

OSC: Statement of Work (Reviews and Background)

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INTRODUCTION

Solar cells are an essential component in the process of transitioning the energy systems of the globe away from the use of fossil fuels and toward the use of renewable technologies. This movement is being driven by the goal of reducing the world's reliance on nonrenewable sources of energy. This shift is currently under way and will be completed over the course of the following several decades. The overall amount of energy that comes from the sun is many thousands of times greater than the amount of energy that is required to meet the demands of the population of the whole globe. This means that the sun is the primary source of energy for the planet. The sun is responsible for the direct production of solar energy, as well as the indirect production of other types of renewable energy, such as wind, wave, hydroelectricity, and biomass. In addition, the sun is directly responsible for the creation of solar energy. One example of this type of technology is photovoltaics, which stands out due to its capacity to transform sunlight into an electric current in a straightforward manner. There are a number of various methods in which the technology that underpins solar cells can be put into practice. Because they can be manufactured on flexible substrates from solutions and at low temperatures with the help of roll-to-roll manufacturing processes that are reasonably inexpensive, organic solar cells represent an alternative to the more typical silicon sun cells. Because of this, organic solar cells are an appealing alternative to the more conventional silicon solar cells. In addition to this, not only are they simple to carry, but they can also be manufactured without the use of any raw materials that may pose a risk to the environment. Organic solar cells have a shorter lifetime and a poorer power conversion efficiency (PCE) compared to silicon solar cells. Organic solar cells also degrade more quickly. This is due to the fact that organic solar cells are constructed using a distinct kind of material. In contrast, organic solar cells have a power conversion efficiency (PCE) of around 5% and a lifetime of approximately 5 years, whereas typical silicon photovoltaic panels have a PCE of approximately 17% and a lifetime of nearly 25 years.

Because of the multiple benefits that they offer, organic solar cells (OSCs) with the bulk heterojunction (BHJ) active layer have caught the curiosity of a substantial number of people. This is due to the fact that organic sun cells (OSCs) are relatively new. A low weight, a low degree of opaqueness in appearance, and a high degree of mechanical flexibility are only some of the advantages that come with this material. It is now possible for organic solar cells, also known as OSCs, to attain power conversion efficiencies (PCEs) of more than 18 percent because to the fast advancement of organic photovoltaic materials and device engineering. The fact that OSCs have been able to attain such high levels of power conversion efficiency is what made this accomplishment conceivable. Despite the significant advancements that have been achieved in PCE development, the low operational stability of OSCs restricts the scope of the practical applications that may be carried out with them. This is the situation in spite of the substantial effort that has been made in bettering PCE. It is important to take note of the fact that the morphology of the photoactive layer degrades with prolonged exposure to a variety of environmental components (such as light, oxygen, and heat), since this continues to be a primary barrier in the path toward the commercialization of OSC technology.

LITERATURE REVIEW

Xiang Xu will have accomplished all of these by the year 2021. The majority of the progress made in organic solar cells in recent years may be credited to the discovery of new non-fullerene acceptors (NFAs). These acceptors have allowed for tremendous advancement in the field. "non-fullerene acceptors" is what "NFAs" stands for in abbreviated form. Although it has been reported that OSCs' power conversion efficiency has grown to around 18%, the technology has not yet attained commercialization due to concerns over its stability. This piques the curiosity of scientists, who proceed to conduct more research on the topic. As a result, we will investigate the processes that are responsible for instability in OSC devices, offer a comprehensive overview of the

advancements that have been made to OSC stability over the course of the previous several years, and suggest a number of remedies that can improve the stability. Even though highly stable OSCs have been recorded with a lifespan that may be projected to encompass several decades, the most obvious disadvantage of these OSCs is their inadequate PCE. This is the case even though the longevity of these OSCs has been reported. Therefore, it is absolutely necessary to do research on OSCs that are both exceptionally effective and astonishingly constant. We think that the suggestions gained from this study, which should assist to stabilize OSCs, will make it easier in the future to commercialize technologies related to OSCs, and this is something that we are excited about.

