Hz, 1 H), 7.16 (t, *J* = 8.4 Hz, 1 H), 6.48 (s, 1 H), 1.33 (s, 9 H).

3-Tert-butyl-5-(2-bromophenyl)-1-(tetrahydro-2H-pyran-2-yl)-1H-pyrazole (3). A mixture of compound **2** (2.09 g, 7.5 mmol), 3,4-dihydro-2H-pyran (2.52 g, 30 mmol), *p*-toluenesulfonic acid monohydrate (140 mg, 0.75 mmol) and CH₂Cl₂ (50 mL) was refluxed overnight under nitrogen. The mixture was washed with NaHCO₃, brine and water and dried over anhydrous Na₂SO₄. The solid was next purified by column chromatography using hexane: EA = 20:1 as the eluent. Yield: 77%. ¹H NMR (400 MHz, CDCl₃): δ 7.65 (dd, *J* = 7.8, 1.1 Hz, 1 H), 7.43 (dd, *J* = 7.6, 1.7 Hz, 1 H), 7.35 (td, *J* = 7.5, 1.3 Hz, 1 H), 7.27–7.22 (m, 1 H), 6.16 (s, 1 H), 4.93–4.87 (m, 1 H), 4.00–3.96 (m, 1 H), 3.39–3.33 (m, 1 H), 2.46–2.40 (m, 1 H), 2.03–1.99 (m, 1 H), 1.88–1.83 (m, 2 H), 1.67–1.64 (m, 1 H), 1.45–1.41 (m, 1 H), 1.33 (s, 9 H).

5-(4-Tert-butyl-2-bromophenyl)-3-(4-methoxyphenyl)-1H-pyrazole (4). Following the same procedure described for **2**, treatment of **A** and dimethylbutan-2-one afforded a white product **4** in 75% yield. ¹H NMR (400 MHz, CDCl₃): δ 7.69 (d, *J* = 8.8 Hz, 2 H), 7.66 (d, *J* = 1.9 Hz, 1 H), 7.52 (d, *J* = 8.1 Hz, 1 H), 7.38 (dd, *J* = 8.2, 1.9 Hz, 1 H), 6.95 (d, *J* = 8.8 Hz, 2 H), 6.82 (s, 1 H), 3.84 (s, 3 H), 1.33 (s, 9 H).

5-(4-Tert-butyl-2-bromophenyl)-1-(tetrahydro-2H-pyran-2-yl)-3-(4-methoxyphenyl)-1H-pyrazole (5). Following the procedure described for **3**, combination of **4** with 3,4-dihydro-2H-pyran afforded a white product **5** in 85% yield. ¹H NMR (400 MHz, CDCl₃): δ 7.80 (d, *J* = 8.9 Hz, 2 H), 7.68–7.67 (m, 1 H), 7.39 (d, *J* = 1.1 Hz, 2 H), 6.91 (d, *J* = 8.9 Hz, 2 H), 6.53 (s, 1 H), 4.98–4.96 (m, 1 H), 4.04–4.01 (m, 1 H), 3.82 (s, 3 H),

3.46–3.40 (m, 1 H), 2.59–2.57 (m, 1 H), 2.10–2.05 (m, 1 H), 1.97–1.93 (m, 1 H), 1.74–1.68 (m, 2 H), 1.63–1.60 (m, 1 H), 1.35 (s, 9 H). MS [FAB], m/z 470.2, M^+ .

N-(4-(5-(4-*Tert*-butyl-2-bromophenyl)-1*H*-pyrazol-3-yl)phenyl)-4-methoxy-*N*-(4-methoxyphenyl)benzenamine (6). Following the procedure described for 2, condensation of compounds A and B afforded a white product 6 in 63% yield. ^1H NMR (400 MHz, acetone- d_6): δ 7.71 (d, $J = 1.9$ Hz, 1 H), 7.68–7.64 (m, 3 H), 7.49 (dd, $J = 8.2$, 2.0 Hz, 1 H), 7.07–6.90 (m, 11 H), 3.78 (s, 6 H), 1.34 (s, 9 H).

