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Organic Photovoltaics: A Chemical Approach

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Abstract. The design of new organic functional molecules able to harvest sunlight and efficiently undergo photon to current energy conversion processes is at the forefront of chemical challenges. In this review the fundamental contribution of chemistry to the multidisciplinary field of organic photovoltaics is presented in a systematic way through the wide variety of organic compounds synthesized to be successfully used in photovoltaic devices.

Juan Luis Delgado obtained his PhD (2004) from the Universidad de Castilla-la Mancha under the supervision of Prof. Fernando Langa. Afterwards he joined the group of Prof. Jean-François Nierengarten, initially at the ECPM-CNRS (Strasbourg, France) and later on at the LCC (CNRS, Toulouse, France), where he worked as a postdoctoral fellow (2005-2006). He then joined the group of Prof. Nazario Martín in December 2006 as a “Juan de la Cierva” postdoctoral fellow. Currently he holds a “Ramón y Cajal” research contract at IMDEA-Nanociencia, where he is focused on the design of new carbon-based energy storing materials for the development of efficient organic photovoltaic devices.

Pierre-Antoine Bouit was graduated in 2005 from the Ecole Nationale Supérieure de Paris (ENSCLP, France). He obtained a PhD in 2008 at the Ecole Normale Supérieure de Lyon (ENS-Lyon, France) under the supervision of Dr O. Maury and Dr C. Andraud on the synthesis of near infrared absorbing dyes for nonlinear optics. Then he joined the group of Prof. Nazario Martín at UCM in 2008 as IMDEA post-doc. His current research interests are the synthesis of eTTF and [60]fullerene based materials for the preparation of Photovoltaic devices (Bulk HeteroJunctions and Dyes Sensitized Solar Cells).

Salvatore Filippone is a Ramón y Cajal researcher at Complutense University. He obtained his PhD degree at University of Pavia (Italy). His thesis was conducted under the supervision of Prof. Faita on metal catalyzed asymmetric synthesis. In 2000, he moved to Ecole Normale Superieure of Paris for a postdoctoral fellowship on biological evaluation of fullerene derivatives, working with Prof. Rassat and Prof. Bensasson. Since April 2002, he joined Prof. Nazario Martín’s group, working mainly on new chemical reactivity of fullerenes involving asymmetric approaches, and molecular wires.

Mª Ángeles Herranz gained a PhD from Complutense University of Madrid (UCM) in 2001. Then she moved to the USA to work as a postdoctoral researcher, first at the University of Miami (2001) and then at Clemson University (2002-2004). In March 2004 she joined the group of Prof. Nazario Martín at UCM, initially holding a Ramón y Cajal research associate position and since 2007 as Assistant Professor. She became an Associate Professor of Chemistry in 2009. Her current research interests primarily focuses on the chemistry and electronic properties of carbon nanostructures and in single-molecule electronics.

Nazario Martín is full professor of Chemistry at Complutense University and vice-Director of IMDEA-Nanoscience Institute.

His research interests span a range of targets with emphasis on the chemistry of carbon nanostructures in the context of electron transfer processes, photovoltaic applications and nanoscience. He is a member of the Editorial Board of Chemical Communications, a member of the international advisory board of The Journal of Organic Chemistry and ChemSusChem, and the Regional Editor for Europe of Fullerences, Nanotubes and Carbon Nanostructures. He is a member of the Real Academia de Doctores de España, a fellow of The Royal Society of Chemistry and the President of the Spanish Royal Society of Chemistry.

Introduction

Energy is currently the most important problem facing mankind. The “fire age” in which our civilization has been based from the very beginning is approaching to its end. Human being has been burning a wide variety of materials since earlier times and with the advent of carbon-based fossil fuels in the last two centuries, its combustion has become nowadays a major problem due to the huge amounts of carbon dioxide emissions produced all over the world. Because of the resulting pollution, global warming and degradation of the planet, a new era based on non contaminating renewable energies is currently a priority. In this regard, the Sun considered as a giant nuclear fussion reactor represents the most powerful source of energy available in our Solar system and, therefore, its use for providing energy to our planet is among the most important challenges nowadays in science. Actually, the energy received from Sun, calculated in 120,000 TW (5 % ultraviolet; 43 % visible and 52 % infrared), surpass in several thousands that consumed in the planet along a year.¹
and since the former silicon-based device prepared by Chapin in 1954 exhibiting an efficiency around 6%; different semiconducting materials (inorganic, organic, molecular, polymeric, hybrids, quantum dots, etc) have been used for transforming sun light into chemical energy. Among them, photo- and electro-active organic materials are promising due to key advantages such as the possibility of processing directly from solution, thus affording lighter, cheaper and flexible all-organic PV devices. Actually, these organic systems have received a lot of attention and a variety of excellent reviews and special issues have recently been published. They are mainly focussing on the race to increase efficiency as well as on those aspects mostly related with the understanding of those essential optical and electronic processes able to transform photon energy into electricity, thus providing a general overview on the state of the art in organic PV solar cells. Therefore, the aim of this review is to emphasize the contribution of chemistry to this field by covering those aspects related with the nature and properties of those type of compounds used to construct “all-organic” PV devices. Furthermore, the systematic organization of those compounds used for different purposes and tasks in organic solar cells should help to the chemical community to think about and to rationalize further alternative compounds which, eventually, result in a better performance of the PV devices.

