



Review

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High-performance and multifunctional organic photovoltaic devices

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Abstract: Organic photovoltaic devices (OPVs) are emerging as a promising renewable energy source for the future. Their unique advantages, such as semitransparency, light weight, superior flexibility, and low cost, enable a wide range of applications. However, compared to silicon-based photovoltaics, OPVs still face challenges for further improving their efficiency. Additionally, there is a need to explore their potential of multi-functionality for practical application in various scenarios. This review summarizes the recent achievements in optimizing device performance and enhancing power-conversion efficiency, particularly via tuning the intermolecular interaction to reduce the electron-vibration coupling and non-radiative charge recombination (denoted as the “dilution effect”). Moreover, the representative development of ultra-thin Ag transparent electrode-based OPVs with multi-functional capabilities (such as semitransparency, flexibility, stretchability, and better aesthetics) have also been covered. Therefore, this review aims to provide a broad landscape on the recent development of OPV and to unlock the full potential of OPVs.

Key words: Organic photovoltaic; Multi-component; Semitransparent; Flexibility

1 Introduction

Reducing global energy consumption from fossil fuels requires innovative and cost-effective renewable energy technologies. Photovoltaics (PVs) can fulfil this need amply if deployed over a large enough area (Maka and Alabid, 2022). As a result, terrestrial PV technology has emerged as the most rapidly growing electrical energy source in the past decade, and now accounts for ~ 6% of the total electrical supply in China (Victoria et al., 2021; Zhang et al., 2023). No doubt, this will continue soaring. Installing more PV panels in remote and sunny regions could theoretically meet the growing demand for energy, but implementation is restricted by several practical factors (Zhang et al., 2023; Zhang et al., 2023; Zhong et al., 2022). First, the area suitable for PV plants is

limited in most energy-demanding countries. Second, the maintenance and distribution costs, as well as adverse environmental effects, make numerous plants less practical. Third, the traditional structure of energy conveyance and consumption requires averting the direct current into alternating current, transportation along grids, and conversion to direct current for most household appliances. This process sacrifices a considerable proportion of the energy produced. In addition, the energy generated by PV panels needs to be conserved in the daytime and released at night. These drawbacks make terrestrial PV-based electrical systems less efficient.

One approach to tackling these issues is through developing PV technology that can be installed near homes for instant use. This can be achieved by integrating PV panels with surfaces used in daily life, e.g. vehicles, buildings, clothing, and wearable electronics, in ways that will not greatly impact function or aesthetics. Applications in outer space are also a possibility. Organic photovoltaic devices (OPVs) based on highly conjugated conductive polymers exhibit inherent stability against high radiation doses, and the excellent photovoltaic performance retention of OPVs when operating at

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high-altitude platform stations makes them promising for applications in outer space (Kumar et al., 2010; Datt et al., 2022). To further advance the sustainability of photovoltaic technology, ongoing research on and enhancements in device design and performance are essential. Future endeavors in photovoltaic technology will strive to enhance functionality, including features like semi-transparency, flexibility, and aesthetics, to seamlessly integrate solar power into our everyday lives (Liu et al., 2022; Huang et al., 2023; Tak et al., 2017; Traverse et al., 2017). In this context, organic semiconductor-based PVs have a unique advantage, due to their intrinsic optoelectronic and processing properties. For example, the organic semiconductor has a distinct absorption profile, which makes it a rare but valuable candidate for achieving selective absorption for semitransparent PV application. The organic semiconductor is also famous for its flexibility and stretchability in device applications, which further widens its applicability in curved structures and dynamically flexible and stretchable applications. Furthermore, organic semiconductors can be manufactured in vivid colors for aesthetic applications.