S. Peng Wang (2020) Organic solar cells, also known as OSCs, have attracted a significant deal of interest over the course of the last several decades due to the multiple desirable qualities that they possess. Their cheap cost, versatility, semitransparency, nontoxicity, and limited environmental effect make them excellent for large-scale roll-to-roll production. Other properties that make them good for this type of production include their semitransparency. In addition to this, the manufacture of organic solar cells on a massive scale is feasible. In recent years, there has been substantial progress achieved in the field of organic semiconductor capacitors, sometimes known as OSCs. In addition to a mix of high-performance active layer materials, electrodes, and interlayers, these organic semiconductor capacitors have innovative device topologies. The development of active layer materials, which has included the introduction of novel acceptors and donors, is a significant element in the huge growth in PCE in OSCs, and it is one of the most essential factors. It has been demonstrated that the conversion rate from input power to output power has improved, leading to a higher overall efficiency. In this line of study, a comprehensive exploration of the realm of high-performance acceptors is undertaken. Acceptors may be broken down into several different categories, such as polymeric non-fullerene acceptors (NFAs), tiny molecule acceptors, and fullerene derivatives. During this period, we will also demonstrate high-efficiency donor materials that have been adapted specifically for fullerene- and NFA-based OSCs. In addition, this article discusses the most recent findings from research on ternary and tandem OSCs, both of which have been inspired by the continuous development of donor and acceptor materials. This article offers an in-depth review of the current developments in question.

Descriptions

Series Resistance in Organic Solar Cells-

It is important to note that a series barrier will have an opposite effect on the properties of the solar cells. The series blockage that was present in the dim characteristics has been eliminated, and the identical circuit model can be fused in two different ways; this does not affect the dull characteristics in any way..

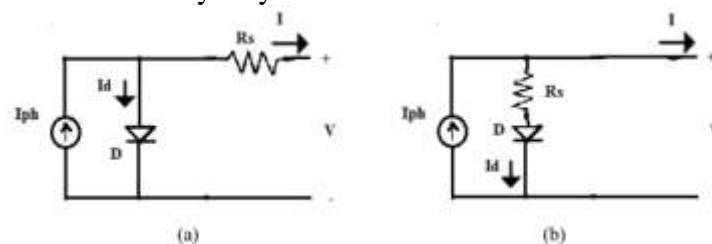


Fig.: solar cell equivalent model when (a) [(a)] series resistance is placed in series of both current source I_{ph} and the diode D (b) [(b)] series resistance is placed in series of diode D but in parallel of current source I_{ph}

In figure (a), the series obstruction is connected in series with both the current source I_{ph} and the diode D . On the other hand, in figure (b), the series obstruction is connected in series with the diode D but not with the current source I_{ph} . In the following part, boundary extraction is completed for the two models shown in figure by making use of the dark characteristics; after that, the recreated light characteristics are fitted back in by making use of the extricated borders..

Statement Of The Problem

Even though significant progress has been made in the field of organic solar cells over the past few decades thanks to a lot of hard effort, there are still certain phenomena that are not fully understood and there is opportunity for additional development. Some examples of such phenomena are as follows:

- Production of excitons
- Exciton diffusion
- Transport of carriers from one layer to the next Electron-hole separation caused by the buildup of the field Transport of carriers to the electrodes

Advantages of Organic Solar Cells over Inorganic Solar Cells

It has been demonstrated that organic solar cells (OSC), as opposed to their inorganic equivalents, offer three important benefits over their inorganic counterparts [15,16]. To begin, in terms of the materials, the limits of the silicon wafer technology result in the fabrication of materials that are ponderous, inflexible, brittle, and bulky. This is due to the fact that silicon wafers are made of silicon. In addition, the color possibilities that are available for these materials are somewhat restricted. On the other hand, the materials that are used for OSC are not only light in weight, but they are also thin, flexible, and changeable in color, and they can be molded into any pattern that you can dream of. This allows for OSC to be utilized in a wide variety of applications. Second, the process of synthesizing inorganic solar cells is not only laborious and time-consuming, but it also requires a significant amount of energy, mandates that the process be carried out at extremely high temperatures, and imposes limits on the largest possible size of the fabrications that are feasible.

All of these factors make it difficult to manufacture inorganic solar cells. On the other hand, the OSC technology is straightforward, easy to understand, does not call for a significant amount of energy, can be fabricated at room temperature, does not have any restrictions on the size of fabrication, and has high speed production in addition to numerous possibilities of printing and coating fabrication techniques (including inkjet printing, screen printing, and spray printing). In conclusion, roofs and open fields are the only viable uses for inorganic solar cells. The uses of organic solar cells (OSC), on the other hand, allow an almost infinite number of options. These applications include things like windows, displays, smart glasses, cars, and the charging of mobile devices and laptops. On rooftops and in open fields, inorganic solar cells are utilized to generate electricity..

