Betterment of optoelectronic and photovoltaic properties is reported through the embodiment of a planar, conjugated naphthalene diimide functionality in a given donor-acceptor small molecular oligothiophene, when used in solution-processable bulk heterojunction solar cells.
Improvement of optoelectronic and photovoltaic properties through the insertion of a naphthalenediimide unit in donor-acceptor oligothiophenes†

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Improvements to the optoelectronic and photovoltaic properties of 2-((5’-(4-(diphenylamino)phenyl)-[2,2’-bithiophen]-5-yl)methylene)malononitrile (R1; a donor-acceptor small molecular oligothiophene) are reported through the insertion of planar, conjugated naphthalenediimide moiety. The novel donor material was successfully synthesised and exhibited a power conversion efficiency of 3.01% under AM 1.5 G irradiation (100 mW cm–2) when tested in solution-processable bulk heterojunction solar cells with PC61BM as an acceptor material.

In recent times, solution-processable bulk heterojunction (BHJ) solar cells have gained interest among many academic and industrial research groups as they have the potential to be lightweight, flexible and cost effective alternatives to the traditional solar-cells, besides, possess capability for simple mass production.1 In the past 5 years, significant improvements have been achieved in the development of BHJ solar cells through a combination of efforts including the design of more active materials, morphology control, and device fabrication processes.2 Traditionally, semiconducting polymers such as poly(3-hexylthiophene) (P3HT) are used as electron donor materials and soluble fullerene derivatives such as [6,6]-phenyl-C61-butyric acid methyl ester (PC61BM) are used as electron acceptor materials.3 Recent reports, however, have described improvements in BHJ device performance by using small molecules and this in turn has drawn attention away from the more established polymeric semiconductors like P3HT.4 These small molecules possess an advantage of being able to be synthesised and purified as single molecular entities, thus removing the major bottlenecks of device variability that can arise from material inconsistencies.

As with polymeric structures, the design requirements of small organic semiconductors include broad absorption profile, high charge mobility, multiple reversible redox potentials and appropriate energy difference in the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO) energy levels. It has been established that low band gap materials can be generated by incorporating an electron donor–π-bridge–electron acceptor (D–π–A or D–A) motif within the structure.5,2a As previously discussed, reports of BHJ devices using small organic molecules as donor components have recently emerged and have resulted in devices exhibiting power conversion efficiencies (PCEs) in excess of 9%.5 Even though this progress is inspiring, ample scope still exists to develop novel light-harvesting materials that possess broad and efficient optical absorption, low HOMO energy levels (~5.0 to -5.5 eV) and adequate solubility required for solution processability.4 One successful strategy to fulfil such requirements is the exploration of D–A based compounds that show charge transfer absorption. Key examples of this approach have been reported by various research groups.7 Thus, there is considerable interest in exploring new D–A combinations and in synthesising molecules that can be processed from solution. In this communication, we report a strategy to investigate D–A design.

![Reference compound (R1)](Fig. 1) Newly designed S1

We chose a triarylamine end-group as the donor component, a dicyanovinyl acceptor group and introduced a planar, conjugated naphthalenediimide (NDI) functionality in order to vary (1) the optoelectronic and photovoltaic properties, and (2) the processability of small organic molecules (Fig. 1). The NDI unit has interesting physical and electronic properties, and chromophores based on the NDI framework have been used as active components for organic electronic applications.6,6c An additional advantage of the NDI functionality is that a variety of alkyl groups can be located on the nitrogen atoms, thus allowing tuning of solubility. In the present case, the -octyl group was the substituent of choice for producing highly soluble compound with excellent film forming properties without crystallization occurring in the film, see compound 2-((5-(5-(5-(4-(diphenylamino)phenyl)thiophen-2-yl)-2,7-dioctyl-1,3,6,8-tetraazoo-1,2,3,6,7,8-hexahydro-benzo[lmn][3,8]phenanthrolin-4-yli thiophen-2-yl)methylene)malono nitrile (S1); Fig. 1. Newly designed S1 is deemed to exhibit enhanced solubility and a large red
shift of lambda maximum when compared with the reference compound R1. It is worth mentioning that compound S1 is the first reported example where NDI unit has been incorporated in the D–A module for BHJ applications.

