



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

[1,2,5]Thiadiazolo[3,4-d]Pyridazine as an Internal Acceptor in the D-A--A Organic Sensitizers for Dye-Sensitized Solar Cells

Citation for published version:

Chmovzh, TN, Knyazeva, EA, Tanaka, E, Popov, VV, Mikhalchenko, L, Robertson, N & Rakitin, OA 2019, '[1,2,5]Thiadiazolo[3,4-d]Pyridazine as an Internal Acceptor in the D-A--A Organic Sensitizers for Dye-Sensitized Solar Cells', *Molecules*, vol. 24, no. 8, pp. 1588. <https://doi.org/10.3390/molecules24081588>

Digital Object Identifier (DOI):

[10.3390/molecules24081588](https://doi.org/10.3390/molecules24081588)

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

Published In:

Molecules

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



Article

[1,2,5]Thiadiazolo[3,4-*d*]Pyridazine as an Internal Acceptor in the D-A- π -A Organic Sensitizers for Dye-Sensitized Solar Cells

Timofey N. Chmovzh¹, Ekaterina A. Knyazeva^{1,2}, Ellie Tanaka³, Vadim V. Popov²,
Ludmila V. Mikhailchenko¹, Neil Robertson^{3,*} and Oleg A. Rakitin^{1,2,*}

¹ N. D. Zelinsky Institute of Organic Chemistry, Russian Academy of Sciences, 119991 Moscow, Russia; orakitin@ioc.ac.ru (O.A.R.); tim1661@yandex.ru (T.N.C.); katerina_knyazev@ioc.ac.ru (E.A.K.); mlv@ioc.ac.ru (L.V.M.)

² Nanotechnology Education and Research Center, South Ural State University, 454080 Chelyabinsk, Russia; popov.ioc@gmail.com

³ EaStCHEM School of Chemistry, University of Edinburgh, Edinburgh EH9 3FJ, UK; Ellie.Tanaka@ed.ac.uk

* Correspondence: Correspondence: Neil.Robertson@ed.ac.uk (N.R.); orakitin@ioc.ac.ru (O.A.R.); Tel.: +44-131-650-4755 (N.R.); +7-499-135-5327(O.A.R.)

Academic Editor: Panayiotis A. Koutentis

Received: 4 April 2019; Accepted: 19 April 2019; Published: 22 April 2019

Abstract: Four new D-A- π -A metal-free organic sensitizers for dye-sensitized solar cells (DSSCs), with [1,2,5]thiadiazolo[3,4-*d*]pyridazine as internal acceptor, thiophene unit as π -spacer and cyanoacrylate as anchoring electron acceptor, have been synthesized. The donor moiety was introduced into [1,2,5]thiadiazolo[3,4-*d*]pyridazine by nucleophilic aromatic substitution and Suzuki cross-coupling reactions, allowing design of D-A- π -A sensitizers with the donor attached to the internal heterocyclic acceptor not only by the carbon atom, as it is in a majority of DSSCs, but by the nitrogen atom also. Although low values of power conversion efficiency (PCE) were found, a few important consequences were identified: (i) poor PCE data can be attributed to high electron deficiency of the internal [1,2,5]thiadiazolo[3,4-*d*]pyridazine acceptor due to lower light harvesting by the dye; (ii) the manner in which the donor was attached to the internal acceptor (by carbon or nitrogen) did not play an essential role in the photovoltaic properties of the dyes; (iii) dyes based on the novel donor 2,3,4,4a,9,9a-hexahydro-1*H*-1,4-methanocarbazolyl and 9-(*p*-tolyl)-2,3,4,4a,9,9a-hexahydro-1*H*-carbazole moieties showed similar photovoltaic properties to dyes based on the well-known 4-(*p*-tolyl)-1,2,3,3a,4,8b-hexahydrocyclopenta[*b*]indolyl building block, which opens the door for further optimization potential of new dye families.

Keywords: sulfur-nitrogen heterocycles; [1,2,5]thiadiazolo[3,4-*d*]pyridazine; dye-sensitized solar cells; power conversion efficiency

1. Introduction

In the recent years, solution processable solar cells, including dye-sensitized solar cells (DSSCs) [1–4], bulk heterojunction donor-acceptor blends [5,6], quantum dot solar cells [7,8], organic–inorganic hybrid perovskite solar cells [9–11], and tandem solar cells [12,13] have attracted much attention to in the search to produce low-cost electricity and portable energy. Among them, DSSCs, as a new kind of green energy device, display important properties such as easy fabrication, relatively low production cost, low toxicity, and good flexibility of molecular design [14–16]. Among all the components of DSSCs, the photosensitizers play a key role, being responsible for light harvesting and then electron transfer to a wide band gap semiconducting oxide (typically TiO₂). Hence, the

optoelectronic properties of sensitizing dyes are crucial to the photovoltaic performance of DSSCs. Therefore, the molecular engineering of sensitizing dyes is one of the most efficient routes by which to advance the performance of DSSCs. One of the most successful design strategies of metal-free organic dyes involves the use of D- π -A architectures, due to their easy synthesis and reliable performance [17–20]. Recently, Zhu and Tian proposed a new concept with a D-A- π -A configuration for designing a generation of stable and efficient organic sensitizers [21,22]. Compared to D- π -A dyes, D-A- π -A sensitizers, where the auxiliary acceptor is inserted between the donor and the π -bridge, showed broadened absorption, optimized energy levels, enhanced stability, and efficient intramolecular charge transfer for high performance DSSCs [23]. Some electron-withdrawing groups, such as diketopyrrolopyrrole [24], benzothiadiazole [25], selenadiazolopyridine [26], benzotriazole [27], quinoxaline [28], isoindigo [29], and many others, have been inserted into the D-A- π -A framework. However, so far, a deep understanding of the relationship between the structure of the internal acceptor group and the photovoltaic properties of the DSSCs is still lacking. For example, it has been found that D-A- π -A type dye molecules containing the more electron-withdrawing benzothiadiazole auxiliary unit show higher light-harvesting efficiency than those with benzotriazole moiety, thus resulting in an enhanced photocurrent (J_{sc}) of the corresponding DSSCs [30].

There are a few examples of D-A- π -A dyes where the donor moiety is attached to the acceptor by the nitrogen atom. Comparison of the photovoltaic properties for two sensitizers of D-A- π -A structure, **KM-11** [31] and **D2** [32], (Figure 1) showed that there is no difference between them in photovoltaic efficiency, and the challenge to use a donor fragment attached by the nitrogen atom remains interesting. The introduction of the N-donor fragment into benzene fused with thiadiazole or similar heterocycles does not look simple and promising, since it requires use of palladium-catalyzed Buchwald-Hartwig or the rarely employed Ullman strategy (compared with introduction of a C-donor by similar palladium-catalyzed Suzuki or Stille cross-couplings). Nevertheless, this strategy looks quite promising for highly reactive S_NAr substitution reactions of electron-deficient heterocycles.

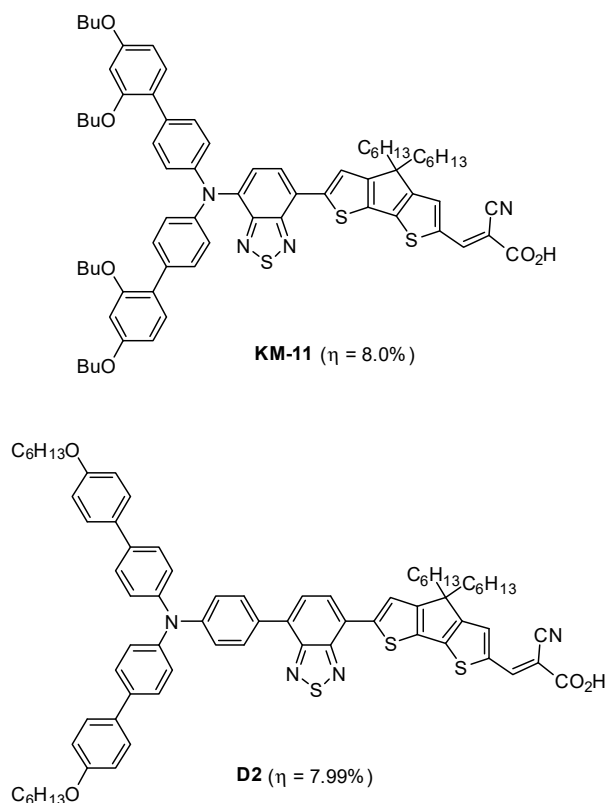


Figure 1. Comparing N- and C-donor attached D-A- π -A dyes.

Recently, we reported the synthesis of the highly electron-deficient [1,2,5]thiadiazolo[3,4-*d*]pyridazine building block, which was evaluated as one of the strongest electron-acceptor systems [33]. It was found that substitution chemistry of 4,7-dibromo[1,2,5]thiadiazolo[3,4-*d*]pyridazine (aromatic nucleophilic and palladium-catalysed cross-couplings) was a powerful tool for the selective formation of various mono- and bis-derivatives of strong electron-accepting heterocycles. Conditions for selective aromatic substitution of one bromine atom by oxygen and nitrogen nucleophiles [34], as well as by Suzuki-Miyaura coupling by arylboronic acids, were found. These results have driven us to the conclusion that this might be a good basis for the synthesis of dyes for DSSCs.

In this work, we therefore aimed to obtain new organic dyes based on the [1,2,5]thiadiazolo[3,4-*d*]pyridazine unit. Taking into account the ease of introduction of N-nucleophiles into [1,2,5]thiadiazolo[3,4-*d*]pyridazine by nucleophilic aromatic substitution, we aimed to design and prepare D-A- π -A sensitizers with the donor attached to the internal heterocyclic acceptor not only by the carbon atom, as it is in a majority of DSSCs, but by the nitrogen atom also. It is known that an indoline group (in **TIM1**) has been proven to endow stronger electron-donating ability than other donor moieties, such as triphenylamine or carbazole [23]. We believed that introduction of similar cycloalkyl groups (such as hexahydrocarbazole in **TIM2** and **TIM4**, and tetrahydromethanocarbazole in **TIM3**) would also increase the donor ability in the corresponding dyes. Herein, the first few dyes using this new internal acceptor (Figure 2) were synthesized, characterized, and tested. The photo-physical and photo-electric properties of these dyes were systemically studied to identify useful design criteria for the discovery of further new dyes within this family.