Challenges for Organic Solar Cell Technology

Silicon is the material that is employed by the vast majority of businesses in the commercial solar industry, making it the material of choice in this sector. This is due to the fact that silicon has a high power conversion efficiency, which is why it is the material that the majority of businesses choose to work with. Inorganic solar cells have a number of drawbacks, the most prominent of which is the excessively high cost of their production, which is in addition to the manufacturing processes, which are difficult and demand a substantial amount of power. The rapid growth in demand for organic solar cells over the past several years is partially attributable to the ease with which these cells may be manufactured. To this point, the private sector has not demonstrated a significant level of interest in the OSC technology that is being developed.

This might be a result of the low power-conversion efficiency (PCE) of the technology as well as its short lifetime. The open-circuit voltage (OSC) is dependent on the heterojunction, which is generated whenever donor and acceptor materials are brought into close contact to one another. The OSC technology has a variety of hurdles, some of which include restrictions on the absorption efficiency of donor materials; blockages to the dispersion of excitons; difficulty in charge dissociation and transfer; and charge transfer. The length of the exciton diffusion, the structure of the crystal, and the collection of photogenerated carriers are all areas that have the potential to be improved by applying various approaches to handle these difficulties. These problems may be solved in a number of different ways. These advancements are made possible by the utilization of polymer-based nanocomposites, plasmonics, and tandem structures, in that order. In addition, hybrid nanocomposites, which

are made up of both organic and inorganic materials, have the potential to be a synergistic combination that makes the most of the one-of-a-kind characteristics of each of the distinct components. [25,26,27]. This is due to the fact that hybrid nanocomposites contain components that are both organic and inorganic in nature. When compared to traditional solar cells, hybrid solar cells operate in a manner that is a great deal more analogous to that of an OSC. High power conversion efficiencies (PCEs) may be attained by hybrid solar cells thanks to the presence of both n-type inorganic material and p-type semiconductor polymer within their composition. PCEs are a measurement of how efficiently a solar cell is able to convert the energy it receives from the sun into a form that can be utilized.

Materials for Organic Solar Cells

It is possible for the organic materials used in the manufacture of organic semiconductors to be derivatives of very small molecules or semiconducting polymers [28]. Organic semiconductors have been shown to contain conjugated organic moieties, which can take the form of dendrimers, pigments, dyes, polymers, and even very small molecules. There is a possibility that organic semiconductors will prove to be an effective replacement for OSC. In order for an OSC to operate effectively, there must be present both a kind of material known as an electron pair donor and a type of material known as an electron pair acceptor. In this procedure, the donor material is typically comprised of conjugated polymers. Poly(1,4-phenylenevinylene) (PPV) and its derivatives garnered a great deal of attention in the very early stages of research due to the fact that they exhibited appropriate optoelectronic features. Later on, polythiophene and its derivatives, notably poly(3-hexylthiophene), also known as P3HT, began to see broad application (as a donor material) due to the material's stability, high carrier mobility, and soluble nature. In particular, poly(3-hexylthiophene) began to see widespread application (as a donor material) because of its high carrier mobility. In reference to the OSC technology, the following is a list of the structures of some key donor materials, which may be found in Figure In spite of the fact that fullerene and its derivatives were the most essential components of OSC technology in the beginning very small molecules such as 3,9-bis(2-methylene-(3-(1,1-dicyanomethylene)-indanone) have subsequently gained the lead.. The three-dimensional architectures of many important fullerene and non-fullerene acceptor materials are depicted in Figure 1, which may be found here..

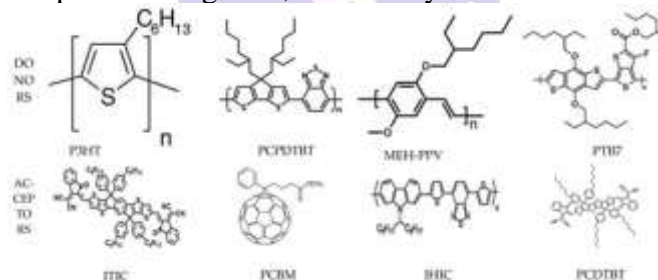


Figure: The structures of some important donor and acceptor materials of OSC.