N-(4-(5-(4-*Tert*-butyl-2-bromophenyl)-1-(tetrahydro-2*H*-pyran-2-yl)-1*H*-pyrazol-3-yl)phenyl)-4-methoxy-*N*-(4-methoxyphenyl)benzenamine (7). Following the procedure described for 3, treatment of 6 with 3,4-dihydro-2*H*-pyran afforded a white product 7 in 90% yield. ^1H NMR (400 MHz, CDCl_3): δ 7.66 (d, $J = 7.5$ Hz, 1 H), 7.50–7.46 (m, 3 H), 7.39–7.33 (m, 1 H), 7.11 (d, $J = 8.7$ Hz, 1 H), 7.05–7.03 (m, 3 H), 6.94 (d, $J = 7.5$ Hz, 2 H), 6.86–6.79 (m, 4 H), 6.50 (s, 1 H), 5.23–5.20 (m, 1 H), 4.17–4.13 (m, 1 H), 3.78 (s, 6 H), 3.64–3.60 (m, 1 H), 2.69–2.62 (m, 1 H), 2.08–2.04 (m, 1 H), 1.84–1.72 (m, 4 H). MS [FAB], m/z 667.3, M^+ .

Compound WY-1. A solution of compound 3 (433 mg, 1.22 mmol) in dry THF (5 mL) was treated with *n*-BuLi (731 μL , 2.5 M in *n*-hexane) under nitrogen at -78°C . After 30 min, a solution of compound C (551 mg, 0.87 mmol) in THF (5 mL) was added dropwise. The mixture was stirred for 30 min at -78°C , and allowed to warm up to RT. After 12 h, the solution was concentrated and the residue was extracted with CH_2Cl_2 and washed with brine and water in sequence and finally dried over anhydrous Na_2SO_4 . The alcohol intermediate was obtained by column chromatography using hexane: EA = 4:1 as the eluent. It was next added to a mixture of concentrated HCl solution (1.5 mL) and acetic acid (20 mL). After stirring for 1 h at RT, the mixture was quenched with ice water and neutralized with $\text{NaHCO}_3(\text{aq})$. The crude product was extracted with CH_2Cl_2 and washed with water. The organic layer was dried over anhydrous Na_2SO_4 and purified by silica gel column chromatography eluting with a mixture of hexane: EA = 8:1, giving a pale green product in 45% yield. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 7.59–7.56 (m, 3 H), 7.34 (t, $J = 7.6$ Hz, 1 H), 7.17 (t, $J = 7.6$ Hz, 1 H), 6.81–6.72 (m, 19 H), 6.33 (s, 1 H), 6.00 (s, 2 H), 3.64 (s, 12 H), 1.19 (s, 9 H). ^{13}C NMR (101 MHz, acetone): δ 167.06, 155.87, 148.04, 147.87, 146.08, 145.74, 140.52, 134.01, 130.64, 128.46, 127.55, 126.15, 122.62, 121.45, 120.22, 120.20, 115.65, 114.56, 92.99, 76.33, 54.70, 32.60, 30.21. MS [FAB], m/z 816.4, M^+ . Anal. Calcd. for $\text{C}_{54}\text{H}_{48}\text{N}_4\text{O}_4$: C, 79.39; H, 5.92; N, 6.86. Found: C, 78.52; H, 5.90; N, 6.16.

Compound WY-2. Following the procedure described for WY-1, treatment of C with 5 afforded a pale yellow product in 54% yield. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 7.65–7.56 (m, 5 H), 7.44 (d, $J = 8.0$ Hz, 1 H), 6.92 (d, $J = 12.9$ Hz, 2 H), 6.79–6.74 (m, 12 H), 6.66–6.64 (m, 9 H), 6.10 (s, 1 H), 3.74 (s, 3 H), 3.60 (s, 12 H), 1.18 (s, 9 H). ^{13}C NMR (101 MHz, acetone): δ 159.44, 156.11, 155.90, 151.48, 148.07, 147.57, 147.10, 145.96, 140.53, 134.26, 127.92, 127.10, 126.53, 125.96, 125.56, 124.82, 121.74, 120.31, 120.04, 119.63, 116.36, 114.49, 113.80, 93.04, 76.63, 54.68, 34.74, 30.57. MS [FAB], m/z 922.5, M^+ . Anal. Calcd. for $\text{C}_{61}\text{H}_{54}\text{N}_4\text{O}_5$: C, 79.37; H, 5.90; N, 6.07. Found: C, 78.66; H, 5.91; N, 6.14.