Thus, this is not a comprehensive review on the chemistry of the so-called “all-organic” or “excitonic” (absorption of light results in the formation of excitons, basically a neutral electron-hole pair) solar cells but, on the contrary, a presentation of the most relevant chemical compounds which currently define the different classes of “all-organic” PV cells. Thus, we have organized this review in four main parts: i) Basic processes in organic solar cells; ii) Bulk-heterojunction plastic solar cells based on semiconducting π-conjugated polymers and fullerenes; iii) Molecular bulk-heterojunction solar cells based on π-conjugated oligomers or dyes and fullerenes; iv) Other carbon nanostructures, namely endohedral fullerenes and carbon nanotubes (CNTs). Actually, the presence of pristine fullerenes (C_{60} and C_{70}), or most commonly a fullerene derivative (namely PCBM: 1-(3-methoxycarbonyl)propyl-1-phenyl[6,6]C_{61}), is a common feature in sections ii) and iii). Therefore, it is not surprising the recent interest in other available related carbon nanostructures such as endofullerenes and CNTs, which is covered in iv).

i) Basic Processes in Organic Solar Cells

Organic solar cells are constituted by semiconducting organic materials formed by contacting electron donor and acceptor compounds (p/n type). A further improvement in the construction of organic photovoltaic devices (OPV) consists on the realization of interpenetrated networks of the donor and acceptor materials. In such bulk heterojunction (BHJ) solar cells, the dramatic increasing of contacting area between D/A materials leads to a significant increasing of the number of generated excitons as well as their dissociation into free charge carriers and hence on the power conversion efficiency (Figure 1).

Transformation of solar energy into electricity occurs through a series of optical and electronic processes which basically involves: i) optical absorption of sun light and formation of the exciton; ii) exciton migration to the donor-acceptor interface; iii) exciton dissociation into charges (electron and holes), and iv) charge transport and collection at the electrodes. All these steps are not totally understood at present and a number of research groups are currently dedicated to unravel essential aspects in the search for better energy transformation efficiencies. In contrast to inorganic semiconductors which upon light excitation form free electron and hole carriers, organic compounds form excitons whose dissociation into free carriers is not straightforward. Due to their low dielectric constants, it requires reaching the donor-acceptor interface to dissociate into free charges. The driving force for this exciton dissociation being provided by the energy difference between the molecular orbitals of the donor and acceptor.

The transport of the thus generated free charges toward the electrodes represents another important issue. Organic materials typically show charge carrier mobilities significantly lower (around 10^{-3} to 1 cm^2 V^{-1} s^{-1} ranging from amorphous to crystalline materials) than those of inorganic semiconductors (around 10^{-5} to 10^{-3} cm^2 V^{-1} s^{-1}). Since charge carriers mobility is strongly dependent on the molecular organization of the material, in order to achieve better efficiencies, a good control on the morphology of the donor-acceptor materials at a nanometer scale is necessary. Furthermore, an efficient charge carriers mobility is essential to prevent charge recombination (geminal or bimolecular) processes which result in lower energy conversion efficiencies.

In summary, an appropriate choice of the donor and acceptor materials is critical to ensure a good match between them in terms of optical, electronic and morphological properties, which eventually determine the effective photocurrent and performance of the PV device. Therefore, the rational design of new materials able to improve some specific demands within the PV device is critical for the successful development of competitive solar cells. In the following sections, we will discuss those materials mainly used in the preparation of “all-organic” solar cells according to their chemical nature and composition.

Figure 1. Basic chemical components for a typical Organic Photovoltaic (OPV) Solar Cell.

ii) Bulk-heterojunction “plastic” solar cells
The term “plastic solar cells” has been coined for those photovoltaic devices using polymers in their constitution. Thus, although a variety of donor and acceptor polymers have previously been used for constructing PV devices, the highest efficiencies have been achieved by mixing electron donor polymers with fullerenes as acceptors. Polymer:fullerene (BHJ) based solar cells are considered suitable candidates to obtain low-cost renewable energy from a large-area, flexible, plastic material.