It is feasible for the organic materials employed in the production of organic semiconductors to be derivatives of extremely tiny molecules or semiconducting polymers [28]. This is one of the ways that organic semiconductors may be created. It has been demonstrated that organic semiconductors contain conjugated organic moieties, which can exist in the form of dendrimers, pigments, dyes, polymers, and even very tiny molecules. These conjugated organic moieties can take any of these forms. There is a chance that organic semiconductors will show promise as a viable alternative to organic semiconductor components (OSC). In order for an OSC to function properly, it is necessary to have both a kind of substance known as an electron pair donor and a type of substance known as an electron pair acceptor in the circuit at the same time. In this method, the donor material will frequently be made up of conjugated polymers. In the very early phases of research, Poly(1,4-phenylenevinylene) (PPV) and its derivatives attracted a great deal of attention due to the fact that they demonstrated acceptable optoelectronic properties [29,30]. Later on, polythiophene and its derivatives, most notably poly(3-hexylthiophene), also known as P3HT, began to have widespread application (as a donor material) due to the material's stability, high carrier mobility, and soluble nature [31,32,33]. In particular, poly(3-hexylthiophene) is the

derivative that has received the most attention. Due to the great carrier mobility that it possessed, poly(3-hexylthiophene) in particular started to see extensive application (as a substance for donor cells). In relation to the OSC technology, the following is a list of the structures of a selection of important donor materials; these structures may be found in Figure In the beginning [34,35,36,37,38], fullerene and its derivatives were the most fundamental components of OSC technology. However, extremely tiny molecules such as 3,9-bis(2-methylene-(3-(1,1-dicyanomethylene)-indanone) have since taken the lead. Figure 1, which may be seen at this link, provides a visual representation of the three-dimensional structures of a number of significant fullerenes and non-fullerene acceptor materials..

Fabrication of Organic Solar Cells

Single-layer organic photovoltaics (OPVs) are just what they sound like: organic photovoltaics that only have one active layer. On the other hand, organic photovoltaics that are built with many active layers are referred to as hetero-junctions or multiple layers [88]. Organic photovoltaics that only include a single active layer are referred to as single-layer OPVs. This is also a common abbreviation for the term "single-layer organic photovoltaics." Due to the fact that they only have one layer, OPVs with only one layer do not have particularly outstanding performance in terms of PCE. When OPVs are built with a high number of layers or heterojunctions, there is a rise in the ON-State Current (OSC), which may be measured. This phenomenon can be observed. It is possible to see this phenomena. The operation of a heterojunction organic photovoltaic device can be broken down into four distinct steps, which are as follows: (i) the generation of an electron-hole pair (exciton) by the absorption of the photon; (ii) the diffusion of the exciton to the interface; (iii) the generation of free carriers by the dissociation of excitons; and (iv) carrier transport and collection at the electrode. Each of these steps is described in more detail below. The following provides a more in-depth explanation of each of these stages. The details of each of these steps are broken down into the following sections for your convenience. The first generation of organic semiconductor chips, known as OSCs, consisted of a single layer that had an extremely low power carrier efficiency (PCE). This was a direct result of the inefficiency of the dissociation exciton, which caused the layer to have such a low efficiency. This was due to the fact that organic semiconductors are single-layer devices, which is one of the characteristics of these materials.. (Figure 2A).

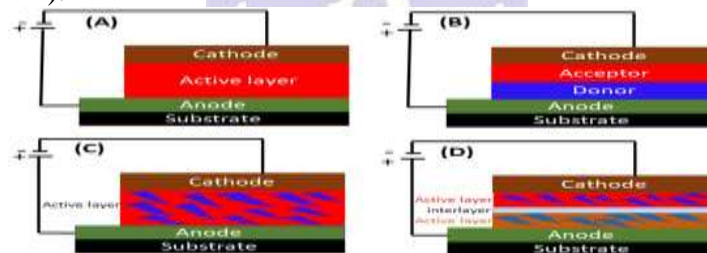


Figure The following stages are involved in the production of an OSC: (A) the single active layer; (B) the bilayer heterojunction; (C) the bulk heterojunction; and (D) the tandem heterojunction.

Further advancements in OSC technology led to the development of heterojunction cells as the final product. It is possible to further subdivide these cells into bilayer heterojunctions, bulk heterojunctions, and tandem heterojunctions. In a configuration known as a bilayer or planar heterojunction (Figure 2B), layers of donor and acceptor materials are positioned between two electrodes. A low PCE is also an issue for planar heterojunctions, which is caused by their short diffusion length and the restricted mobility of the excitons that they form. This difficulty is created by the fact that planar heterojunctions are very simple structures. In order to achieve an optimal power conversion efficiency (PCE), it is essential to synchronize the diffusion length with the dimensions of the bilayer heterojunction. Because of this, the final product will be films that are extremely thin and have a low optical absorption.