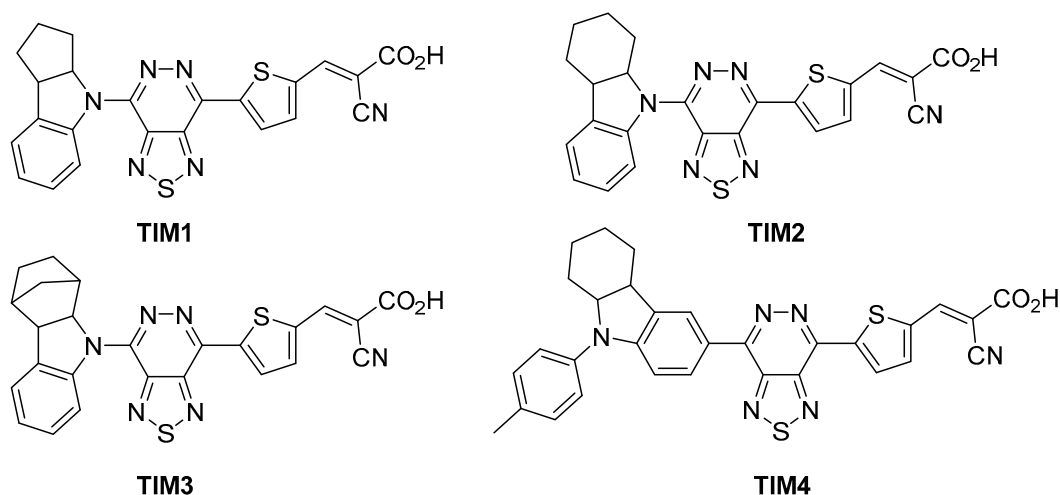
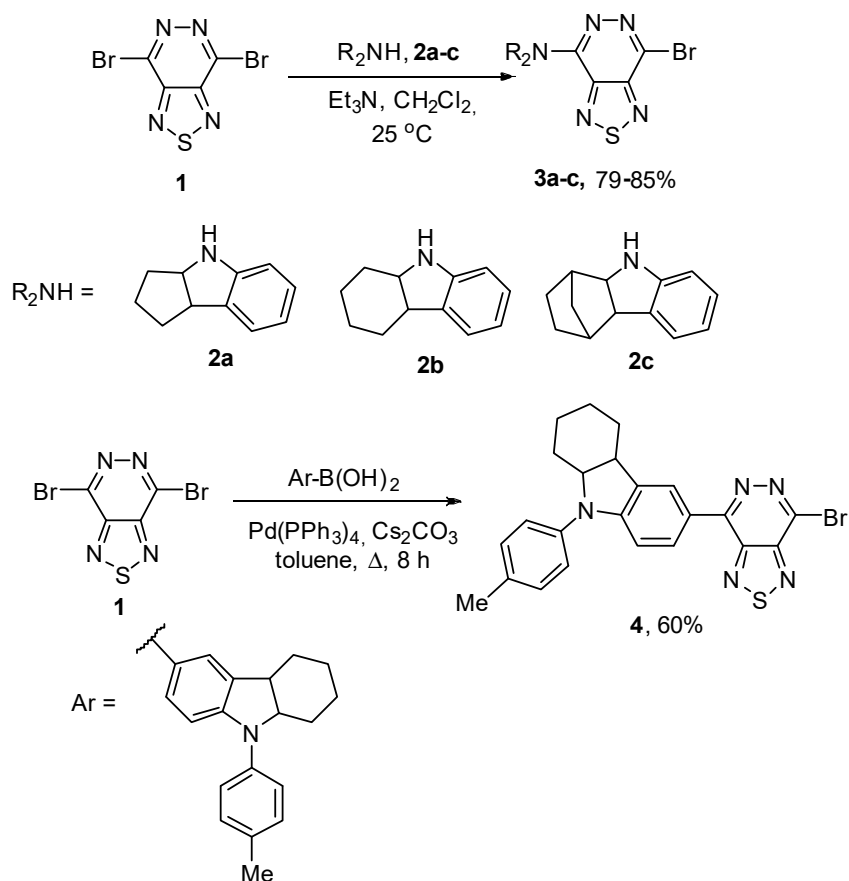


Figure 2. Chemical structures of the dyes synthesized.

2. Results and Discussion

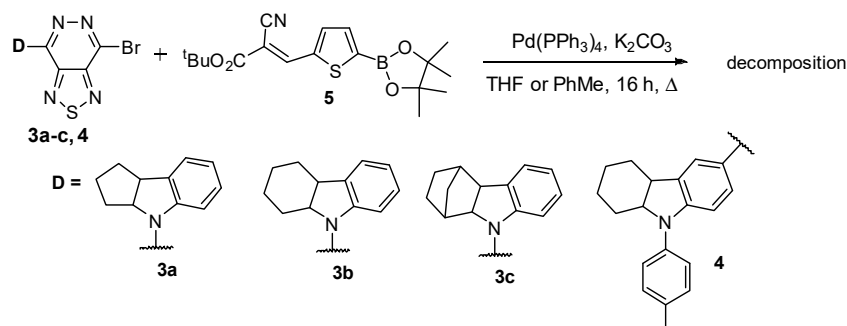
2.1. Synthesis and Characterization

Mono-adducts **3a–c** and **4** were synthesized by nucleophilic aromatic substitution with appropriate amines [34] and by Suzuki cross-coupling reaction with boronic acid, following the procedures described [33] (Scheme 1).



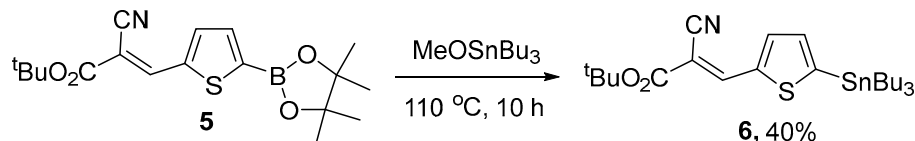
Scheme 1. Synthesis of mono-adducts **3a–c** and **4**.

Unexpectedly, the second Suzuki cross-coupling between mono-adducts **3a–c** and **4** with *tert*-butyl ester **5** in the presence of $\text{Pd}(\text{PPh}_3)_4$ as a catalyst and aqueous solution K_2CO_3 in tetrahydrofuran (THF) or toluene, which was successfully used for 4-bromo[1,2,5]thiadiazolo[3,4-*c*]pyridines and 4-bromo[1,2,5]selenadiazolo[3,4-*c*]pyridines [26], failed, and only decomposition products were detected (Scheme 2).



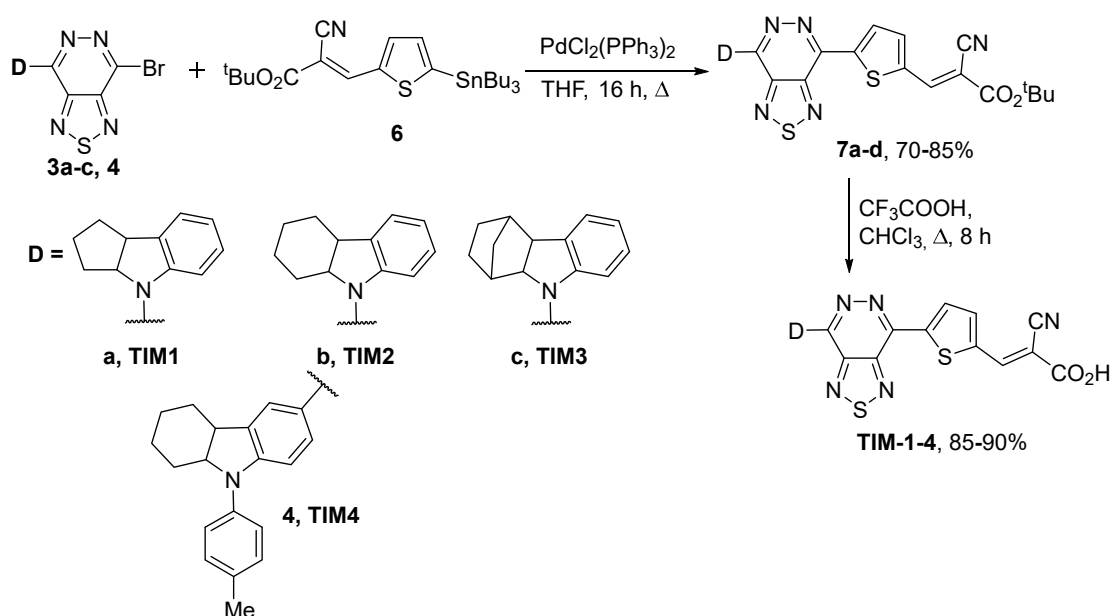
Scheme 2. Reaction of mono-adducts **3a–c** and **4** with *tert*-butyl ester **5**.

We assumed that mono-adducts are insufficiently stable by the action of water or a base (K_2CO_3) and the Stille coupling in non-aqueous and non-basic conditions may be successful. It was recently described that aryltributylstannanes can be easily prepared from arylboronic acids and tributyltin methoxide [35] We have found that thienylboronic ester **5** reacted with tributyltin methoxide under argon to afford thiophenetriethylstannane **6** in moderate yield (Scheme 3).



Scheme 3. Synthesis of *tert*-butyl 2-cyano-3-(5-(tributylstannyl)thiophen-2-yl)acrylate **6**.

Mono-adducts **3a–c** and **4** were subjected to Stille cross-coupling reactions with tributylstannane **6** in the presence of $PdCl_2(PPh_3)_2$ as a catalyst in non-aqueous and non-basic conditions in THF, to afford bis-adducts **7a–d** in high yields. Final hydrolysis of compounds **7a–d** with CF_3CO_2H resulted in the formation of the target dyes in high yields (Scheme 4). All dyes were purified by column chromatography before measurements of the physical and electrochemical properties and solar cell device fabrication.



Scheme 4. Synthesis of dyes.

2.2. Photophysical and Electrochemical Properties

The response region in sunlight for DSSCs is determined primarily by the UV–vis absorption of the sensitizer. Therefore, we initially characterized the spectral response of the **TIM** series in EtOH at 5×10^{-5} mol L^{-1} (Figure 3). The absorption peaks (λ_{max}) and their corresponding molar absorption coefficients (ϵ) are listed in Table 1.

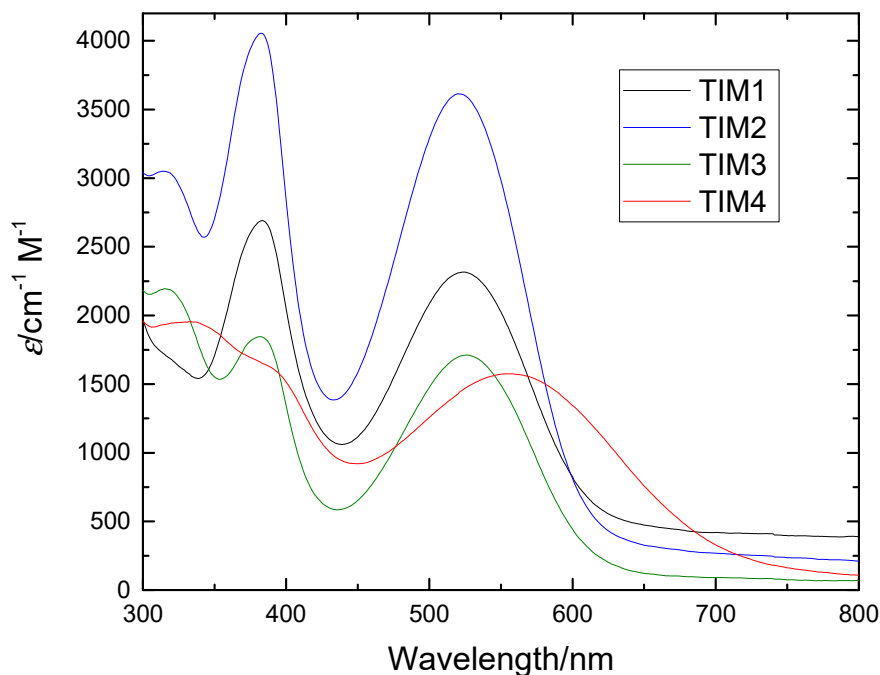


Figure 3. UV-visible absorption of the **TIM** series in EtOH at 5×10^{-5} mol mL⁻¹.