Compound WY-3. Following the procedure described for WY-1, treatment of C with 7 afforded a pale yellow product in 36% yield. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 7.58 (d, $J = 8.3$ Hz, 2 H), 7.56–7.51 (m, 3 H), 7.42 (dd, $J = 8.1$, 1.6 Hz, 1 H), 6.97–6.96 (m, 4 H), 6.88–6.85 (m, 4 H), 6.78–6.72 (m, 15 H), 6.66–6.63 (m, 8 H), 6.09 (s, 1 H), 3.70 (s, 6 H), 3.60 (s, 12 H), 1.16 (s, 9 H). ^{13}C NMR (101 MHz, acetone): δ 156.22, 156.03, 155.83, 151.35, 148.21, 148.10, 147.60, 147.08, 146.13, 140.74, 140.57, 134.32, 127.93, 126.68, 126.45, 126.04, 125.93, 125.67, 121.98, 120.34, 120.21, 120.18, 119.49, 116.46, 114.68, 114.51, 93.06, 76.82, 54.82, 54.71, 34.66, 30.78. MS [FAB], m/z 1119.5, M^+ . Anal. Calcd. for $\text{C}_{74}\text{H}_{65}\text{N}_5\text{O}_6$: C, 79.33; H, 6.25; N, 5.85. Found: C, 78.50; H, 6.24; N, 5.89.

Fabrication of perovskite solar cells. FTO glass substrates (2.2 mm, $8 \Omega/\square$, NSG, Japan) were etched by a laser engraver (LMF-020F, Taiwan) to obtain the required electrode patterns. The etched sheets were then cleaned by commercial detergent solution (PK-LCG46, USA) and DI-water in sequence under ultrasonic bath for 30 minutes. The electrodeposited TiO_2 blocking layer (ED-BL) was anodically electrodeposited on clean FTO in an aqueous solution containing 250 mM TiCl_4 (20% in 2 N HCl, Acros Organics) for 300 seconds⁴⁵. Subsequently, 200 nm-thick mesoporous TiO_2 layer was deposited by spin-coating commercial TiO_2 paste (30 nm, G24 power Ltd. United Kingdom) diluted in anhydrous ethanol by a 1:5 weight ratio at 6000 rpm for 30 seconds, followed by heating at 120°C and 500°C for 5 minutes and 30 minutes, respectively. The resultant film was treated in a 40 mM TiCl_4 aqueous solution at 70°C for 30 minutes and followed by annealing at 450°C for 30 minutes. After cooling down to RT, the perovskite layer was applied following the published sequential deposition method⁴². In brief, 1.2 M PbI_2 (99.999%, Sigma-Aldrich) was dissolved in a mixture of DMF (*N,N*-dimethylformamide, 99.8%, Merck) and DMSO (Dimethyl sulfoxide, 99.9%, J. T. Baker). Next, the ED-BL coated TiO_2 scaffold was infiltrated by PbI_2 solution by spin-coating at 6000 rpm for 10 seconds and followed by heating at 70°C for 30 minutes. The substrates were immersed into a solution containing 63 mM methylamine iodide (MAI)⁴⁶ in 2-propanol for 1 minute, then dried and baked on 70°C for 20 minutes. Finally, the hole-transporting layer was deposited on top of the perovskite layer by spin-coating at 3000 rpm for 30 seconds using of HTM solution, which contained 75 mM of the studied HTM, 20 mM of bis(trifluoromethane)sulfonimide lithium salt (Li-TFSI, Sigma-Aldrich) and 120 mM of 4-*tert*-butylpyridine (TBP, 96%, Sigma-Aldrich) in chlorobenzene. After storing the device at dry atmosphere overnight, a gold metal film was thermally evaporated onto HTM layer as the back electrode.