Compound S1 was synthesized in moderate to high yields by reacting 5-(9-(5-(4-(diphenylamino)phenyl)thiophen-2-yl)-2,7-dioctyl-1,3,6,8-tetraoxo-1,2,3,6,7,8-hexahydrobenzo[lmn]3,8]phenantralin-4-yl)thiophene-2-carbaldehyde with malononitrile, in acetonitrile:chloroform (1:1) solvent mixture at reflux with piperidine as a base (for detailed synthetic procedures, please see the electronic supplementary information (ESI†)). S1 was purified by conventional column chromatography and was fully characterized by means of high resolution mass spectrometry and NMR spectroscopy. High solubility of small molecular semiconductors is an essential feature for the fabrication of solution-processable BHJ devices and S1 meets this criterion as it displays high solubility in many routine solvents, such as chloroform (CHCl₃), chlorobenzene and toluene (for instance, 25 mg/mL in CHCl₃). Thermogravimetric analysis (TGA) revealed that S1 exhibits excellent thermal stability (>250 °C), a finding that strongly supports the fact that the compound will remain active and stable during the high temperature annealing of organic photovoltaic devices (Fig. S1, ESI†). Reference compound R1 was synthesized in our labs following the literature procedure.⁷c

The comparative UV–vis absorption spectral study of S1 and R1 was carried out in CHCl₃ solution (Fig. 2). The use of NDI functionality induces a strong red-shift in the absorption of compound S1 when compared with R1. Typically, S1 gives strong absorption band maximum (λ_max) at 632 nm (ε = 54,820 M⁻¹ cm⁻¹), and R1 gives λ_max at 514 nm (ε = 41,250 M⁻¹ cm⁻¹). With the use of NDI functionality we found >30% enhancement to the peak molar absorptivity of S1 compared with R1. A note of interest is the fact that we observed the same bathochromic absorption shift in thin film spectrum of S1 compared with that of R1 (Fig. 2).

The absorption profile of S1 in pristine state can be described as panchromatic with its absorption extending through the entire visible spectrum and tailing into the near infra-red region (400–1000 nm). This type of control in altering the absorption profile by incorporating a more strongly electron accepting moiety could lead to enhanced light harvesting properties in small molecular semiconductors.

Density functional theory (DFT) calculations using the Gaussian 09 suite of programs⁹ and B3LYP/6-311+G(d,p)//B3LYP/6-31G(d) level of theory indicated that the HOMO orbital density of S1 has a major distribution over triphenylamine donor functionality and the LUMO density was delocalized through both acceptor functionalities, NDI and malononitrile, with almost equal contribution (Fig. 3). Such a segregation of HOMO and LUMO densities is an ideal condition for the intramolecular charge transfer (ICT) transition and is attributed to the presence of a strong acceptor unit, of which NDI is an example, in the given D–A system.

Experimental estimation of HOMO energies was carried out using photo electron spectroscopy in air (PESA) and the LUMO energies were calculated by adding the bandgap to the HOMO values (see Fig. 4 for energy level diagram, and Fig. S2 for PESA curve).

The UV-vis spectra indicated that insertion of NDI unit reduces the band gap of S1 when compared with R1. PESA and UV–vis spectra indicated that the band gaps of these materials are all in the range required of donor materials for BHJ devices and are significantly narrower in magnitude than 1.8 eV measured for P3HT. The LUMO of S1 was the lowest of two due to the bathochromic shift observed in the thin film spectrum when compared with R1. The optical and electrochemical properties of both the materials are summarized in Table S1 (see ESI†).

![Fig. 2 Absorption spectra of compounds S1 and R1 (a) in CHCl₃ solutions and (b) for pristine as-casted films (films of S1 and R1 were spin-casted at 3000 rpm for 1 min to give a film thickness of ~55 nm).](image1)

![Fig. 3 Orbital density distribution for the HOMOs (lower) and LUMOs (upper) of S1. DFT calculations were performed using the Gaussian 09 suite of programs and B3LYP/6-311+G(d,p)//B3LYP/6-31G(d) level of theory.](image2)