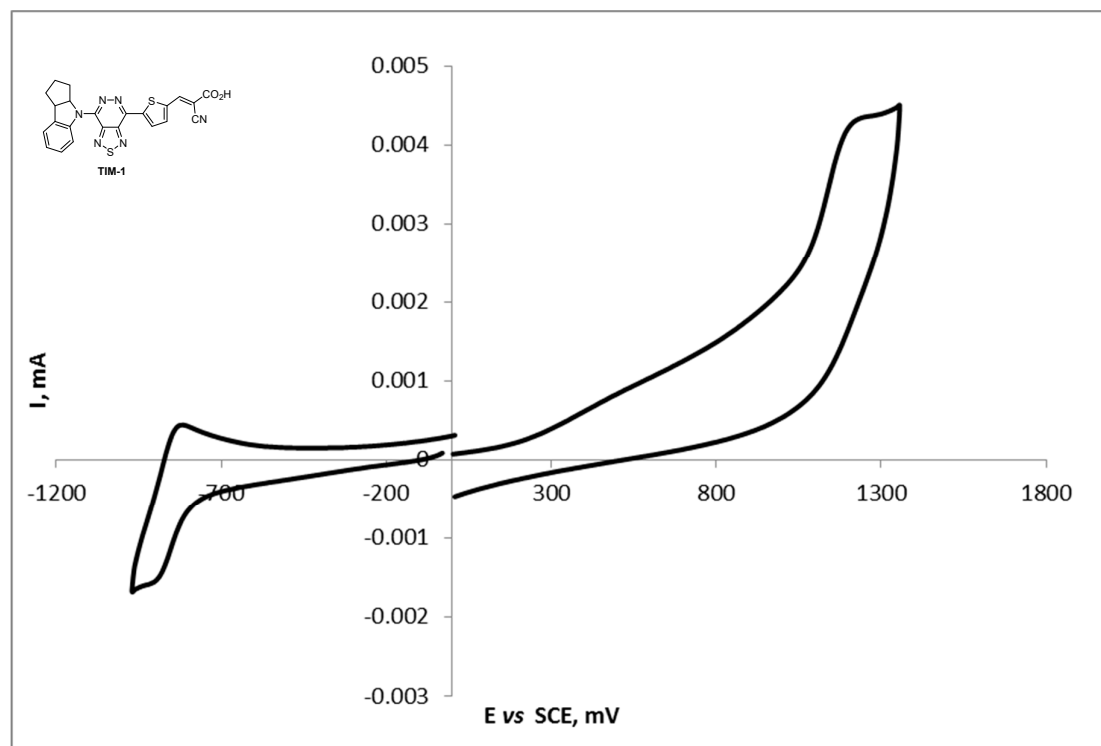
As shown in Figure 3, three compounds of the **TIM** series had two pronounced absorption maxima. Firstly, the absorption in the 380 nm area mainly corresponded to the π - π^* electronic transition. For the **TIM4** compound, this maximum was weakly expressed, but the presence of a plateau in the region of 370–410 nm suggests the presence of the same electronic effects as for the other three dyes of the **TIM** series. Secondly, the absorption bands between 520–560 nm were assigned to an intramolecular charge transfer (ICT) process between the donor and anchor/acceptor group, which produced an efficient charge-separated excited state. We found that, for dyes in which the donor fragment is connected to the pyridazine ring by a nitrogen atom, both absorption maxima wavelengths differed only slightly. The size of the substituent in dihydroindole in the dye **TIM1–3** affected the absorption intensity: the highest extinction coefficient was observed for the derivative with a cyclohexane ring (**TIM2**), and the smallest for the skeleton derivative **TIM3**. The dye **TIM4**, in which the donor fragment was connected to the pyridazine cycle by a carbon atom, had a hypsochromic shift of the long-wavelength absorption maximum compared to the analogues with C–N bonds (**TIM1–3**). This indicates an increase in the conjugation chain in the **TIM4** compound, due to the presence of an additional p-tolyl group. It is worth noting that the extinction coefficient for this dye was also low, as for the derivative norbornene (**TIM3**). In general, the extinction coefficients of all the dyes were low independently from the coupling motif of the donor group to the acceptor heterocycle. The reason for low extinction coefficients is probably in the specific nature of [1,2,5]thiadiazolo[3,4-*d*]pyridazine, which is due to the electron deficient properties of the pyridazine ring [33].

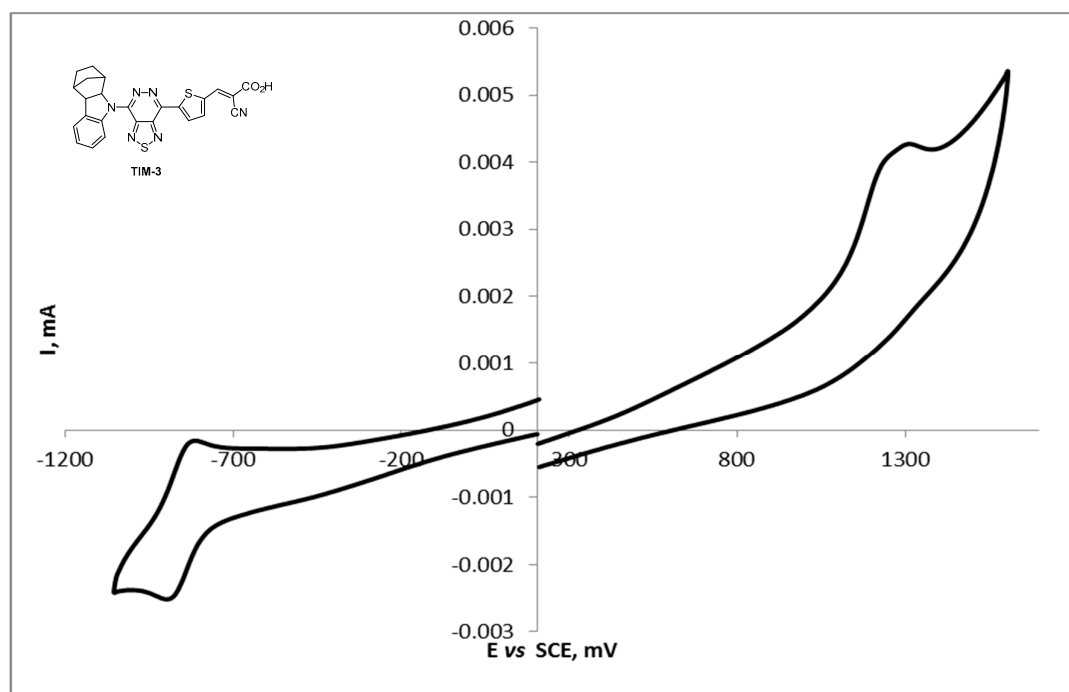
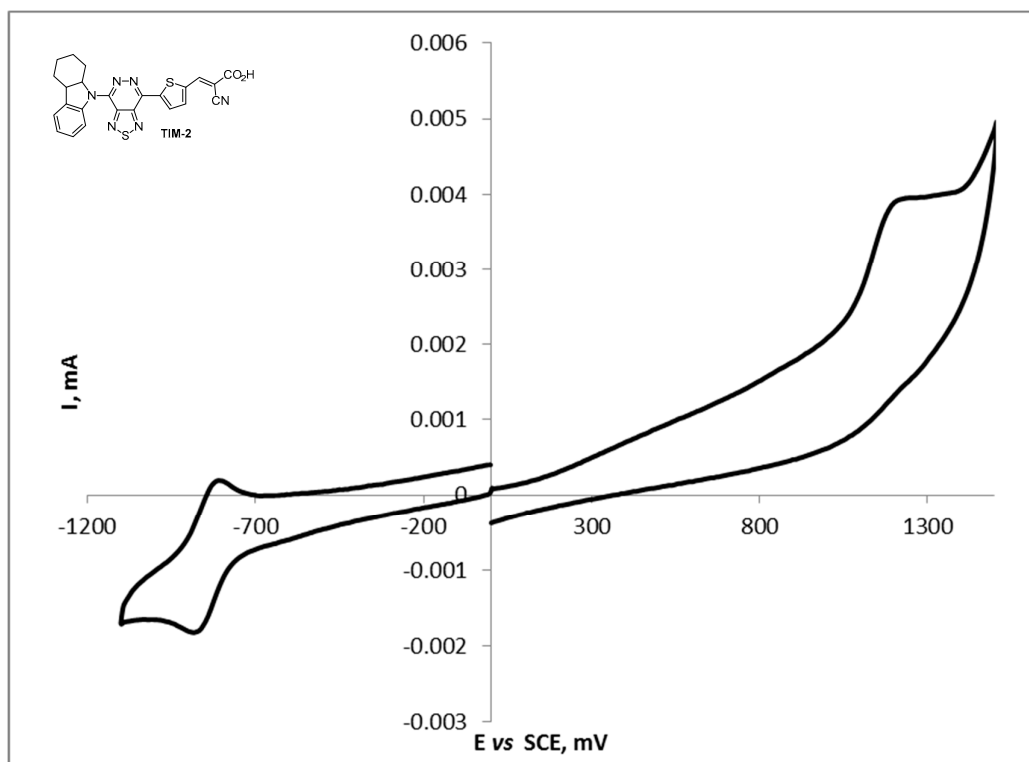
Table 1. The absorption peaks (λ_{\max}) and their corresponding molar absorption coefficients (ϵ) of the TIM series.

Dye	$\lambda_{\max 1}$ [nm] ^a	$\epsilon_{\max 1} \times 10^3$ [M ⁻¹ ·cm ⁻¹] ^a	$\lambda_{\max 2}$ [nm] ^a	$\epsilon_{\max 2} \times 10^3$ [M ⁻¹ ·cm ⁻¹] ^a	λ_{onset} [nm] ^a	$E_{\text{gap}}^{\text{opt}}$ [eV] ^b
TIM1	383	2.7	524	2.3	631	1.968
TIM2	382	4.1	520	3.6	617	2.013
TIM3	382	1.8	526	1.7	620	2.003
TIM4	-	-	554	1.6	722	1.720

^a Absorption peaks (λ_{\max}) and molar extinction coefficients (ϵ_{\max}) in EtOH; ^b Calculated by $1.242/\lambda_{\text{onset}}$.