Currently, the relatively low device efficiency they offer is still the major obstacle for the practical application of OPVs. Their intrinsic limitations include high exciton-binding energy, short exciton-diffusion distance, and severe non-radiative charge recombination. (Veldman et al., 2008; Yuan et al., 2011). The innovation of D:A heterojunction structure and bulk-heterojunction structure successfully compensates for the high exciton-binding energy and short exciton-diffusion length, and enables efficient charge generation. As a result, high photon-to-electron conversion efficiency was achieved by Park et al., with internal quantum efficiency around 100% (Park et al., 2009). As shown in Table 1, it is clear that specific device parameters, i.e. the external quantum efficiency (EQE) and the fill factor (FF), are quite similar to those of inorganic counterparts. However the open-circuit voltage (V_{OC}) loss is still large, which has become the most critical challenge in device efficiency. It has been generally recognized that this severe V_{OC} loss is caused by non-radiative charge recombination (Vandewal et al., 2009). As a matter of fact, the non-radiative decay

process occurs throughout the whole photon-to-electron process: (1) the exciton generated upon light incidence is quenched via thermal relaxation. (2) The subsequent exciton separation results in a low-energy charge-transfer state (CT), which is regarded as the dominant species for non-radiative charge recombination (according to the energy-gap law). (3) Following CT separation, polarons will form and ultimately be collected by an electrode to drive the out-circuit appliance. Meanwhile, a charge back-transfer process will occur and cause triplet states, which will be quenched to ground state via the non-radiative mode. Vandewal et al. (2017) systematically studied the electron-vibration coupling in fullerene-based OPVs, and pessimistically concluded that the intrinsic electron-vibration coupling from the backbone of the active layers inevitably induces large non-radiative charge recombination and subsequent large V_{OC} loss (Benduhn et al., 2017). This sets an upper efficiency limit of 25.5% for OPVs. By comparison, the efficiency limit for a normal PV technique is over 33% (according to the Shockley-Queisser limit) when no non-radiative charge recombination occurs. Moreover, electron-vibration coupling is also significantly involved in the CT state's induced non-radiative charge recombination. Most high-performance OPVs are currently fabricated on glass/indium tin oxide (ITO) substrates with an opaque metal film as the back electrode. This structure offers a good compromise between transparency and conductance. However, it also results in rigid, expensive, fragile, non-transparent, and heavy devices. Consequently, it diminishes many of the usual advantages of OPVs, such as intrinsic flexibility, vibrant color, light weight, and cost-effectiveness (Zhao et al., 2022a; Zheng et al., 2022a). Developing more advanced device structures and non-active materials like interfacial layers and electrodes is necessary to fully explore the potential of OPVs. For instance, achieving superior flexible or stretchable OPVs requires the development of appropriate device structures, transparent electrodes, back electrodes, substrates, and interfacial layers. One notable feature of organic semiconductors for multifunctional photovoltaic applications is their selective absorption profiles. They strongly absorb near-infrared light, while allowing penetration of

most visible photons. This characteristic sufficiently enables integration with windows, allowing simultaneous electricity generation without compromising window functionality (Tak et al., 2017; Wang et al., 2021). Additionally, it facilitates heat control of the building by managing middle-infrared photons (Amirkhani et al., 2019). However, high-performance semi-transparent OPVs require management of optical properties throughout the entire device to maximize selective absorption.

In this review, we summarize the recent work to address electron-vibration-induced non-radiative charge-recombination loss via the dilution effect (or

the phenomenon of intermolecular influence on the electron-vibration coupling) for high device performance, as well as the new concepts and device structures targeted to various applications of OPVs. These understandings and strategies essentially enable the progress high-performance OPVs with certified efficiency approaching 20%, which is currently the best result of single-junction OPVs. These new device structures, including semitransparent and flexible/stretchable structures, also represent the best compromise between device performance and functionality.

Table.1 Device parameters of different single-junction PVs (Zuo et al., 2022)

	GaAs	Si	CdTe	CIGS	OSCs
E_g (eV)	1.43	1.11	1.44	1.12	1.52
V_{OC} (eV)	1.10	0.75	1.01	0.74	0.88
FF (%)	> 85%	~ 85%	> 80%	> 80%	< 80%
V_{OC} loss (eV)	0.33	0.36	0.43	0.38	0.56
EQE (Avg.) (%)	> 95	> 95	> 90	> 90	< 80

2 Effects of electron-vibration coupling

Electron-vibration coupling is regarded as one of the most important channels for non-radiative charge recombination, as reported in our previous work (Zuo et al., 2022). Since this coupling originates from the conjugated backbones of an organic semiconductor, the resultant V_{OC} loss appears to be unavoidable and severe. However, it should be noted that electron-vibration coupling is also intimately linked to the intermolecular interactions/environment, where two major effects, i.e., local polarization and molecular packing, are involved.