Characterization and measurement. The ^1H NMR spectra were measured with a Varian Mercury-400 instrument. Elemental analysis was carried out on a Heraeus CHN-O Rapid Elementary Analyzer. Mass spectra

were recorded on a JEOL SX-102A instrument operating in electron impact (EI) or fast atom bombardment (FAB) mode. The thermogravimetric analysis and differential scanning calorimetry were measured on a Seiko SSC 5000 instrument. UV-Vis spectra were recorded on a Hitachi U-3900 spectrophotometer. Cyclic voltammetry was conducted on a CHI 621 A Electrochemical Analyzer with a conventional three-electrode system consisting of a platinum working electrode, a platinum counter electrode and an Ag/AgCl reference electrode, and using 0.1 M tetrabutylammonium hexafluorophosphate (TBAPF₆) as the supporting electrolyte. All potentials were calibrated against the ferrocene/ferrocenium couple assuming $FcH/FcH^+ = -5.1 \text{ eV}^{47}$. Cross-sectional view of PSC was investigated using a Hitachi SU-8010 field emission scanning electron microscope. The conductivity of HTM was investigated by a computer-controlled Autolab PGSTAT30 Potentiostat/Galvanostat. The I-V characteristic was measured using a Keithley 2400 digital source meter under AM1.5 G illumination at intensity of 100 mW/cm² (Peccell Technologies, PEC-L15). A KG3 monocrystalline silicon photodiode (Oriel, USA) was used to calibrate the light intensity. A 0.2 × 0.5 cm² photo-mask was attached to the front side of the PSC to accurately control the illuminating area. The IPCE was measured using monochromatic light illumination (Peccell Technologies, PEC-20).