To estimate the energies of the frontier orbitals of the compounds studied, cyclic voltammograms of TIM1–4 on a Pt electrode in DMF were obtained (Figure 4). TIM1–4 electro-oxidation (EO) is irreversible even at high potential scan rate (10 Vs⁻¹). The first stage of electric recovery (ER) of dyes TIM1–4 is quasi-reversible at low values of the potential sweep rate (100 ms⁻¹), and the cathode current ER is close to the current of a single-electron electrochemical reaction of ferrocene oxidation. To estimate the energy of the frontier orbitals, the potentials of the first stages of ER and EO TIM1–4 were measured with respect to the internal standard, the reversible pair ferrocene/ferrocenium (Fc/Fc⁺), the absolute potential of which was taken as -5.1 eV [36]. The measured redox potentials are collected in Table 2. ER TIM1–3 potentials had similar values (-1.28, -1.26, and -1.27 V, respectively) that is, TIM1–3 substituents probably contributed almost the same to the energy of the E_{LUMO} . The ER TIM4 potential (-1.10 V) was 260–280 mV more positive than in the case of TIM1–3, which can be explained by an increase in conjugation in the TIM4.





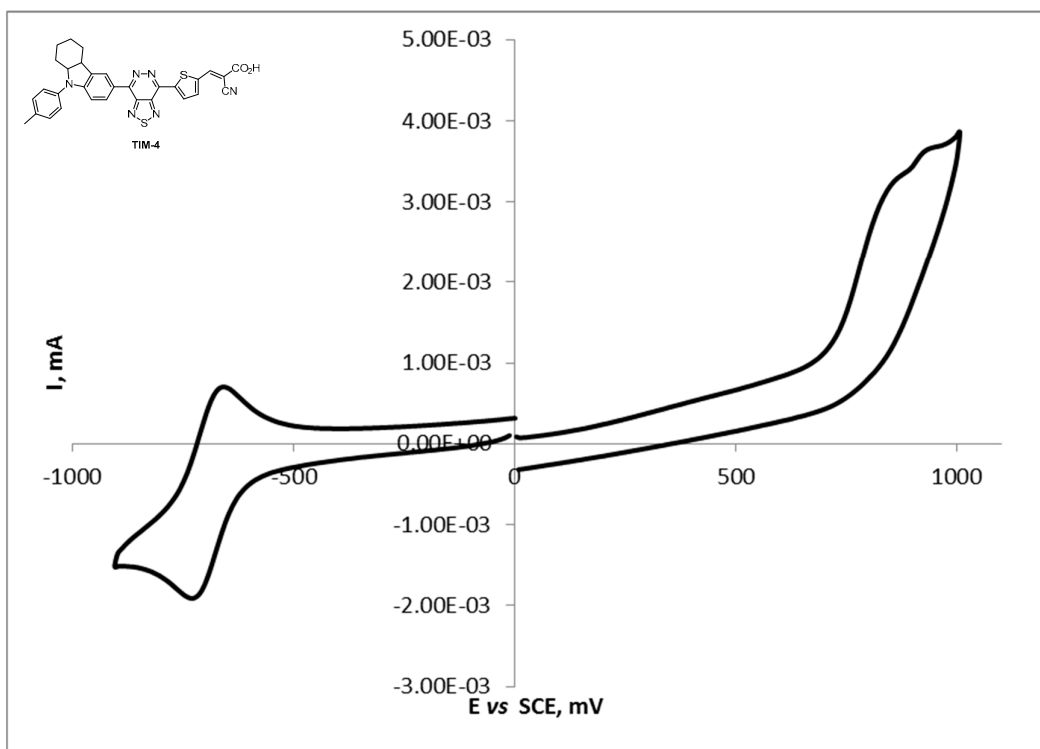


Figure 4. Cyclic voltammograms showing reduction and oxidation of **TIM1–4**. Scan rate 100 mVs^{-1} , electrolyte $0.1 \text{ M Bu}_4\text{NClO}_4$ in DMF.

Since the oxidation is irreversible and the formal potential was not determined, the values E_{LUMO} and E_{HOMO} were calculated by values of the potential onset of peaks ER ($E_{\text{onset}}^{\text{red}}$) and EO ($E_{\text{onset}}^{\text{ox}}$) (Table 2) according to the equations (1) and (2) [36]:

$$E_{\text{HOMO}} (\text{eV}) = - |e| (E_{\text{onset, Fe/Fe}^+}^{\text{ox}} + 5.1) \quad (1)$$

$$E_{\text{LUMO}} (\text{eV}) = - |e| (E_{\text{onset, Fe/Fe}^+}^{\text{red}} + 5.1) \quad (2)$$

The values of E_{LUMO} of all studied compounds were more positive than the energy level of TiO_2 semiconductor (-4.2 eV) [37], and all values of E_{HOMO} were more negative than the level of I^-/I_3^- (-5.2 eV) [38]. That is, on formal grounds, they meet the thermodynamic requirements for dyes. Moreover, the resulting compounds are characterized by a relatively narrow HOMO-LUMO gap $E_{\text{gap}}^{\text{CV}}$: **TIM1** (-1.82 eV) > **TIM2** (-1.80 eV) > **TIM3** (-1.82 eV) > **TIM4** (-1.31 eV), which, according to the literature [39], should enable good efficiency of solar cells.

The analysis of electrochemical parameters of dyes, which showed high efficiency values, indicated that the excess of the E_{LUMO} level over the energy level of TiO_2 should be not less than $0.10\text{--}0.15 \text{ eV}$ [40]. With regard to the efficiency of the regeneration of the dye, it is determined by the magnitude of E_{HOMO} compared to the level of I^-/I_3^- . In different studies, this value has been considered to be optimal from 0.25 [41] to 0.75 eV [42]. Among the synthesized dyes **TIM1–4**, only the E_{HOMO} (-5.31 eV) for **TIM4** may not have a negative enough value.

Table 2. Electrochemical properties of the dyes **TIM1–4** in DMF solution.

Dye	E_{ox} [V] vs. Fc/Fc ⁺ ^a	E_{red} [V] vs Fc/Fc ⁺ ^a	E_{HOMO} [eV] ^b	E_{LUMO} [eV] ^b	E_{gap}^{CV} [eV] ^c
TIM1	0.54	−1.28	−5.64	−3.82	−1.82
TIM2	0.54	−1.26	−5.64	−3.84	−1.80
TIM3	0.52	−1.27	−5.62	−3.83	−1.79
TIM4	0.21	−1.10	−5.31	−4.00	−1.31

^a Here E_{ox} and E_{red} are a linear extrapolation of the low reduction potential side of the first oxidation or reduction wave respectively to the base line relative to Fc/Fc⁺, respectively; ^b Energies of frontier orbitals were calculated according to equations (1) and (2); ^c $E_{gap}^{CV} = E_{LUMO} - E_{HOMO}$.

The E_{gap}^{CV} values obtained from the absorption onset of the **TIM** series dyes are consistent with the data obtained using CV. The **TIM4** compound, containing a C–C bond between the donor and the internal acceptor, has the lowest E_{gap}^{CV} value compared to the three other dyes **TIM1–3** (−1.31 eV and −1.79 eV, −1.82 eV, respectively). Such a significant decrease in energy between HOMO and LUMO is probably due to an increase in the conjugation chain in the **TIM4** compound as compared to derivatives with C–N bonds, which is a consequence of the introduction of the *p*-tolyl substituent into the donor fragment.

2.3. DSSC Performance

To evaluate the dye adsorption on the TiO₂ film, diffuse reflectance was measured for the sensitized photoanodes prior to assembling the device. All samples were irradiated from the non-FTO side to simulate the actual device operation. In order to avoid transmittance effects and purely investigate the absorption, the reference alumina block was placed behind each sample during the measurement. In principle, the lower values in diffuse reflectance relate to higher absorption. The recorded spectra in Figure 5 show a clear drop in the diffuse reflectance around 400 nm and 550 nm, which indicates maximum light absorption of this spectral range by each dye. Similar to the trend in Figure 3, **TIM1–3** display a similar spectral profile, while **TIM4** shows a broader absorption from 600–800 nm, extending to the near-IR. For **TIM1–3**, the diffuse reflectance value of the longer-wavelength peak increased in the order of **TIM2**, **TIM1**, **TIM3**, which is in good accordance with the absorption profile of the solutions. These data indicate that the sensitization was performed in an adequate manner without any unwanted dye aggregation. The results from Figure 3 and Figure 5 imply that the best dye performance is expected for **TIM2**, in terms of light harvesting.

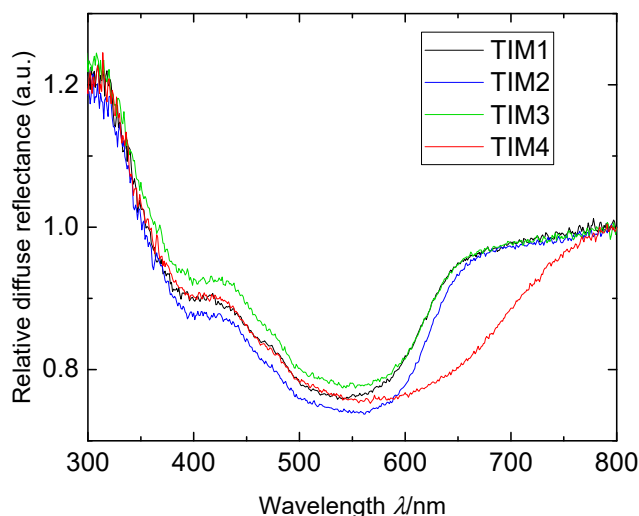


Figure 5. Diffuse reflectance of **TIM1–4**-coated TiO₂ photoanodes. The light was irradiated from the non-FTO side. The irradiated area was fixed with a metal aperture to 0.0625 cm² and an alumina block was placed at the back of the sample to exclude any transmittance. The *y*-axis shows the relative diffuse reflectance, by shifting the tail of the longer-wavelength peak (800 nm) to 1.

The photovoltaic characteristics of each dye in the assembled DSSCs are listed in Figure 6 and Table 3. The overall performance was low, probably due to the low extinction coefficients derived from Figure 3. Comparisons within the series however, may give pointers towards future dye design and performance. Firstly, **TIM4** showed the poorest performance, which is probably due to the E_{HOMO} mismatch suggested earlier. For **TIM1–3**, interestingly, both the current and voltage improved in the order of **TIM2** < **TIM1** < **TIM3**.

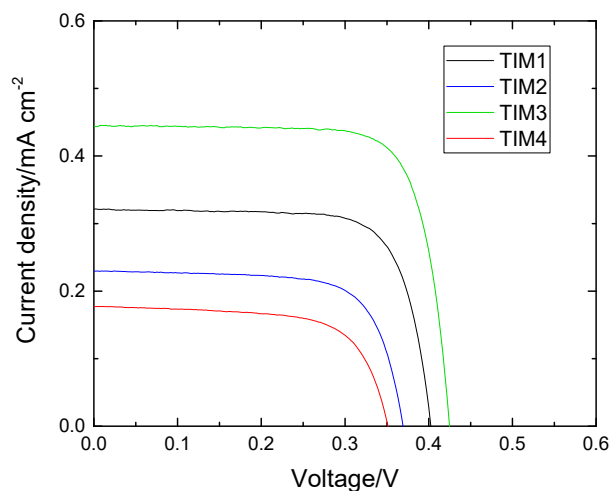


Figure 6. *J*-*V* curves of the dye-sensitized solar cells using **TIM1–4** series as the light absorber.