Fullerenes and their derivatives possess important electronic properties like small reorganization energy, high electron affinity, ability to transport charge and stability, that makes them (see below) one of the best candidates to act as electron acceptor components in BHJ photovoltaic PV devices. On the other hand, MDMO-PPV [poly[2-methoxy-5-(3′,7′-dimethyloctoxy)]-1,4-phenylenevinylene] and P3HT [poly(3-hexylthiophene-2,5-diyl)] have been by far the most studied π-conjugated polymers as donor components in BHJ devices. Actually, both polymers have thoroughly been studied from a morphological and electronic standpoint to form BHJ devices by mixing with fullerene derivatives, namely PCBM (19, Figure 4). In fact, the best fullerene/polymer combination has been obtained with PCBM/P3HT (regioregular) mixtures, reaching energy conversion efficiencies in the range of 5%.

In the following, we will briefly describe the most relevant achievements both in the chemistry of polymers as well as on the fullerene chemical modifications for PV purposes.

### a) Polymers for BHJ Photovoltaics

Polymers endowed with fullerenes is a broad interdisciplinary field of interest for a variety of purposes. However, photovoltaics are among the most realistic applications for polymer/fullerene blends.

The chemistry of π-conjugated polymers has been focused on tuning their energy (HOMO and LUMO) levels in order to have a good control on the band gap of the polymer. This is a key issue to modulate the light harvesting properties of the polymer [low band gap polymers (lower than 1.5 eV) absorb in the visible region] as well as the right energy match with the acceptor moiety which, eventually, controls the open circuit voltage (Voc) values, and hence the cell power conversion efficiency.

Since π-conjugated polymers constitute an specific field by their own right, we will discuss the trends currently followed in polymer chemistry to improve the performance of those “classical” MDMO-PPV and P3HT. Some of the most relevant polymers synthesized so far are shown in Figure 2. All of them involve the use of alternating donor and acceptor moieties within the polymer backbone since they allow a fine tuning of the band gap of the polymer, as well as an improvement of planarity and hence of the mobility of the charge carriers. Thus, a variety of soluble electroactive carbo- and heterocycle derivatives such as fluorene, thiophene, cyclopentadithiophene and carbazole have been used as electron rich units, whereas benzo[1,2,5]thiadiazol-7-ylthiophen-2-yl)benzo[c][1,2,5]thiadiazol-7-yl]thiophen-2-yl)-9-octyl-9H
carbazole, HXS-1) bearing two octyloxy chains on the benzothiazole ring and an octyl chain on the carbazole ring. The synthesis of HXS-1 is shown in Scheme 1 to illustrate the synthetic strategy followed for the preparation of this kind of polymers. Thus, the coupling reaction of 4,7-dibromo-5,6-bis(octyloxy)benzo-2,1,3-thiadiazole (10) with 4,4,5,5-tetramethyl-2-thiophen-2-yl)-1,3,2-dioxaborolane (11) using Pd(PPh₃)₄ as the catalyst precursor and aqueous NaHCO₃ and THF as solvent, led to compound 12 in moderate yield.

Bromination of 12 with N-bromosuccinimide afforded 13 whose further Suzuki-Miyaura polycondensation with 2,7-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborol-an-2-yl)-9-octyl-9H-carbazole (14) using Pd(PPh₃)₄ as the catalyst precursor in a biphasic mixture of THF-toluene (5:1)/aqueous NaHCO₃ formed polymer HXS1 as a black solid in good yield.¹⁸

Semiconducting polymers based on alternating thieno[3,4-b]thiophene and benzothiadiazole units such as 7 and 9 show an excellent photovoltaic behaviour. The stabilization of the quinoid structure from the thieno[3,4-b]thiophene moiety leads to remarkable low band gap polymers (around 1.6 eV). A further optimization has been achieved with the new PTB7 in which the presence of the fluorine atom leads to a lower HOMO level, thus enhancing the Vₜₚ parameter. With an average molecular weight around 97.5 KDa and a polydispersity index of 2.1, this soluble polymer exhibits a strong absorption from 550 to 750 nm. Interestingly, by blending this polymer with PC₇₅BM, which nicely complements the absorption in the visible range, has allowed preparing a PV device exhibiting an energy conversion efficiency of 7.40%. This value represents the highest efficiency reported so far for a polymeric solar cell.¹⁸

![Scheme 1. Synthesis of polymer HXS1.]()

Since the development of new π-conjugated donor polymers for improving the efficiency in BHJ photovoltaic devices is currently a broad and very active field, out of the scope of this general presentation, for further details the reader is referred to some of the excellent monographs available in the literature.²⁰

b) Chemically Modified Fullerenes for BHJ Photovoltaics

The most widely used configuration of polymer solar cells is based on the use of a fullerene derivatives as acceptor component. Indeed, fullerenes have been demonstrated to be the ideal acceptor because of their singular electronic and geometrical properties and for the ability of their chemically functionalized derivatives to form a bicontinuous phase network with π-conjugated polymers acting as electron conducting (n type) material.