References

- Kojima, A., Teshima, K., Shirai, Y. & Miyasaka, T. Organometal Halide Perovskites as Visible-Light Sensitizers for Photovoltaic Cells. *J. Am. Chem. Soc.* **131**, 6050–6051 (2009).
- De Wolf, S. *et al.* Organometallic Halide Perovskites: Sharp Optical Absorption Edge and Its Relation to Photovoltaic Performance. *J. Phys. Chem. Lett.* **5**, 1035–1039 (2014).
- Wang, Y., Zhang, Y., Zhang, P. & Zhang, W. High intrinsic carrier mobility and photon absorption in the perovskite CH₃NH₃PbI₃. *Phys. Chem. Chem. Phys.* **17**, 11516–11520 (2015).
- Wehrenfennig, C., Eperon, G. E., Johnston, M. B., Snaith, H. J. & Herz, L. M. High Charge Carrier Mobilities and Lifetimes in Organolead Trihalide Perovskites. *Adv. Mater.* **26**, 1584–1589 (2014).
- Motta, C., El-Mellouhi, F. & Sanvito, S. Charge carrier mobility in hybrid halide perovskites. *Sci. Rep.* **5**, 12746 (2015).
- Stranks, S. D. *et al.* Electron-Hole Diffusion Lengths Exceeding 1 Micrometer in an Organometal Trihalide Perovskite Absorber. *Science* **342**, 341–344 (2013).
- Xing, G. *et al.* Long-Range Balanced Electron- and Hole-Transport Lengths in Organic-Inorganic CH₃NH₃PbI₃. *Science* **342**, 344–347 (2013).
- Yang, W. S. *et al.* High-performance photovoltaic perovskite layers fabricated through intramolecular exchange. *Science* **348**, 1234–1237 (2015).
- Bi, D. *et al.* Efficient luminescent solar cells based on tailored mixed-cation perovskites. *Sci. Adv.* **2**, 1501170 (2016).
- Saliba, M. *et al.* Cesium-containing triple cation perovskite solar cells: improved stability, reproducibility and high efficiency. *Energy Environ. Sci.* **9**, 1989–1997 (2016).
- Snaith, H. J. & Grätzel, M. Electron and Hole Transport through Mesoporous TiO₂ Infiltrated with Spiro-MeOTAD. *Adv. Mater.* **19**, 3643–3647 (2007).
- Gong, G. *et al.* Dopant-free 3,3'-bithiophene derivatives as hole transport materials for perovskite solar cells. *J. Mater. Chem. A* **4**, 3661–3666 (2016).
- Swetha, T. & Singh, S. P. Perovskite solar cells based on small molecule hole transporting materials. *J. Mater. Chem. A* **3**, 18329–18344 (2015).
- Yu, Z. & Sun, L. Recent Progress on Hole-Transporting Materials for Emerging Organometal Halide Perovskite Solar Cells. *Adv. Energy Mater.* **5**, 1500213 (2015).
- Ameen, S. *et al.* Perovskite Solar Cells: Influence of Hole Transporting Materials on Power Conversion Efficiency. *Chem. Sus. Chem.* **9**, 10–27 (2016).
- Tong, X., Lin, F., Wu, J. & Wang, Z. M. High Performance Perovskite Solar Cells. *Adv. Sci.* **3**, 1500201 (2016).
- Jeon, N. J. *et al.* Efficient Inorganic–Organic Hybrid Perovskite Solar Cells Based on Pyrene Arylamine Derivatives as Hole-Transporting Materials. *J. Am. Chem. Soc.* **135**, 19087–19090 (2013).
- Nishimura, H. *et al.* Hole-Transporting Materials with a Two-Dimensionally Expanded π -System around an Azulene Core for Efficient Perovskite Solar Cells. *J. Am. Chem. Soc.* **137**, 15656–15659 (2015).
- Wu, Y. *et al.* Highly compact TiO₂ layer for efficient hole-blocking in perovskite solar cells. *Appl. Phys. Express* **7**, 052301 (2014).
- Molina-Ontoria, A. *et al.* Benzotrithiophene-Based Hole-Transporting Materials for 18.2% Perovskite Solar Cells. *Angew. Chem., Int. Ed.* **55**, 6270–6274 (2016).
- Zimmermann, I. *et al.* High-Efficiency Perovskite Solar Cells Using Molecularly Engineered, Thiophene-Rich, Hole-Transporting Materials: Influence of Alkyl Chain Length on Power Conversion Efficiency. *Adv. Energy Mater.* **7**, 1601674 (2017).
- Choi, H., Do, K., Park, S., Yu, J.-S. & Ko, J. Efficient Hole Transporting Materials with Two or Four N,N-Di(4-methoxyphenyl) aminophenyl Arms on an Ethene Unit for Perovskite Solar Cells. *Chem. Eur. J.* **21**, 15919–15923 (2015).
- Wu, F., Liu, J., Wang, G., Song, Q. & Zhu, L. m-Methoxy Substituents in a Tetraphenylethylene-Based Hole-Transport Material for Efficient Perovskite Solar Cells. *Chem. Eur. J.* **22**, 16636–16641 (2016).
- Rakstys, K. *et al.* Highly Efficient Perovskite Solar Cells Employing an Easily Attainable Bifluorenylidene-Based Hole-Transporting Material. *Angew. Chem., Int. Ed.* **55**, 7464–7468 (2016).
- Zhao, X. *et al.* A novel one-step synthesized and dopant-free hole transport material for efficient and stable perovskite solar cells. *J. Mater. Chem. A* **4**, 16330–16334 (2016).
- Zhang, J. *et al.* Constructive Effects of Alkyl Chains: A Strategy to Design Simple and Non-Spiro Hole Transporting Materials for High-Efficiency Mixed-Ion Perovskite Solar Cells. *Adv. Energy Mater.* **6**, 1502536 (2016).
- Yan, Q., Guo, Y., Ichimura, A., Tsuji, H. & Nakamura, E. Three-Dimensionally Homoconjugated Carbon-Bridged Oligophenylenevinylene for Perovskite Solar Cells. *J. Am. Chem. Soc.* **138**, 10897–10904 (2016).
- Park, S. *et al.* A [2,2]paracyclophane triarylamine-based hole-transporting material for high performance perovskite solar cells. *J. Mater. Chem. A* **3**, 24215–24220 (2015).
- Zhang, J. *et al.* The Role of 3D Molecular Structural Control in New Hole Transport Materials Outperforming Spiro-OMeTAD in Perovskite Solar Cells. *Adv. Energy Mater.* **6**, 1601062 (2016).
- Shi, Y. *et al.* Two methoxyaniline-substituted dibenzofuran derivatives as hole-transport materials for perovskite solar cells. *J. Mater. Chem. A* **4**, 5415–5422 (2016).
- Li, Z. A. *et al.* Rational Design of Dipolar Chromophore as an Efficient Dopant-Free Hole-Transporting Material for Perovskite Solar Cells. *J. Am. Chem. Soc.* **138**, 11833–11839 (2016).
- Wang, Y.-K. *et al.* Dopant-Free Spiro-Triphenylamine/Fluorene as Hole-Transporting Material for Perovskite Solar Cells with Enhanced Efficiency and Stability. *Adv. Funct. Mater.* **26**, 1375–1381 (2016).
- Li, M.-H. *et al.* Novel spiro-based hole transporting materials for efficient perovskite solar cells. *Chem. Commun.* **51**, 15518–15521 (2015).