The outcome that **TIM3**, with the lowest light harvesting ability, exhibited the highest J_{sc} implies that the electron donating properties of the derivatives may be different. In addition, it is noteworthy that a distinct voltage difference was observed between **TIM1–3**, since the E_{HOMO} levels were shown to be identical for these dyes. The reasons behind the poorer results for **TIM1** and **TIM2** may be attributed to poor electron injection from the dye to the TiO₂ and high charge recombination rate.

Table 3. Summary of the device performance for the champion cells. The data for **N719** dye are shown as a reference.

Dye	Voc [V]	J_{sc} [mA cm ⁻²]	FF	PCE [%]
N719	0.67	16.35	0.76	8.29
TIM1	0.39	0.32	0.75	0.09
TIM2	0.37	0.23	0.71	0.06
TIM3	0.42	0.45	0.76	0.14
TIM4	0.35	0.20	0.68	0.05

3. Experimental Section

3.1. Materials and Reagents

The reagents were purchased from commercial sources and used as received. 4,7-Dibromo[1,2,5]thiadiazolo[3,4-*d*]pyridazine (**1**) [34], 4-bromo-7-(1,3,3a,8b-tetrahydrocyclopenta[*b*]indol-4(2*H*)-*yl*)[1,2,5]thiadiazolo[3,4-*d*]pyridazine (**3a**) [34], 4-bromo-7-(2,3,4,4a-hexahydro-1*H*-carbazol-9

(9aH)-yl)[1,2,5]thiadiazolo[3,4-*d*]pyridazine (**3b**) [34], 4-bromo-7-(2,3,4,4a-tetrahydro-1H-1,4-methanocarbazol-9(9aH)-yl)[1,2,5]thiadiazolo[3,4-*d*]pyridazine(**3c**) [34], 4-bromo-7-[9-(*p*-tolyl)-2,3,4,4a,9,9a-hexahydro-1H-carbazol-6-yl][1,2,5]thiadiazolo[3,4-*d*]pyridazine (**4**) [33], *tert*-butyl 2-cyano-3-(4-(4,4,5,5-tetramethyl-1, and 3,2-dioxaboran-2-yl)thiophenyl-yl)acrylate (**5**) [43] were prepared according to the published methods and characterized by NMR spectra. All synthetic operations were performed under a dry argon atmosphere. Solvents were purified by distillation from the appropriate drying agents.

3.2. Analytical Instruments

Melting points were determined on a Kofler hot-stage apparatus and are uncorrected. ¹H- and ¹³C-NMR spectra were taken with a Bruker AM-300 machine (at frequencies of 300.1 and 75.5 MHz, respectively) in CDCl₃ solutions, with TMS as the standard. *J* values are given in Hz. MS spectra (EI, 70 eV) were obtained with a Finnigan MAT INCOS 50 instrument. High-resolution MS spectra were measured on a Bruker micrOTOF II instrument using electrospray ionization (ESI). The measurement was operated in a positive ion mode (interface capillary voltage −4500 V) or in a negative ion mode (3200 V); mass range was from *m/z* 50 to *m/z* 3000 Da; external or internal calibration was done with Electrospray Calibrant Solution (Fluka Chemicals Ltd., Gillingham, UK). A syringe injection was used for solutions in acetonitrile, methanol, or water (flow rate 3 μL·min^{−1}). Nitrogen was applied as a dry gas; interface temperature was set at 180 °C. IR spectra were measured with a Bruker “Alpha-T” instrument (Bruker, Billerica, MA, USA) in KBr pellets.

3.3. General Procedure for Fabrication and Characterization of DSSCs

Fluorine doped tin oxide (FTO) conductive glass (2 mm thick, 7 Ω sheet resistance, Solaronix, Aubonne, Switzerland) was cleaned by sonication in 2% Decon 90 (Decon Laboratories, Hove, UK) in water, deionized water, acetone, and ethanol and treated with UV/O₃ for 20 min. The substrates were treated with 40 mM TiCl₄ at 80 °C for 30 min to form the blocking TiO₂ layer. TiO₂ paste 18NR-T and 18NR-AO (Dyesol, Queanbeyan, Australia) were deposited onto the blocking layer by screen printing, to obtain a film thickness of c.a. 7 μm (3 μm 18NR-T transparent layer +4 μm 18NR-AO scattering layer). The TiO₂ anodes (area = 0.2728 cm²) were annealed in a furnace at 325 °C for 5 min, 375 °C for 5 min, 450 °C for 15 min, and finally 500 °C for 15 min (ramp 10 °C/min). Upon cooling to r.t., the treatment with 40 mM TiCl₄ was repeated and the films were annealed at 500 °C for 30 min. The anodes were kept a 150 °C to release any trapped moisture before immersing in a 0.5 mM ethanol solution of **TIM** dyes with 5 mM chenodeoxycholic acid (CDCA) as an additive. Devices employing **N719** dye were also prepared for comparison. After an overnight sensitization, the anodes were rinsed with ethanol and dried in air. FTO glass for the cathodes was cleaned by sonication in 0.1 M HCl in ethanol, deionized water, acetone, and ethanol. The substrates were platinized by doctor-blading Platisol T/SP (Solaronix, Aubonne, Switzerland) and annealed at 450 °C for 10 min. The solar cells were assembled by binding the two electrodes together with a piece of Surlyn (25 μm, Du Pont, Wilmington, USA) at 125 °C. Electrolyte was vacuum filled into a pre-drilled hole and the hole was sealed by Surlyn and a cover glass. The electrolyte was composed of 0.1 M LiI, 0.05 M I₂, 0.6 M 1,2-dimethyl-3-propylimidazolium iodide (DMPII), and 0.5 M 4-*tert*-butylpyridine in anhyd. acetonitrile.

Current–voltage (*I*–*V*) curves were measured with a potentiostat (Metrohm, Autolab, Herisau, Switzerland) and a class AAA solar simulator (SLB300A, Sciencetech, London, Ontario, Canada). The intensity of the incident light was calibrated to AM1.5G sunlight (100 mW cm^{−2}) using a Si reference cell. The solar cells were masked with a metal aperture to define the active area to 0.0625 cm².

3.4. Detailed Experimental Procedures and Characterization Data

3.4.1. Optical characterization

Solution UV–visible absorption spectra were recorded using a Jasco V-670 UV/Vis/NIR spectrophotometer (JASCO, Mary’s Court Easton, MD, USA) controlled with SpectraManager software. All samples were measured in a 1 cm quartz cell at room temperature with 5 × 10^{−5} mol mL^{−1} concentration in EtOH.

3.4.2. Electrochemical characterization

Electrochemical measurements were carried out in a dry argon atmosphere using an IPC Pro MF potentiostat. The redox properties of compounds were determined using cyclic voltammetry in a three-electrode electrochemical system. A three-electrode system consisting of platinum as working electrode with an area of 0.8 mm², platinum wire as counter electrode, and saturated calomel electrode (SCE) as reference electrode was employed. The reduction and oxidation potentials were determined in DMF, using 0.1 mol L⁻¹ n-Bu₄NClO₄ as the supporting electrolyte. Cyclic voltammetry (CV) measurements used scan rates of 0.1 V·s⁻¹. The first reduction/oxidation potentials were referenced to the internal standard redox couple Fc/Fc⁺. Ferrocene was added to each sample solution at the end of the experiment, and employed for calibration.

3.4.3. Synthesis and characterization of compounds

tert-Butyl 2-cyano-3-(5-(tributylstannyl)thiophen-2-yl)acrylate (**6**): A mixture of *tert*-butyl 2-cyano-3-(4-(4,4,5,5-tetramethyl-1,3,2-dioxaboran-2-yl)thiophenyl-yl) acrylate **5** (0.3 mmol) and Bu₃SnOMe (0.3 mmol) was heated at 110 °C in an argon atmosphere for = 10 h in a sealed vessel. After completion of the chemical reaction (monitored by TLC), the mixture was purified by silica gel column chromatography (eluent: CH₂Cl₂/hexane, 1:2). Colorless oil, yield 63 mg (40%), R_f = 0.6 (CH₂Cl₂/ petroleum ether = 1:1). IR ν_{max} (KBr, cm⁻¹): 2958, 2930, 2853, 2218, 1717, 1590, 1457, 1407, 1370, 1300, 1280, 1238, 1159, 1060, 939, 841, 752, 668, 503. ¹H NMR (300 MHz, CDCl₃): 0.92 (t, J = 7.2 Hz, 9H), 1.16–1.21 (m, 5H), 1.30–1.42 (m, 7H), 1.54–1.62 (m, 15H), 7.28 (d, J = 3.4 Hz, 1H), 7.90 (d, J = 3.4 Hz, 1H), 8.27 (s, 1H). ¹³C-NMR (75 MHz, CDCl₃): δ 11.2, 13.6, 27.2, 28.1, 28.9, 83.3, 99.9, 116.4, 136.6, 136.8, 141.5, 144.9, 152.2, 162.1. HRMS (ESI-TOF), m/z: calcd for C₂₄H₃₉NO₂S¹²⁰SnNa [M + Na]⁺, 548.1618, found 548.1624.

General procedure for the of cross-coupling reaction of mono-adducts **3(a–c)**, **4**, and stannane **6**

To a solution of mono-substituted products **3a–c**, **4** (0.25 mmol) in anhydrous THF (5 mL) were added PdCl₂(PPh₃)₂ (3% mol) and stannane **6** (0.3 mmol). The resulting cloudy yellow mixture was stirred and degassed by argon and refluxed under argon for 8 h. After cooling, an additional amount of stannane **6** (0.3 mmol) and PdCl₂(PPh₃)₂ (3% mol) were added and the reaction mixture was refluxed for 8 h. On completion (monitored by TLC), the mixture was washed with water and the organic layer was extracted with CH₂Cl₂ (3 × 50 mL). The combined organic layers were washed with brine, dried over MgSO₄ and concentrated under reduced pressure. The crude product was purified by column chromatography.

tert-Butyl 2-cyano-3-(5-(7-(1,3,3a,8b-tetrahydrocyclopenta[b]indol-4(2H)-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylate (**7a**): Dark red solid, yield 108 mg (85%), R_f = 0.3 (CH₂Cl₂). Mp 89–91 °C. Eluent - CH₂Cl₂:hexane, 1:1 (v/v). IR ν_{max} (KBr, cm⁻¹): 2955, 2931, 2865, 2216, 1713, 1585, 1503, 1457, 1424, 1367, 1287, 1255, 1244, 1211, 1152, 1109, 1051, 840, 752, 521. ¹H-NMR (300 MHz, CDCl₃): δ 1.44–1.52 (m, 1H), 1.62 (s, 9H), 1.68–1.81 (m, 2H), 2.05–2.17 (m, 2H), 2.20–2.35 (m, 1H), 4.05–4.10 (m, 1H), 5.97–6.02 (m, 1H), 7.12 (t, J = 7.3 Hz, 1H), 7.23 (d, J = 7.3 Hz, 1H), 7.30 (t, J = 8.4 Hz, 1H), 7.93 (d, J = 4.1 Hz, 1H), 8.22 (s, 1H), 8.47 (d, J = 4.1 Hz, 1H), 8.87 (d, J = 8.4 Hz, 1H). ¹³C-NMR (75 MHz, CDCl₃): δ 23.8, 28.0, 34.1, 36.7, 45.8, 67.9, 83.5, 101.2, 115.9, 119.2, 124.0, 124.7, 127.7, 129.4, 136.4, 136.7, 136.9, 142.3, 143.3, 143.4, 145.1, 147.3, 149.3, 149.4, 161.6. HRMS (ESI-TOF), m/z: calcd for C₂₇H₂₅N₆O₂S₂ [M + H]⁺, 529.1475, found 529.1473.