On the other hand, heterojunctions can be either distributed or bulk, and in either case, they are composed of combinations of materials that can act as either donors or acceptors (Figure

2C). Excitons have an easier time diffusing to the interface and achieving high power carrier efficiency when bulk heterojunctions are used, as opposed to planar heterojunctions. PCBM is the combination of donor and acceptor materials that is used in bulk heterojunctions more frequently than any other [89,90]. P3HT is an abbreviation for polycrystalline bulk metal-organic framework. Up to an 18 percent increase in power conversion efficiency (PCE) has been shown in bulk heterojunctions as a result of recent research [91,92,93]. In single-layer, bilayer, and bulk heterojunction structures, the capacity to absorb light is limited to a narrow band of the visible light spectrum. Depending on the structure, this band is either in the short- or the long-wavelength region. When using a tandem heterojunction, one has the possibility to make use of a large wavelength region that is based on two sub-cells whose absorption spectra are complimentary [94,95]. This gives an excellent chance to maximize one's profit potential. Interlayers divide the sub-cells from one another and are responsible for aligning the quasi-Fermi level of both cells (Figure 2D). The sub-cells are separated from one another by the interlayers. As a result of the many benefits that were just gone over, tandem solar cells have lately attracted a lot of interest in recent times. An obstacle for tandem inorganic solar cells is the enormous expenditure that must be paid in order to accomplish the requisite crystal growth and lattice matching. This expense is a significant barrier. On the other hand, tandem OSC is experiencing an increase in both attention and application due to the fact that its manufacture is simple and cost-effective. This is one of the reasons why. For bulk heterojunctions, the input materials have to have absorbance spectra that are a perfect match for one another, correct energy level alignment, nanoscale phase separation, and high charge mobility.

The Working Principles Behind Organic Solar Cells

The operation of organic semiconductor sun cells (OSC) is completely different from that of conventional silicon-based inorganic solar cells, both in terms of the material properties of the OSC and the design of the OSC. Organic semiconductor sun cells, in contrast to their normal inorganic counterparts, do not immediately result in the production of free charge carriers upon the absorption of light. This is because organic semiconductor solar cells include a double layer structure. Excitons are pairs of electrons and holes that have no net charge whatsoever and are created as a byproduct of this process. It is a pair of electrons and holes that is extremely tightly linked and has a life span on the order of nanoseconds. The excitonic property of the organic semiconductor material has a considerable impact on both the design and the geometry of organic photovoltaic (OPV) devices. To be able to deal with the limitations and challenges that are presented by the materials and the production process, it is very necessary to have a fundamental understanding of how OPV works. The exciton must first dissociate into its component components in order for the photovoltaic effect to be created. This is a prerequisite for the process. Excitons have a binding energy that ranges anywhere between 0.3 and 1.0 eV. In addition to the energy required to bind the excitons together, an evident amount of more energy is needed in order to dissociate the excitons and free the carrier. This shows a requirement for a large quantity of energy as compared to its inorganic equivalent, which means that it is necessary. Excitons are created as a direct consequence of the acceptor absorbing part of the light that is passing through it. If the amount of thermal energy generated is less than the amount of energy required to bind an exciton, the electron and hole states will ultimately unite into a single state. On the other hand, if the thermal energy is greater, it will surpass the exciton binding energy if it is compared to that value. Because of this, free charges will be generated, which will ultimately result in the electron and hole becoming structurally distinct from one another. Because of this, photogenerated carriers begin to accumulate on the electrodes, with electrons beginning to gather at the cathode and holes beginning to gather at the anode. The operational idea that underpins OSC is illustrated in Figure 3, which may be found here. The act of absorbing and processing light from the sun results in the development of an electron-hole pair, which is also referred to as an exciton. Because of the high energy absorption, the donor's valence electrons jump from their highest occupied molecular orbital (HOMO) level to their least unoccupied molecular orbital (LUMO) level. This causes the excitons to split into electrons

and holes at the HOMO level. HOMO and LUMO are both abbreviations for molecular orbital levels. After that, electrons hop from the lower unoccupied shell (LUMO) of the donor to the lower unoccupied shell (LUMO) of the acceptor, which has a lower energy level than the donor's LUMO. When the holes from the HOMO of the donor are moved to the anode, the circuit is complete. The circuit is completed when the electrons from the LUMO of the acceptor are transferred to the cathode, where they are collected. In this context, some of the most important considerations to take into account are the alignment of the energy levels of the donor materials and the acceptor materials, the absorption profiles, the mobility of charges, and the miscibility of the materials.