34. Xu, B. *et al.* A low-cost spiro[fluorene-9,9[prime or minute]-xanthene]-based hole transport material for highly efficient solid-state dye-sensitized solar cells and perovskite solar cells. *Energy Environ. Sci.* **9**, 873–877 (2016).
35. Maciejczyk, M., Ivaturi, A. & Robertson, N. SFX as a low-cost 'Spiro' hole-transport material for efficient perovskite solar cells. *J. Mater. Chem. A* **4**, 4855–4863 (2016).
36. Murray, A. T. *et al.* Modular design of SPIRO-OMeTAD analogues as hole transport materials in solar cells. *Chem. Commun.* **51**, 8935–8938 (2015).
37. Otalora, C. A., Loaiza, A. F. & Gordillo, G. Study of Electrical Transport Properties of Thin Films Used as HTL and as Active Layer in Organic Solar Cells, through Impedance Spectroscopy Measurements. *Adv. Mater. Sci. Eng.* **2016**, 7 (2016).
38. Gaur, A. & Kumar, P. An improved circuit model for polymer solar cells. *Prog. Photovoltaics Res. Appl.* **22**, 937–948 (2014).
39. Liu, Y. *et al.* A dopant-free organic hole transport material for efficient planar heterojunction perovskite solar cells. *J. Mater. Chem. A* **3**, 11940–11947 (2015).
40. Garcia-Belmonte, G. *et al.* Charge carrier mobility and lifetime of organic bulk heterojunctions analyzed by impedance spectroscopy. *Org. Electron.* **9**, 847–851 (2008).
41. Garcia-Belmonte, G., Bisquert, J. & Popkurov, G. S. Determination of the electronic conductivity of polybithiophene films at different doping levels using *in situ* electrochemical impedance measurements. *Appl. Phys. Lett.* **83**, 2178–2180 (2003).
42. Burschka, J. *et al.* Sequential deposition as a route to high-performance perovskite-sensitized solar cells. *Nature* **499**, 316–319 (2013).
43. Dressler, J. J., Miller, S. A., Meeuwse, B. T., Riel, A. M. S. & Dahl, B. J. Synthesis of dilactone bridged terphenyls with crankshaft architectures. *Tetrahedron* **71**, 283–292 (2015).
44. Newton, G. N. *et al.* Lability-Controlled Syntheses of Heterometallic Clusters. *Angew. Chem., Int. Ed.* **53**, 2941–2944 (2014).
45. Su, T.-S., Hsieh, T.-Y., Hong, C.-Y. & Wei, T.-C. Electrodeposited Ultrathin TiO₂ Blocking Layers for Efficient Perovskite Solar Cells. *Sci. Rep.* **5**, 16098 (2015).
46. Hsieh, T.-Y., Wei, T.-C., Wu, K.-L., Ikegami, M. & Miyasaka, T. Efficient perovskite solar cells fabricated using an aqueous lead nitrate precursor. *Chem. Commun.* **51**, 13294–13297 (2015).
47. Xu, B. *et al.* 1,1,2,2-Tetrachloroethane (TeCA) as a Solvent Additive for Organic Hole Transport Materials and Its Application in Highly Efficient Solid-State Dye-Sensitized Solar Cells. *Adv. Energy Mater.* **5**, 1402340 (2015).

Acknowledgements

This work was supported by Ministry of Science and Technology, Taiwan, under the grant numbers MOST 105-2119-M-007-005 and 105-2119-M-007-001.

Author Contributions

Y. Chi and T.C. Wei developed the concept. Y. Wang synthesized and characterized the compound. T.S. Su carried out hole mobility measurement and device fabrication. H.Y. Tsai conducted NMR characterization. Y. Wang, T.S. Su and T.C. Wei drafted the article and Y. Chi polished the content to the present form. All authors reviewed the paper.

Additional Information

Supplementary information accompanies this paper at doi:10.1038/s41598-017-08187-4

Competing Interests: The authors declare that they have no competing interests.

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