tert-Butyl 2-cyano-3-(5-(7-(2,3,4,4a-tetrahydro-1H-carbazol-9(9aH)-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylate (**7b**): Dark red solid, yield 105 mg (80%), R_f = 0.3 (CH₂Cl₂). Mp 199–201 °C. Eluent - CH₂Cl₂:hexane, 1:1 (v/v). IR ν_{max} (KBr, cm⁻¹): 2925, 2854, 2216, 1701, 1593, 1506, 1457, 1425, 1369, 1290, 1268, 1258, 1233, 1210, 1152, 1092, 1057, 1018, 910, 837, 826, 757, 722, 522, 425. ¹H-NMR (300 MHz, CDCl₃): δ 1.33–1.43 (m, 3H), 1.62 (s, 9H), 1.67–1.74 (m, 2H), 1.92–2.06 (m, 1H), 2.25–2.30 (m, 1H), 2.47 (d, J = 14.1 Hz, 1H), 3.71–3.75 (m, 1H), 5.85–5.90 (m, 1H), 7.21 (t, J = 7.3 Hz, 1H), 7.32 (d, J = 7.3 Hz, 1H), 7.38 (t, J = 8.0 Hz, 1H), 8.00 (d, J = 4.0 Hz, 1H), 8.26 (s, 1H), 8.54 (d, J = 4.0 Hz, 1H), 8.78 (d,

$J = 8.0$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, CDCl_3): δ 20.9, 22.9, 24.3, 28.1, 29.8, 40.3, 64.0, 83.6, 101.2, 116.1, 120.7, 122.6, 124.7, 127.5, 129.4, 135.4, 136.8, 137.0, 142.5, 142.6, 143.0, 145.3, 147.5, 149.3, 149.5, 161.8. HRMS (ESI-TOF), m/z : calcd for $\text{C}_{28}\text{H}_{27}\text{N}_6\text{O}_2\text{S}_2$ $[\text{M} + \text{H}]^+$, 543.1631, found 543.1622.

tert-Butyl 2-cyano-3-(5-(7-(2,3,4,4a-tetrahydro-1H-1,4-methanocarbazol-9(9aH)-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylate (**7c**): Dark red solid, yield 103 mg (75%), $R_f = 0.3$ (CH_2Cl_2). Mp 218–220 °C. Eluent - CH_2Cl_2 :hexane, 1:1 (v/v). IR ν_{max} (KBr, cm^{-1}): 2956, 2924, 2853, 2216, 1716, 1586, 1502, 1457, 1423, 1368, 1292, 1284, 1250, 1211, 1153, 1108, 1037, 813, 752, 522. $^1\text{H-NMR}$ (300 MHz, CDCl_3): 1.10 (d, $J = 10.7$ Hz, 1H), 1.39 (d, $J = 10.06$ Hz, 1H), 1.54–1.57 (m, 1H), 1.62 (s, 9H), 1.64–1.72 (m, 3H), 2.43 (d, $J = 20.0$ Hz, 2H), 3.53 (d, $J = 7.8$ Hz, 1H), 5.45 (d, $J = 7.8$ Hz, 1H), 7.11 (t, $J = 7.3$ Hz, 1H), 7.23 (d, $J = 7.3$ Hz, 1H), 7.30 (t, $J = 8.2$ Hz, 1H), 7.94 (d, $J = 4.2$ Hz, 1H), 8.23 (s, 1H), 8.49 (d, $J = 4.2$ Hz, 1H), 8.90 (d, $J = 8.2$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, CDCl_3): δ 25.9, 28.0, 28.1, 31.9, 43.6, 43.7, 50.6, 69.8, 83.6, 101.3, 116.0, 119.0, 124.3, 124.5, 127.9, 129.5, 135.5, 136.7, 137.1, 142.5, 143.4, 144.9, 145.2, 147.4, 149.4, 149.7, 161.7. HRMS (ESI-TOF), m/z : calcd for $\text{C}_{29}\text{H}_{27}\text{N}_6\text{O}_2\text{S}_2$ $[\text{M} + \text{H}]^+$, 555.1620, found 555.1631.

tert-Butyl 2-cyano-3-(5-(7-(9-(*p*-tolyl)-2,3,4,4a,9,9a-hexahydro-1H-carbazol-6-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylate (**7d**): Violet-green solid, yield 110 mg (70%), $R_f = 0.3$ (CH_2Cl_2). Mp 110–112 °C. Eluent - CH_2Cl_2 :hexane, 1:1 (v/v). IR ν_{max} (KBr, cm^{-1}): 2924, 2853, 2217, 1718, 1603, 1587, 1513, 1474, 1452, 1399, 1270, 1246, 1154, 812, 523. $^1\text{H-NMR}$ (300 MHz, CDCl_3): δ 1.40–1.56 (m, 4H), 1.62 (s, 9H), 1.76–1.77 (m, 2H), 1.92–2.00 (m, 2H), 2.40 (s, 3H), 3.38–3.45 (m, 1H), 4.23–4.29 (m, 1H), 6.87 (d, $J = 8.5$ Hz, 1H), 7.21–7.28 (m, 4H), 8.01 (d, $J = 4.2$ Hz, 1H), 8.27 (s, 1H), 8.66 (d, $J = 1.5$ Hz, 1H), 8.69 (d, $J = 4.2$ Hz, 1H), 8.72 (d, $J = 1.5$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, CDCl_3): δ 21.0, 21.1, 22.3, 26.0, 27.7, 28.0, 40.2, 65.1, 83.8, 102.5, 108.5, 115.8, 123.4, 124.0, 124.8, 130.1, 131.7, 131.8, 134.4, 135.6, 136.5, 139.0, 139.1, 144.9, 146.2, 146.4, 148.5, 149.1, 152.6, 152.8, 161.4. HRMS (ESI-TOF), m/z : calcd for $\text{C}_{35}\text{H}_{33}\text{N}_6\text{O}_2\text{S}_2$ $[\text{M} + \text{H}]^+$, 633.2101, found 633.2084.

General procedure for hydrolysis of ethers **7(a–g)**

To the solution of ether **7a–d** (0.2 mmol) in chloroform (10 mL), trifluoroacetic acid (4 mmol) was added and the mixture was refluxed for 8 h. After cooling, the mixture was concentrated under residue pressure. The crude product was purified by column chromatography.

2-Cyano-3-(5-(7-(1,3,3a,8b-tetrahydrocyclopenta[b]indol-4(2H)-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylic acid (**TIM-1**): Dark red solid, yield 85 mg (90%), $R_f = 0.1$ (EtOAc). Mp 253–255 °C. Eluent – MeOH:EtOAc, 1:1 (v/v). IR ν_{max} (KBr, cm^{-1}): 2959, 2928, 2215, 1685, 1576, 1522, 1508, 1425, 1363, 1293, 1262, 1211, 1140, 1051, 1021, 755, 665, 521. $^1\text{H-NMR}$ (300 MHz, DMSO-d_6): δ 1.29–1.34 (m, 1H), 1.65–1.69 (m, 1H), 1.98–2.03 (m, 1H), 2.10–2.50 (m, 3H), 4.08–4.12 (m, 1H), 5.96–6.01 (m, 1H), 7.12 (t, $J = 7.2$ Hz, 1H), 7.29 (t, $J = 8.1$ Hz, 1H), 7.34 (d, $J = 7.2$ Hz, 1H), 7.79 (d, $J = 3.9$ Hz, 1H), 8.13 (s, 1H), 8.42 (d, $J = 3.9$ Hz, 1H), 8.77 (d, $J = 8.1$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, DMSO-d_6): δ 23.2, 33.5, 35.9, 44.9, 67.2, 111.5, 117.8, 118.8, 123.7, 123.9, 127.0, 128.6, 134.5, 136.1, 138.4, 139.3, 142.3, 142.5, 142.9, 143.2, 148.6, 149.1, 162.7. HRMS (EI-MS): calcd. for $\text{C}_{23}\text{H}_{16}\text{N}_6\text{O}_2\text{S}_2\text{Na}$ $[\text{M} + \text{Na}]^+$, 495.0668, found 495.0668.

2-Cyano-3-(5-(7-(2,3,4,4a-tetrahydro-1H-carbazol-9(9aH)-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylic acid (**TIM-2**): Dark red solid, yield 87 mg (90%), $R_f = 0.1$ (EtOAc). Mp 138–140 °C. Eluent – MeOH:EtOAc, 1:1 (v/v). IR ν_{max} (KBr, cm^{-1}): 2927, 2855, 2215, 1680, 1510, 1459, 1380, 1295, 1212, 1152, 848, 803, 726, 520. $^1\text{H-NMR}$ (300 MHz, DMSO-d_6): δ 1.34–1.37 (m, 3H), 1.60–1.63 (m, 2H), 1.91–1.95 (m, 1H), 2.18–2.24 (m, 1H), 2.42 (d, $J = 13.9$ Hz, 1H), 3.71–3.73 (m, 1H), 5.80–5.85 (m, 1H), 7.18 (t, $J = 7.2$ Hz, 1H), 7.32 (t, $J = 8.3$ Hz, 1H), 7.39 (d, $J = 7.2$ Hz, 1H), 7.84 (d, $J = 3.9$ Hz, 1H), 8.20 (s, 1H), 8.43 (d, $J = 3.9$ Hz, 1H), 8.65 (d, $J = 8.3$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, DMSO-d_6): δ 20.3, 21.8, 23.3, 27.3, 40.4, 62.8, 109.9, 118.4, 121.5, 122.4, 123.7, 126.7, 128.6, 135.0, 135.2, 137.9, 140.3, 142.3, 142.4, 142.5, 143.1, 148.6, 149.0, 163.6. HRMS (EI-MS): calcd. for $\text{C}_{24}\text{H}_{18}\text{N}_6\text{O}_2\text{S}_2\text{Na}$ $[\text{M} + \text{Na}]^+$, 509.0825, found 509.0815.