![Fig. 4 Energy level diagram depicting the band gaps of S1 and R1 in comparison with P3HT and PC₆₇BM.](image3)
Initial screening into the potency of S1 as a donor material (p-type) was carried out by using it in a solution-processed BHJ device with conventional n-type soluble fullerene derivative PC61BM, and its performance under simulated sunlight and monochromatic light illumination was characterized. BHJ architectures typically deliver higher power conversion efficiencies (PCEs) by maximising the surface area of interface between the p- and n-type materials in the active layer. The blend solution of S1 and PC61BM was used to cast an active layer on top of the PEDOT:PSS surface. The BHJ device architecture used was ITO/PEDOT:PSS (38 nm)/active layer/Ca (20 nm)/Al (100 nm) where the active layer was a solution processed blend of S1 and PC61BM. For S1, a promising PCE of 3.01% was achieved when the film was spin-coated from a chlorobenzene solution as a 1:1 blend with PC61BM. By contrast, the maximum PCE obtained for a device based on R1 was 1.10%, when fabricated under similar conditions. The comparative current–voltage curves for the optimized blends of S1 and R1 with PC61BM are shown in Fig. 5.

The optimized devices based on S1 showed lower open circuit voltages ($V_{oc}$) than the devices based on R1. These values are consistent with the measured HOMO values where the more positive HOMO for compound S1 would predict a lower $V_{oc}$. Devices based on S1 showed higher photocurrent than the devices based on R1, a finding that is consistent with the observed bathochromic-shift in the absorption spectrum of S1 compared with R1. The BHJ devices based on S1 yielded promising performance and the photovoltaic cell parameters, $V_{oc}$, short circuit current density ($J_{sc}$), fill factor (FF) and PCE, reached 0.90 V, 7.90 mA/cm$^2$, 0.43 and 3.01%, respectively. Overall, the insertion of NDI functionality into the studied D–A module resulted in around 80% increase in the photocurrent density and enhanced the PCE by a factor of >2, thus promoting the use of an electron accepting and highly conjugated functionality as an interesting structural concept for the design and development of highly efficient BHJ materials. With regards to the processing conditions of blend solutions, it has been shown that there exists a strong relationship between the degree of crystallization and solvent selection on the cell performance. Our attempts to fabricate devices using low boiling solvents, such as CHCl$_3$, resulted in very poor photovoltaic performance. This was primarily due to low film quality. The use of high-boiling solvent is further preferable from a processing point of view.

The incident-photon-to-current conversion efficiency (IPCE) curves of the best BHJ devices based on these small molecules under monochromatic light are shown in Fig. 6. The analysis of the IPCE measurements of these BHJ devices reveals a significantly higher and broader peak IPCE of S1 (~42% @ 640 nm) compared to R1 (38% @ 515 nm), which is presumably caused by the synergistic effect of excellent light-harvesting and tuned electrochemical properties. The IPCE measurements reported here are comparable to the film absorptions of S1 and R1 which indicate that the current arises mainly from the p-type donor material. Note that S1 shows light harvesting and electrochemical properties that are largely superior to R1, which translates to higher conversion efficiency.

The active layer morphology was examined by an atomic force microscope (AFM) technique in tapping mode. The actual surface morphology of blend film of S1:PC61BM (1:1, w/w; as cast) is depicted in Fig. 7. The as casted film exhibited crystalline grains with a root-mean-square (RMS) roughness of 0.32 nm. The crystalline grains may be beneficial to ordered structure formation and charge transport in thin film. The hole only mobility in the BHJ blend film was measured by a space-charge limited current (SCLC) method with device structure of ITO/PEDOT:PSS/Active layer/Au. A hole mobility of 8.5x10$^{-6}$ cm$^2$ V$^{-1}$ s$^{-1}$ was obtained for the blend of S1 and PC61BM.

Even though the progress in the development of small molecular donors is on surge, the material reported in this paper has highlighted the potential for efficiency improvements in a given D–A system. The discovery of such materials exhibiting promising optoelectronic properties opens up a way to develop such motifs (based on the
insertion of highly conjugated, planar functionality) and paves the way for such materials to be used for other organic electronic applications such as organic field-effective transistors.

In conclusion, we have demonstrated the first use of a NDI functionality to improve the optoelectronic and photovoltaic properties of D–A small molecules in organic solar cells. In a direct comparison we have shown that the incorporation of this electron accepting, highly conjugated and planar system lowers the band gap of an oligothiophene dye, S1, and shows higher PCE compared with an analogue. R1. S1 showed excellent thermal stability and solubility in common organic solvents, strong optical absorption over the whole visible region, low lying HOMO energy level and their use in conjunction with alternative accepting functionalities.

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Notes and references

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