A variety of chemically modified fullerenes were initially synthesized for blending with semiconducting polymers (namely PPV derivatives) and to prepare photovoltaic devices. These fullerene derivatives were covalently linked to different chemical species such as electron acceptors,²¹ electron donors,²² π-conjugated oligomers,²³ dendrimers,²⁴ etc. (Figure 3). However, in general, the obtained blends resulted in PV devices exhibiting low energy conversion efficiencies.²⁵

![Figure 3. Some examples of modified fullerenes bearing different organic addends (15-18) used to prepare photovoltaic devices.]()

The best known and most widely used fullerene derivative as acceptor for PV devices is [6,6]-phenyl-C₆₁ butyric acid methyl ester 19 (PCBM), firstly prepared by Hummelen and Wudl in 1995.²⁶ Since its first reported application in solar cells,²⁷ it has been by far the most widely used fullerene,
being considered as a benchmark material for testing new devices.

**Figure 4.** PCBM and other representative PCBM analogues involving a thienyl group ([60]ThCBM) or a [70]fullerene (PC_{71}BM).

This has led to the synthesis of many other PCBM analogues in an attempt to increase the efficiencies of the cells by improving the processability, the stability or PV parameters such as the open circuit voltage (Voc) by rising the LUMO energies of the fullerene acceptor (Figure 4). In this regard, only small shifts (< 100 meV) of the LUMO level have been obtained by attaching a single substituent on the fullerene sphere, even by using electron-donating groups. In contrast, significantly higher Voc values have been achieved through the polyaddition of organic addends to the fullerene cage (~100 mV raising the LUMO per saturated double bond). Recently, an externally verified power-conversion efficiency of 4.5% has been reported by Hummelen et al. employing a regiosomeric mixture of PCBM bisadducts as result of an enhanced open-circuit voltage, while maintaining a high short-circuit current (J_{sc}) and fill factor (FF) values. The cyclopropanation reaction to superior fullerenes to form PCBM analogues is more complex than for C_{60}. Indeed, the less symmetry and the presence of more than one reactive double bond is often responsible for the formation of regiosomeric mixtures. Nevertheless, the loss of symmetry of C_{60} induces a stronger absorption, even in the visible region. As a result, PC_{71}BM is considered a suitable candidate for more efficient polymer solar devices. Moreover, such devices performed the highest verified efficiency determined so far in a BHJ solar cell, with an internal quantum efficiency approaching 100%. Analogously, PC_{84}BM has been obtained as a mixture of three major isomers. The stronger electron affinity and the diminished solubility gave rise, however, to poor power conversion efficiencies.

Although the PCBM is the acceptor that best performances guarantees to the moment, it does not mean that they are necessarily the optimum fullerene derivatives. Therefore, a variety of other fullerene derivatives have been synthesized in order to improve the device efficiency or to achieve a better understanding on the dependence of the cell parameters from the structure of the acceptor.

Among the different modified fullerenes prepared so far, diphenylmethanofullerene 22 (DPM12) prepared by Martín et al. is another successful type of methanofullerene endowed with two alkyl chains to drastically improve the solubility of the acceptor in the blend and reaching efficiencies in the range of 3% (Figure 5). Although the LUMO energy level for DPM12 is the same as that for PCBM, an increase for the Voc of 100 mV for the DPM12 over PCBM has been observed. This is currently an important issue for improving the design of future fullerene-based acceptors. Devices based on dihydroxynaphthylfullerene benzyl alcohol benzoic ester (23) synthesized by Fréchet et al. reported one of the highest power conversion efficiency (PCE) (up to 4.5%) for a non PCBM based polymer–fullerene solar cell. An alternative molecular approach to control both the length scale of polymer/fullerene phase segregation and the electron mobility in the fullerene network is based on the employ of a self assembly fullerene shuttlecock derivative (24), able to form one-dimensional wirelike domain within the active layer.