2-Cyano-3-(5-(7-(1,2,3,4,4a,9a-hexahydro-9H-1,4-methanocarbazol-9-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylic acid (TIM3): Dark red solid, yield 85 mg (86%), $R_f = 0.1$ (EtOAc). Mp > 260 °C. Eluent – MeOH:EtOAc, 1:1 (v/v). IR ν_{\max} (KBr, cm^{-1}): 2929, 2856, 2216, 1686, 1576, 1507, 1423, 1211, 1189, 1140, 840, 660. $^1\text{H-NMR}$ (300 MHz, DMSO- d_6): δ 1.09 (d, $J = 10.7$ Hz, 1H), 1.17 (d, $J = 10.06$ Hz, 1H), 1.58–1.62 (m, 4H), 2.43–2.47 (m, 2H), 3.60 (d, $J = 7.9$ Hz, 1H), 5.49 (d, $J = 7.9$ Hz, 1H), 7.09 (t, $J = 7.3$ Hz, 1H), 7.27 (d, $J = 7.3$ Hz, 1H), 7.33 (t, $J = 8.2$ Hz, 1H), 7.78 (d, $J = 4.0$ Hz, 1H), 8.08 (s, 1H), 8.42 (d, $J = 4.0$ Hz, 1H), 8.78 (d, $J = 8.2$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, DMSO- d_6): δ 24.9, 27.3, 31.3, 42.8, 42.9, 49.6, 68.9, 111.9, 117.6, 119.2, 123.7, 124.4, 127.3, 128.7, 135.1, 135.2, 138.5, 139.3, 142.5, 142.6, 143.2, 144.7, 148.7, 149.4, 162.2. HRMS (EI-MS): calcd. for $\text{C}_{25}\text{H}_{18}\text{N}_6\text{O}_2\text{S}_2\text{Na}$ [$\text{M} + \text{Na}$] $^+$, 521.0825, found 521.0820.

2-Cyano-3-(5-(7-(9-(p-tolyl)-2,3,4,4a,9,9a-hexahydro-1H-carbazol-6-yl)-[1,2,5]thiadiazolo[3,4-d]pyridazin-4-yl)thiophen-2-yl)acrylic acid (TIM4): Violet solid, yield 98 mg (85%), $R_f = 0.1$ (EtOAc), Mp > 260 °C. Eluent – MeOH:EtOAc, 1:1 (v/v). IR ν_{\max} (KBr, cm^{-1}): 2926, 2854, 2213, 1718, 1603, 1561, 1513, 1450, 1399, 1376, 1272, 1135, 1108, 811, 522. $^1\text{H-NMR}$ (300 MHz, DMSO- d_6): δ 1.42–1.61 (m, 4H); 1.53–1.67 (m, 2H); 1.90–2.01 (m, 2H); 2.34 (s, 3H); 3.39–3.46 (m, 1H); 4.29–4.35 (m, 1H); 6.82 (d, $J = 8.6$ Hz, 1H); 7.22–7.29 (m, 4H); 7.83 (d, $J = 4.1$ Hz, 1H); 8.09 (s, 1H); 8.49–8.53 (m, 3H), 8.57 (d, $J = 4.1$ Hz, 1H). $^{13}\text{C-NMR}$ (75 MHz, DMSO- d_6): δ 20.3, 20.4, 21.7, 25.1, 27.2, 40.4, 63.8, 107.8, 119.1, 122.7, 124.0, 124.1, 126.5, 130.0, 130.5, 131.4, 133.3, 135.0, 135.1, 138.7, 139.1, 140.7, 141.6, 146.2, 147.7, 148.4, 151.2, 151.7, 162.2. HRMS (EI-MS): calcd. for $\text{C}_{31}\text{H}_{24}\text{N}_6\text{O}_2\text{S}_2\text{Na}$ [$\text{M} + \text{Na}$] $^+$, 599.1294, found 599.1287.

4. Conclusions

In summary, we have designed four new D-A- π -A metal-free organic sensitizers with [1,2,5]thiadiazolo[3,4-d]pyridazine as the internal acceptor, thiophene unit as the π -spacer, and cyanoacrylate as the anchoring electron acceptor. Despite the variation of the donor fragment and manner of its attachment to the internal acceptor, low PCE data were measured for all dyes. There are a few conclusions that can be made from the results obtained. First of all, comparison of the PCE for **TIM1–TIM3** drives us to conclude that there is no significant impact originating from the *N*-donor; they showed quite similar PCE values, with a slightly better result for the 2,3,4,4a,9,9a-hexahydro-1H-1,4-methanocarbazolyl moiety (**TIM3**). These data are in good agreement with the data obtained by us earlier [44], with sensitizers based on 9-(*p*-tolyl)-2,3,4,4a,9,9a-hexahydro-1H-carbazole (used in **TIM4**) showing similar results to the well-known 4-(*p*-tolyl)-1,2,3,3a,4,8b-hexahydrocyclopenta[*b*]indolyl building block (see, for example, dye **WS-2** [4]). Secondly, the comparison of PCE data for **TIM2** and **TIM4** showed practically identical values, indicating a small difference in attachment of the donor fragment to the internal acceptor moiety, which may open another interesting possibility for flexibility of the chemical structures of the dyes. Finally, the relatively low performance suggests that the internal [1,2,5]thiadiazolo[3,4-d]pyridazine acceptor was responsible for poor PCE data due to lower light harvesting by the dye, which is not beneficial when designing D-A- π -A dyes for DSSCs. This is not surprising, since benzo[*c*][1,2,5]thiadiazoles showed much better PCE values than [1,2,5]thiadiazolo[3,4-*c*]pyridines; apparently, employing high electron-accepting internal building blocks is unfavorable to the improvement of PCE.

Supplementary Materials: Results of quantum calculations and characterization data including ^1H and ^{13}C NMR spectra for the compounds **6**, **7a–d**, **TIM1–3**.

Author Contributions: O.A.R. and N.R. conceived and designed the study; T.N.C., E.T., L.V.M. and V.V.P. performed the experiments; E.A.K. and E.T. analysed the data; all authors contributed to writing and editing the paper.

Funding: This research was funded by the Russian Science Foundation (grant number 15-13-10022).

Acknowledgments: We gratefully acknowledge financial support from the Russian Science Foundation (grant number 15-13-10022). E.T. thanks JASSO for a PhD studentship. V.V.P. is grateful to South Ural State University. Authors thank I. S. Golovanov (N.D. Zelinsky Institute of Organic Chemistry) for quantum chemical calculations.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Oregan, B.; Grätzel, M. A low-cost, high-efficiency solar cell based on dye sensitized colloidal TiO₂ films. *Nature* **1991**, *353*, 737–739, doi:10.1038/353737a0.
2. Qin, C.; Numata, Y.; Zhang, S.; Yang, X.; Islam, A.; Zhang, K.; Chen, H.; Han, L. Novel Near-Infrared Squaraine Sensitizers for Stable and Efficient Dye-Sensitized Solar Cells. *Adv. Funct. Mater.* **2014**, *24*, 3059–3066, doi:10.1002/adfm.201203384.
3. Yella, A.; Lee, H.; Tsao, H.N.; Yi, C.; Chandiran, A.K.; Nazeeruddin, M.K.; Diao, E.W.; Yeh, C.; Zakeeruddin, S.M.; Gratzel, M. Porphyrin-Sensitized Solar Cells with Cobalt(II/III)-Based Redox Electrolyte Exceed 12 Percent Efficiency. *Science* **2011**, *334*, 629–634, doi:10.1126/science.1209688.
4. Knyazeva, E.A.; Rakitin, O.A. Influence of structural factors on the photovoltaic properties of dye-sensitized solar cells. *Russ. Chem. Rev.* **2016**, *85*, 1146–1183, doi:10.1070/RCR4649.
5. W. C. Zhao, S.S. Li, H.F. Yao, S.Q. Zhang, Y. Zhang, B. Yang, J.H. Hou, Molecular Optimization Enables over 13% Efficiency in Organic Solar Cells *J. Am. Chem. Soc.* **2017**, *139*, 7148–7151, doi:10.1021/jacs.7b02677.
6. Zhao, W.; Qian, D.; Zhang, S.; Li, S.; Inganas, O.; Gao, F.; Hou, J. Fullerene-Free Polymer Solar Cells with over 11% Efficiency and Excellent Thermal Stability. *Adv. Mater.* **2016**, *28*, 4734–4739, doi:10.1002/adma.201600281.
7. Du, J.; Du, Z.; Hu, J.; Pan, Z.; Shen, Q.; Sun, J.; Long, D.; Dong, H.; Sun, L.; Zhong, X.; Wan, L. Zn-Cu-in-Se Quantum Dot Solar Cells with a Certified Power Conversion Efficiency of 11.6%. *J. Am. Chem. Soc.* **2016**, *138*, 4201–4209, doi:10.1021/jacs.6b00615.
8. Pan, Z.; Rao, H.; Mora-Seró, I.; Bisquert, J.; Zhong, X. Quantum dot-sensitized solar cells. *Chem. Soc. Rev.* **2018**, *47*, 7659–7702, doi:10.1039/C8CS00431E.
9. Rong, Y.; Hu, Y.; Mei, A.; Tan, H.; Saidaminov, M.I.; Seok, S.I.; McGehee, M.D.; Sargent, E.H.; Ha, H. Challenges for commercializing perovskite solar cells. *Science*, **2018**, *361*, eaat8235, doi:10.1126/science.aat8235.
10. Jena, A. K.; Kulkarni, A.; Miyasaka, T. Halide Perovskite Photovoltaics: Background, Status, and Future Prospects. *Chem. Rev.* **2019**, *119*, 3036–3103. doi:10.1021/acs.chemrev.8b00539.
11. Saliba, M.; Correa-Baena, J.-P.; Wolff, C.M.; Stollerfoht, M.; Phung, N.; Albrecht, S.; Neher, D.; Abate, A.; How to Make over 20% Efficient Perovskite Solar Cells in Regular (n-i-p) and Inverted (p-i-n) Architectures. *Chem. Mater.*, **2018**, *30*, 4193–4201, doi:10.1021/acs.chemmater.8b00136.
12. Meng, L.; Zhang, Y.; Wan, X.; Li, C.; Zhang, X.; Wang, Y.; Ke, X.; Xiao, Z.; Ding, L.; Xia, R.; Yip, H.-L.; Cao, Y.; Chen, Y. Organic and solution-processed tandem solar cells with 17.3% efficiency. *Science*, **2018**, *361*, 1094–1098, doi:10.1126/science.aat2612.
13. Cui, Y.; Yao, H.; Gao, B.; Qin, Y.; Zhang, S.; Yang, B.; He, C.; Xu, B.; Hou, J. Fine-tuned photoactive and interconnection layers for achieving over 13% efficiency in a fullerene-free tandem organic solar cell. *J. Am. Chem. Soc.* **2017**, *139*, 7302–7309, doi:10.1021/jacs.7b01493.
14. Liang, M.; Chen, J. Arylamine organic dyes for dye-sensitized solar cells. *Chem. Soc. Rev.* **2013**, *42*, 3453–3488, doi:10.1039/C3CS35372A.
15. Yang, J.B.; Ganesan, P.; Teuscher, J.; Moehl, T.; Kim Y. J.; Yi, C.Y.; Comte, P.; Pei, K.; Holcombe, T.W.; Nazeeruddin, M.K.; Hua, J.L.; Zakeeruddin, S.M.; Tian, H.; Grätzel, M. Influence of the donor size in D- π -A organic dyes for dye-sensitized solar cells. *J. Am. Chem. Soc.* **2014**, *136*, 5722–5730, doi:10.1021/ja500280r.
16. Hung, W.; Liao, Y.; Lee, T.; Ting, Y.; Ni, J.; Kao, W.; Lin, J.T.; Wei, T.; Yen, Y. Eugenol Metal-Free Sensitizers with Double Anchors for High Performance Dye-Sensitized Solar Cells. *Chem. Commun.* **2015**, *51*, 2152–2155, doi:10.1039/C4CC09294E.
17. Li, L.L.; Diao, E.W.G. Porphyrin-sensitized solar cells. *Chem. Soc. Rev.* **2013**, *42*, 291–304, doi:10.1039/C2CS35257E.
18. Huang, Z.-S.; Hua, T.; Tian, J.; Wang, L.; Meier, H.; Cao, D. Dithienopyrrolobenzotriazole-based organic dyes with high molar extinction coefficient for efficient dye-sensitized solar cells. *Dyes Pigm.* **2016**, *125*, 229–240, doi:10.1016/j.dyepig.2015.10.022.
19. Fischer, M.K.R.; Wenger, S.; Wang, M.; Mishra, A.; Zakeeruddin, S.M.; Grätzel, M.; Bäuerle, P. D- π -A sensitizers for dye-sensitized solar cells: linear vs branched oligothiophenes. *Chem. Mater.* **2010**, *22*, 1836–45, doi:10.1021/cm903542v.
20. Ganesan, P.; Yella, A.; Holcombe, T.W.; Gao, P.; Rajalingam, R.; Al-Muhtaseb, S.A.; Grätzel, M.; Nazeeruddin, M.K. Unravel the impact of anchoring groups on the photovoltaic performances of