CONCLUSION

Solar cells are what you name the devices that do the work of turning light into energy. This theory is based on the organic solar cell because of the many benefits it provides, such as low material and energy requirements, low temperature and energy handling requirements, low material necessity, the ability to be used on a flexible substrate, and the capability to be molded to suit a compositional purpose. These benefits serve as the foundation for this theory. Both the light absorption and the exciton dispersion processes fall within the category of optical processes, whereas the remaining phases are driven by electrical forces. Light retention, exciton dissemination, charge movement, charge transport, and charge assortment are the main phases that are involved in the creation of electricity from light in an organic solar cell. These phases are essential to the process. The four most essential types of architectures for organic solar cells are single-layer, double-layer, mixed, and coated organic solar cells. This theory is based exclusively on the MATLAB-based 1D float dispersion electrical display of a bilayer organic solar cell, and it is relied on it alone. MATLAB, a high-performance programming language with particular registration capabilities, is used to carry out the simulation. Calculation, disclosure, and programming are all part of this language's feature set, and it operates in an environment that is free of demanding scenarios. A large amount of numerical documentation is used to present both the problems and the answers. The bilayer organic semiconductor capacitor (OSC) employs two extremely thin layers (100 nm each) of organic material (a contributor and an acceptor) in the space between the cathodes. Each layer is a contributor to the OSC.

1. This test system uses MATLAB programming to successfully work through each and every step involved in the translation of light into electricity by bilayer organic solar cells. This testing device performs exceptionally well for a modest range of voltages, which is typical for areas with solar activity.
2. It has been discovered that the open-circuit voltage is dependent not only on the difference in the cathode work works but also on the force and the charge densities at the terminals. An increase in open-circuit voltage can be achieved either by increasing the infusion hindrance in the polymer or by reducing the charge densities at the anodes. In an open circuit, the voltage rises in a logarithmic fashion with increasing light force.

REFERENCES

- [1] H.Hoppe, N.S.Sariciftci. "Organic solar cells: An overview", J. Mater. Res, (2004) <https://doi.org/10.1557/JMR.2004.0252>
- [2] B. A. Gregg and M. C. Hanna. "Comparing organic to inorganic photovoltaic cells: Theory, experiment and simulation", J. Appl. Phys., 93:3605, (2002). <https://doi.org/10.1063/1.1544413>
- [3] J. Nelson. "The Physics of Solar Cells", Imperial College Press, London, (2003). <https://doi.org/10.1142/p276>
- [4] C. Shuttle, B.O'Regan, A.Ballantyne, J.Nelson, D.Bradley, J.Durrant, "Bimolecular recombination losses in poly thiophene:fullerene solar cells", Phys.Rev.B:Condens.Matter 78, 113201, (2008). <https://doi.org/10.1103/PhysRevB.78.113201>
- [5] J. Guo, H.Ohkita, "Charge generation and recombination dynamics in poly(3-hexylthiophene)/fullerene blend films with different region regularities and morphologies", Adv.Mater., 132, 6154–6164,(2010).

- [6] M. Lenes, M.Morana, C.J.Brabec, P.W.M.Blom, "Recombination-limited photo currents in low bandgap polymer/fullerene solar cells", *Adv. Funct. Mater.* 19 1106–1111, (2009). <https://doi.org/10.1002/adfm.200801514>
- [7] K.J. Li, L.J. Li, J.C. Campbell, "Recombination lifetime of free polarons in polymer/fullerene bulk heterojunction solar cells", *J. Appl. Phys.* 111, 034503 (2012). <https://doi.org/10.1063/1.3680879>
- [8] Jain, A., and Kapoor, A., "A new approach to study organic solar cell using Lambert W-function", *Sol. Energy Mater. Sol. Cells*, 86, 197–205 (2005). <https://doi.org/10.1016/j.solmat.2004.07.004> Mazhari, B., "An improved solar cell circuit model for organic solar cells", *Sol. Energy Mater. Sol. Cells*, 90,1021–1033 (2006). <https://doi.org/10.1016/j.solmat.2005.05.017>
- [9] P. K. Watkins, A. B. Walker, and G. L. B. Verschoor, "Dynamical Monte Carlo modelling of organic solar cells: The dependence of internal quantum efficiency on morphology," *Nano Lett.* 5, 1814–1818 (2005). <https://doi.org/10.1021/nl051098o>