**Figure 5.** Different modified fullerenes used as successful acceptors for PV devices

Although fullerene dimers are known to exhibit low solubilities, recently a series of soluble homo (25: C_{60}-C_{60} and 26: C_{70}-C_{70}) and heterodimers (27: C_{60}-C_{70}) formed from C_{60} and C_{70} derivatives have been prepared and explored as less-known acceptors for PV devices. The synthesis of these pyrroolidino-pyrazolino fullerene dimers has been carried out in a straightforward manner from the formyl-containing pyrazolino[60] or [70]fullerene, which is used as the aldehyde component in the subsequent pyrrolidine formation by 1,3-dipolar cycloaddition of the respective azomethyne ylide with the fullerene. As expected, [70]fullerene dimers showed better absorption in the visible spectrum than the analogues of [60]fullerene. Interestingly, non optimized BHJ solar cells formed by blending with P3HT exhibited quantum conversion efficiencies of 37 %, with a PCE of 1.0 % (Figure 6).
Figure 6. Homo- and hetero-dimers of C₆₀ and C₇₀ (25-27) and bis-cycloadduct 28 bearing two indene units.

A remarkable bis-adduct fullerene derivative formed by two indene units covalently connected to the fullerene sphere of C₆₀ (28) has recently been reported by Hou and Li. Interestingly, the presence of two aryl groups improves the visible absorption compared to the parent PCBM, as well as its solubility (> 90 mg/ml in chloroform) and the LUMO energy level, which is 0.17 eV higher than PCBM.

Surprisingly, PV devices formed with P3HT as semiconducting polymer revealed PCE values of 5.44 % under illumination of AM1.5, 100 mW/cm², thus surpassing to PCBM which afforded an efficiency of 3.88 % under the same experimental conditions.

A major drawback in the synthesis of bis-adducts of fullerenes is that the products formed are constituted by a mixture of regioisomers which are not separated because of the experimental difficulties. This fact, however, does not seem to have a strong influence on the PV parameters and power conversion efficiencies. Nevertheless, from a chemical viewpoint, the synthesis of isomerically pure bis-adducts and multiaadducts of fullerenes is still an open question which should be properly addressed by the chemical community.

Although some of the fullerene derivatives prepared so far exhibit good performances in PV devices, the synthesis of new fullerene derivatives with stronger visible absorption and higher LUMO energy levels than PCBM is currently a challenge for all those chemists engaged in the chemical modification of fullerenes for PV applications.

c) Double-Cable Approach for BHJ Photovoltaics

This strategy is based on the appendage of the acceptor moiety to the π-conjugated polymeric skeleton with the aim to maximize the interface between the two active components, thus preventing the phase separation between them. At the same time, such approach is expected to promote an efficient charge transport to the electrodes through a nanoscale homogeneous distribution of the domains of the donor and the acceptor in the blend.

The first example of a PV cell prepared by using the double-cable approach was carried out by spin coating from poly(p-phenylenevinylene) and poly(p-phenyleneethynylene) covalently connected to methanofullerene units. Photovoltaic parameters (Voc = 0.83 V; Jsc = 0.42 mAcm⁻²; FF = 0.29 under AM1.5 conditions) resulted to be quite promising.

Interestingly, the realization of effective double-cable polymers brings the p/n heterojunction at the molecular level, thus allowing a better control from a chemical viewpoint. However, to date, and despite the beauty of this chemical design, double-cable polymers have yielded poor energy conversion efficiencies. One of the reasons to account for this finding has been the low fullerene loading of the material. Therefore, new polymers have been synthesized such as PTF (29) or PBTC-F (30) with a higher fullerene content (up to 63 wt %) (Figure 7). Other studies involving polythiophenes followed to this one with slightly better performances and efficiencies around 0.60 %.

An interesting alternative approach is based on the use of well-defined rod-coil block copolymers consisting of P3HT donor and C₆₀ acceptor chromophores (P3HT-b-P(SₓAᵧ)-C₆₀) (31) as “surfactant”. The control of the interface morphology of the P3HT:PCBM donor–acceptor phase domains within the composite resulted in an improvement of the efficiency with respect to the analogous solar cell fabricated without the “surfactant”.