- diketopyrrolopyrrole sensitizers for dye-sensitized solar cells. *ACS Sustainable Chem. Eng.* **2015**, *3*, 2389–2396, doi:10.1021/acsschemeng.5b01026.
21. Zhu, W.H.; Wu, Y.Z.; Wang, S.T.; Li, W.Q.; Li, X.; Chen, J.; Wang, Z.S.; Tian, H. Organic D-A- π -A Solar Cell Sensitizers with Improved Stability and Spectral Response. *Adv. Funct. Mater.*, **2011**, *21*, 756–763, doi:10.1002/adfm.201001801.
 22. He, J.; Wu, W.; Hua, J.; Jiang, Y.; Qu, S.; Li, J.; Long, Y.; Tian, H. Bithiazole-bridged dyes for dye-sensitized solar cells with high open circuit voltage performance. *J. Mater. Chem.* **2011**, *21*, 6054–6062, doi:10.1039/C0JM03811C.
 23. Wu, Y.; Zhu, W. Organic sensitizers from D–p–A to D–A–p–A: effect of the internal electron-withdrawing units on molecular absorption, energy levels and photovoltaic performances. *Chem. Soc. Rev.*, **2013**, *42*, 2039–2058, doi:10.1039/c2cs35346f.
 24. Qu, S.; Qin, C.; Islam, A.; Wu, Y.; Zhu, W.; Hua, J.; Tian, H.; Han, L. A novel D-A- π -A organic sensitizer containing a diketopyrrolopyrrole unit with a branched alkyl chain for highly efficient and stable dye-sensitized solar cells. *Chem. Commun.* **2012**, *48*, 6972–6974, doi:10.1039/C2CC31998E.
 25. Huang, Z.S.; Zang, X.F.; Hua, T.; Wang, L.; Meier, H.; Cao, D. 2,3-Dipentylidithieno[3,2-f:2',3'-h]quinoxaline-Based organic dyes for efficient dye-sensitized solar cells: effect of π -bridges and electron donors on solar cell performance. *ACS Appl. Mater. Interfaces.* **2015**, *7*, 20418–20429, doi:10.1021/acsami.5b06404.
 26. Knyazeva, E.A.; Wu, W.; Chmovzh, T.N.; Robertson, N.; Woollins, J.D.; Rakitin, O.A. Dye-sensitized solar cells: Investigation of D-A- π -A organic sensitizers based on [1,2,5]selenadiazolo[3,4-c]pyridine. *Sol. Energy* **2017**, *144*, 134–143, doi:10.1016/j.solener.2017.01.016.
 27. Cui, Y.; Wu, Y.Z.; Lu, X.F.; Zhang, X.; Zhou, G.; Miapah, F.B.; Zhu, W.H.; Wang, Z.S. Incorporating Benzotriazole Moiety to Construct D-A- π -A Organic Sensitizers for Solar Cells: Significant Enhancement of Open-Circuit Photovoltage with Long Alkyl Group. *Chem. Mater.*, **2011**, *23*, 4394–4401, doi:10.1021/cm202226j.
 28. Zhu, H.; Liu, B.; Liu, J.; Zhang, W.; Zhu, W.-H. D–A– π –A featured sensitizers by modification of auxiliary acceptor for preventing “trade-off” effect. *J. Mater. Chem. C* **2015**, *3*, 6882–6890, doi:10.1039/C5TC01195G.
 29. Ying, W.; Guo, F.; Li, J.; Zhang, Q.; Wu, W.; Tian, H.; Hua, J. Series of new D–A– π –A organic broadly absorbing sensitizers containing isoindigo unit for highly efficient dyesensitized solar cells. *ACS Appl. Mater. Interfaces* **2012**, *4*, 4215–4224, doi:10.1021/am300925e.
 30. Lua, F.; Yanga, G.; Xua, Q.; Zhanga, J.; Zhanga, B.; Feng, Y. Tailoring the benzotriazole (BTZ) auxiliary acceptor in a D-A- π -A type sensitizer for high performance dye-sensitized solar cells (DSSCs). *Dyes Pigm.* **2018**, *158*, 195–203, doi:10.1016/j.dyepig.2018.05.044.
 31. Katono, M.; Wielopolski, M.; Marszalek, M.; Bessho, T.; Moser, J.-E.; Humphry-Baker, R.; Zakeeruddin, S.M.; Grätzel, M. Effect of Extended π -Conjugation of the Donor Structure of Organic D–A– π –A Dyes on the Photovoltaic Performance of Dye-Sensitized Solar Cells. *J. Phys. Chem. C* **2014**, *118*, 16486–16493, doi:10.1021/jp411504p
 32. Wang, X.; Yang, J.; Yu, H.; Li, F.; Fan, L.; Sun, W.; Liu, Y.; Koh, Z.Y.; Pan, J.; Yim, W.-L.; Yan, L.; Wang, Q. A benzothiazole–cyclopentadithiophene bridged D–A–p–A sensitizer with enhanced light absorption for high efficiency dye-sensitized solar cells. *Chem. Commun.* **2014**, *50*, 3965–3968, doi:10.1039/C4CC00577E.
 33. Chmovzh, T.N.; Knyazeva, E.A.; Mikhalchenko, L.V.; Golovanov, I.S.; Amelichev, S.A.; Rakitin, O.A. Synthesis of 4,7-dibromo derivative of ultrahigh electron-deficient [1,2,5]thiadiazolo[3,4-d]pyridazine heterocycle and its cross-coupling reactions, *Eur. J. Org. Chem.*, **2018**, *41*, 5668–5677, doi:10.1002/ejoc.201800961.
 34. Chmovzh, T.N.; Knyazeva, E.A.; Lyssenko, K.A.; Popov, V.V.; Rakitin, O.A. Safe synthesis of 4,7-dibromo[1,2,5]thiadiazolo[3,4-d]pyridazine and its S_NAr reactions. *Molecules* **2018**, *23*, 2576, doi:10.3390/molecules23102576.
 35. Oikawa, A.; Kindaichi, G.; Shimotori, Y.; Okimoto, M.; Hoshi, M. Simple preparation of aryltributylstannanes and its application to one-pot synthesis of diaryl ketones. *Tetrahedron* **2015**, *71*, 1705–1711, doi:10.1016/j.tet.2015.01.048.
 36. Cardona, C.M.; Li, W.; Kaifer, A.E.; Stockdale, D.; Bazan, G.C. Electrochemical Considerations for Determining Absolute Frontier Orbital Energy Levels of Conjugated Polymers for Solar Cell Applications, *Adv. Mater.* **2011**, *23*, 2367–2371, doi:10.1002/adma.201004554.

37. Oskam, G.; Bergeron, B.V.; Meyer, G.J.; Searson, P.C. Pseudohalogens for Dye-Sensitized TiO₂ Photoelectrochemical Cells. *J. Phys. Chem. B* **2001**, *105*, 6867–6873, doi:10.1021/jp004411d.
38. Babu, D.D.; Cheema, H.; Elsherbiny, D.; El-Shafei, A.; Adhikari, A.V. Molecular Engineering and Theoretical Investigation of Novel Metal-Free Organic Chromophores for Dye-Sensitized Solar Cells. *Electrochim. Acta* **2015**, *176*, 868–879, doi:10.1016/j.electacta.2015.07.079.
39. Akkuratov, A.V.; Troshin, P.A. Conjugated Polymers with Benzothiadiazole, Benzoxadiazole, and Benzotriazole Moieties as Promising Semiconductor Materials for Organic Solar Cells. *Polym. Sci., Ser. B* **2014**, *56*, 414–442, doi:10.1134/S1560090414040010.
40. Koops, S.E.; O'Regan, B.C.; Barnes, P.R.F.; Durrant, J.R. Parameters influencing the efficiency of electron injection in dye-sensitized solar cells. *J. Am. Chem. Soc.* **2009**, *131*, 4808–4818, doi:10.1021/ja8091278.
41. Hussain, M.; Islam, A.; Bedja, I.; Gupta, R.K.; Han, L.; El-Shafei, A. A comparative study of Ru(II) cyclometallated complexes versus thiocyanated heteroleptic complexes: thermodynamic force for efficient dye regeneration in dye-sensitized solar cells and how low could it be? *Phys. Chem. Chem. Phys.* **2014**, *16*, 14874–14881, doi:10.1039/C4CP00907J.
42. Hardin, B.E.; Snaith, H.J.; McGehee, M.D. The renaissance of dye-sensitized solar cells, *Nature Photonics* **2012**, *6*, 162–169, doi:10.1038/nphoton.2012.22.
43. Fuse, S.; Sugiyama, S.; Maitani, M.M.; Wada, Y.; Ogomi, Y.; Hayase, S.; Katoh, R.; Kaiho, T.; Takahashi, T. Elucidating the Structure–Property Relationships of Donor– π -Acceptor Dyes for Dye-Sensitized Solar Cells (DSSCs) through Rapid Library Synthesis by a One-Pot Procedure. *Chem. Eur. J.* **2014**, *20*, 10685–10694, doi:10.1002/chem.201402093.
44. Mikhailov, M.S.; Gudim, N.S.; Knyazeva, E.A.; Zhang, L.; Mikhalchenko, L.V.; Robertson, N.; Rakitin, O.A. will be published elsewhere.

Sample Availability: Samples of the compounds are available from the authors.



© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).