Figure 7. Some representative examples of fullerene-containing polymers used to the double-cable approach.

iii) Molecular Bulk-Heterojunctions

Despite the outstanding results obtained from BHJ solar cells formed from fullerenes and polymers, the use of commercially available P3HT displays a number of troubles associated to its synthesis, purification, and inherent electronic properties. Factors such as polydispersity, regioregularity and molecular weight have a strong influence on the final performance of the photovoltaic device, which indicates that the control over P3HT structure and purity is an important issue for controlling organic photovoltaics. Concerning its electronic properties, it has been calculated that P3HT can only absorb 46% of the
available solar photons,\textsuperscript{49} owing to a bandgap of 1.90 eV and a relatively narrow absorption band. A promising solution to overcome these limitations could be the use of soluble small conjugated molecules instead of polymers. This strategy would avoid the above mentioned structural problems inherent to polymers, and allow the use of several well-defined small molecules whose electronic properties can be tuned by chemical synthesis, in order to obtain a better photovoltaic performance. Following this approach, in the last years, a number of molecules\textsuperscript{18} such as oligomers, dendrimers, dyes and porphyrines have been used as molecular donors blended with pristine [60]fullerene or [60]fullerene derivatives, in order to prepare new and efficient photovoltaic devices.

a) Conjugated oligomers and dendrimers as donors in BHJ-PV devices

The synthesis and study of new organic monodisperse π-conjugated oligomers is a hot topic in photovoltaics.\textsuperscript{30} Oligothiophenes are among the most promising oligomers, due to their exceptional electronic properties and easy tuning of their properties by introducing functional substituents. Several substituted oligothiophenes have been described and studied for organic photovoltaics. However, most of them show a planar geometry, which tends to aggregate due to π-π stacking interactions. Sun \textit{et al.}\textsuperscript{21} described recently the preparation of new X-shaped oligothiophenes (32-35) as electron donors in BHJ-PV devices blended with PCBM (Figure 8). The authors observed that both the open circuit voltage ($V_{OC}$) and the short circuit current ($J_{SC}$) are highly dependent on the oligothiophene chain length. While increasing the number of oligothiophene rings, the absorption was red-shifted and broadened, therefore the light harvesting properties and the PCE increase from 0.008 % to 0.8 %.

![Figure 8](image8.png)

**Figure 8.** Chemical structures of four X-shaped oligothiophenes.

Other interesting oligothiophene-based compound, was recently described by Tamayo \textit{et al.}\textsuperscript{52} namely oligothiophene diketopyrrolo-pyrrole (36) (SMDPPEH, Figure 9). This appealing oligothiophene is a low bandgap well-defined molecule that displays intense absorption bands at 650 and 720 nm. Blends of this material and PC$_7$BM were prepared in different ratios and the results indicated the highest efficiency described so far for a molecular bulk heterojunction (PCE of 3 \%) being 0.75 V the $V_{OC}$ and 9.2 the $J_{SC}$. Interestingly, the presence of the ethyl-hexyl groups increment the solubility and processability and, as a consequence, no microscale phase segregation was observed in the films.

![Figure 9: Chemical structures of SMDPPEH and C$_{71}$-PCBM](image9.png)

**Figure 9:** Chemical structures of SMDPPEH and C$_{71}$-PCBM

On the other hand, conjugated dendrimers constitute a new kind of well-defined molecular materials with monodisperse structures which are highly controllable due to precise synthetic approaches. In this context Ma \textit{et al.}\textsuperscript{53} have reported on the photovoltaic properties of BHJ-PV devices prepared with blends of PCBM and various generations of dendritic oligothiopenes (DOTs) bearing from 9 to 90 thiophene units (Figure 10 ). Owing to the lower HOMO levels of the DOTs, the devices showed remarkable $V_{OC}$ values (0.93-1.06 V), which are higher by 0.3-0.4 V in comparison with P3HT-PCBM. Other photovoltaic parameters like $J_{SC}$ and external quantum efficiency (EQE), showed a different trend, being their values increased while increasing the size of the DOT. These photovoltaic parameters lead to a maximum power conversion efficiency of 1.72 \% for compound 38, which is composed by 42 units of thiophene.

![Figure 10: Chemical structures of the DOTs containing 21 and 42 thiophene units.](image10.png)

**Figure 10.** Chemical structures of the DOTs containing 21 and 42 thiophene units.

b) Organic Dyes as donors in BHJ-PV devices

Following the use of monodisperse and well defined molecules for the preparation of BHJ-PV devices, in the last years a new and interesting approach has emerged in the photovoltaic field. Organic dyes are single molecules which display high absorption extinction coefficients and electronic properties which can be easily tuned by chemical synthesis.
Roquet et al. have described the synthesis of triphenylamine (TPA) based compounds (39–41) (Figure 11) bearing several units of dicyanovinylene. Photovoltaic studies on these compounds blended with C60 in bulk and bilayers heterojunctions reveal a high value of $V_{OC}$ in the BHJ-PV devices due to their high oxidation potential. Owing to an intramolecular charge transfer band from the TPA core to the dicyanovinylene acceptor groups, and to the above mentioned high $V_{OC}$, an improvement of the power conversion efficiency was observed, while the number of dicyanovinylene acceptor units increase in the molecule (0.49-1.17%).

Figure 11. TPA derivatives bearing one or more dicyanovinylene groups.

Kronemberg et al. have recently described the photovoltaic study of blends of merocyanines and PCBM. These dyes exhibit intense absorption bands ($\varepsilon$ up to 1.4 x 10^5 M^{-1}cm^{-1}) at different wavelengths in the visible spectral region. For several merocyanine dyes (42) power conversion efficiencies around 1% were obtained under standard AM 1.5 conditions. Although the best performance was obtained for MD 304 due to a higher $J_{SC}$ value (6.3 mAcm^{-2}) and a $V_{OC}$ of 0.76 V which lead to a PCE of 1.74%. In order to harvest light in the NIR region, where almost 50% of the sunlight power is displayed, Silvestri et al. described the synthesis of two new squaraine derivatives (43,44) (Figure 12) displaying intense absorption from 700 nm to 800 nm in solution. Photovoltaic studies of blends of these dyes with PCBM revealed a power conversion efficiency up to 1.3%. To the best of our knowledge this is the first efficient device prepared using NIR light harvesting molecular dyes. In order to improve the efficiency of squaraine/PCBM based BHJ, Mayerhöffer et al. studied the charge carrier mobility of a series of squaraine dyes bearing a dicyanovinyl unit, preparing OFET devices. The authors showed that the most efficient photovoltaic device (PCE=1.79%, $J_{SC} = 12.6mAcm^{-2}$) presented good hole-carrier mobility. This remarkable PCE value was attributed to well packed domain of squaraine and PCBM. Besides the introduction of the dicyanovinyl moiety favorably changes the absorption properties of these squaraine dyes, and therefore the capability to harvest solar energy. This study illustrates nicely the importance of tuning both absorption properties and morphology in molecular BHJ cells, as it has previously been shown with polymer-based devices.

Figure 12. 42) Merocyanine MD 304; 43 and 44) Squaraine derivatives used for preparing PV devices by blending with PCBM.

Borondipyromethene (BODIPY) dyes (45, 46) are another kind of organic dye widely used for their outstanding absorption and fluorescence properties in the visible range. Rousseau et al. took advantage of the ease of chemical modification on the bodipy core to tune the optical properties of the dyes and reached 1.3% of efficiency in bodipy:PCBM BHJ devices (Figure 13). Using the same family of dyes, these authors built BHJ devices with both Bodipy donors which display complementary absorption properties in the visible range ($\lambda_{max1}= 590$ nm, $\lambda_{max2}= 675$ nm). This strategy has allowed to solve the problem of the narrow absorption of these organic molecular dyes. An efficiency of 1.7% has been described with this multidonor BHJ, which represents an increase of 0.4% compared to the best efficiency described with a single Bodipy donor. This significant increase nicely illustrate the importance of harvesting sunlight in the whole solar spectrum in order to obtain more efficient photovoltaic systems.

Figure 13. Bodipy dyes used for PV devices by blending with PCBM.

Using a similar strategy, some of us have described a new organic salt (49) by association of two NIR absorbing cyanine dyes (47,48), which exhibit a good solubility in organic solvents (Figure 14). As a result of the combination of the two cyanines, the complex is able to absorb light in a broad range of the solar spectrum (650-950 nm) displaying outstanding light harvesting properties in the NIR region with giant extinction coefficients. In preliminary experiments, the sun to power conversion efficiency obtained for the cyanine-cyanine complex:[60]PCBM (PCE = 0.4%) was higher than the efficiency observed for blends of PCBM:cyanine-47 or PCBM:cyanine-48. This result indicates that the new dye is an appealing system for further studies as light harvester for photovoltaics.
iv) Organic Solar Cells with Innovative Carbon Nanostructures

As mentioned above, one of the fundamental limitations of OPV device PCEs has been the molecular orbital offset between the available donor and acceptor materials. Toward this goal, new acceptor materials systems are being designed and synthesized to optimize the donor/acceptor energy levels in order to increase the device photovoltage. From the variety of nanoscale size shapes with fascinating properties that carbon is able to provide, carbon nanotubes (CNTs) and endohedral fullerenes are of current interest because of their promising applications in next-generation organic photovoltaics, due to their outstanding intrinsic electronic properties.

a) Endohedral fullerenes

Novel acceptor materials with LUMO energies closer to that of the donor, such as the derivatives of metallo endohedral fullerenes M3N@C80, where the choice of metal (M = Sc, Y, Gd, Tb, Dy, Ho, Er, Tm, Lu) allows the fine tuning of the LUMO levels, as well as, the absorptive coefficients of the fullerene species, offer the possibility of considerably increase the PCE of OPV devices.

In preliminary investigations, Sc3N@C80 and Y3N@C80 shown to be applicable as part of electron-donor/acceptor systems in combination with powerful donors such as ferrocene or π-extended tetrafulvalene derivatives (exTTFs) (Figure 15) (50 and 51 respectively). The photophysical investigations revealed a significant stabilization of the radical ion pair state of Sc3N@C80-ferrocene when compared to an analogous C60-ferrocene dyad.

Figure 15. Optimized structures of a ferrocene-\(I_x\)-[5,6]-Sc3N@C80 fulleropyrrolidine (50) (left) and the down isomer of an exTTF derivative of [6,6]-I0-Y3N@C80 (51) (Courtesy of Prof. J. M. Poblet).

These positive results have been recently complemented with the preparation of BHJ based solar cells considering a Lu3N@C80 fulleroid modified with a phenyl butyric acid hexyl ester (52, PCBH) solubilizing group and, blended with P3HT. In photophysical experiments, the efficient charge transfer between P3HT and the Lu3N@C80-PCBM were confirmed. The optimized P3HT/Lu3N@C80-PCBM active layer morphology has resulted in OPV devices with \(V_{oc}\) values as high as 890 mV (260 mV above reference devices made with C60-PCBM) and a similar photocurrent and fill factor compared to P3HT/C60-PCBM reference devices (Figure 16). The higher overall PCE (4.2%) of the devices fabricated is attributed to a better positioned LUMO level that captures more of the energy associated with each absorbed photon. Further investigations, considering comparative analysis of the film absorption and X-ray diffraction, indicate that the optimized blend ratio in a P3HT matrix differs from a molecular equivalent of an optimized P3HT/C60-PCBM active layer, and this is related to the physical differences of the C80 fullerene.

Although additional investigations are necessary in the near future, the potential of endohedral metallofullerene species to enable optimized OPV devices is certainly very promising.

b) Carbon nanotubes

Since their discovery CNTs have been a constant source for scientific motivation and in recent years have emerged as an outstanding class of carbon into OPVs. The most important strategies for implementing CNTs in OPV devices consider the incorporation of single-walled carbon nanotubes (SWCNTs) in dye-sensitized solar cells (DSSC), their combination with...
The most significant organic compounds prepared so far to be used in all-organic solar cells have been presented in a systematic way. Bulk heterojunction plastic solar cells constituted by a semiconducting electron donor π-conjugated polymer and a fullerene as the acceptor moiety are currently the most promising devices for practical applications. The most remarkable achievements in π-conjugated polymers have been presented as well as those modified fullerenes which, by blending with the appropriate polymers, have afforded the best energy conversion efficiencies reported so far, reaching values as high as 7.4 %.

Covalently connected π-conjugated polymers-fullerene forming the so-called double-cable approach represents an elegant methodology to prepare new materials which, up to now, has afforded comparatively very low energy conversion efficiencies.

Molecular bulk heterojunction represents an interesting
alternative to the use of polymers for the preparation of PV devices. These compounds have been classified in those systems formed by π-conjugated oligomers or dendrimers and fullerences, and the most recent dyes blended with fullerences.

Other carbon nanostructures such as endohedral fullerenes and carbon nanotubes are promising materials for PV devices. Although so far only a couple of examples are known on PV devices involving endofullerenes, the high energy conversion efficiency (4.2%), similar to that found for the efficient [60]PCBM make these materials good candidates for further studies. Obviously, the main problem of these endofullerenes is currently related with their availability in larger scale. Regarding the use of CNTs, it is important to note that the fate of these carbon nanoforms for PV devices depends primarily on the separation of the metallic and semiconducting types of CNTs which typically are mixed in the commercially available materials. Although still low efficiencies have been reported with CNTs, the huge potential of these carbon allotropes makes them very appealing in this field.

A most important matter in all the above materials for improving the cell performance is the precise control of the energy levels of donor and acceptor components as well as the network morphology to maximize the mobility of the charge carriers. In this regard the realization of nano-structured materials by means of a bottom-up supramolecular organization is currently in its infancy and, therefore, more work is need for improving the materials used for PV applications.

In summary, organic solar cells are call to play an important role for satisfying the increasing energy demanding of our society. Because of their features of low cost, flexibility and lightness, these new PV cells – which should nicely complement the commercially available silicon cells – are very appealing for a variety of new practical purposes. The outstanding achievements obtained so far put these solar cells closer to the market.

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