2.1 Polarization effect

The excited state of a molecule typically exhibits a larger dipole moment compared to the ground state. The coulombic interactions that occur in excited states cause dramatic change in the surroundings. Therefore, the local surrounding polarizability also becomes involved; the change in the surrounding chemical environment contributes to electron-vibration couplings. A prominent phenomenon in this mechanism is the solvation effect, in which the solute molecules dissolved in solvents with different dipole moments tend to exhibit different optical spectra (Bulović et al., 1998, 1999). It is generally

observed that the absorption and photoluminescence (PL) spectra exhibit an obvious red shift with higher solvent polarity, while the PL quantum efficiency (PLQY) tends to decrease. A similar effect has been observed in the solid state, and is termed the “solid-state-solvation” effect. It is generally adopted as an effective method to tune the excitonic and optical properties of organic semiconductors in thin films (Northey et al., 2017). Usually, dispersing the emitter or dopant in a continuous matrix of a semiconducting material with a larger band gap significantly suppresses non-radiative charge recombination and induces higher luminescence. As a result, the solid-state-solvation effect has been widely explored in organic light-emitting devices (OLEDs), and demonstrated to be an important strategy to achieve high performance. Although the working mechanism for the suppressed non-radiative charge recombination underlying the solid-state-solvation effect is complicated for OLED applications, the change in surrounding polarizability certainly plays a role. This role becomes even more important for OPVs, because the dipole moment for OPV materials is even larger. To absorb more photons in the solar spectra, OPV materials must have much a lower band gap (optimal $E_g \sim 1.30$ eV from the S-Q limit) than OLED materials (visible region $\sim 1.77 - 3.1$ eV). To achieve a lower

band gap, one of the most frequently used strategies is the donor-acceptor alternating strategy, and the internal charge transfer is closely linked to band-gap size. The stronger the intramolecular charge transfer (ICT), the smaller the band gap (Lim et al., 2021; Scharber and Sariciftci, 2021). Therefore, the classic OPV materials typically exhibit larger conjugation and stronger polarity. This affects the properties of the excited states as well as electron-phonon coupling. Because the photon absorption of organic molecules induces a excited state of larger dipole moment, and this will further induces the change of the surrounding molecules. This also relaxes the energy of the excited state, corresponding to the coordinate shift in the overall system.

2.2 Molecular packing effect

Vibration of chemical bonds is the most important factor in electron-vibration coupling, particularly for bonds that directly contribute to the molecular frontier orbitals. Bond vibration can also be directly affected by molecular packing. For example, in solution, free rotation of the phenyl ring of tetraphenylethylene (TPE) occurs, while this motion is suppressed in a solid state, contributing to the famous aggregation-induced emission (AIE) (Hong et al., 2009, 2011). Furthermore, the different molecular aggregations, e.g. the H or J aggregates, can also affect electron-vibration coupling dramatically. Amorphous and crystalline packing structures affect electron-vibration coupling as well. These findings indicate that it is possible to take advantage of the intermolecular effect to suppress electron-vibration coupling and the corresponding non-radiative charge recombination.

3 Dilution effect for multi-component OPVs

The most popular device structure for OPVs is the bulk-heterojunction (BHJ), which contains a nanoscale phase-separated continuous interpenetrating network of donor and acceptor binary blend (Huang et al., 2014). This structure successfully compensates for the high exciton-binding energy and poor ambipolar charge transport of OPV materials, and thus, enables high photocurrent generation and collection efficiency, corresponding to a high external quantum efficiency of over 90% and a high fill

factor (FF) of over 0.8. In order to tailor the intermolecular interaction for electron-vibration coupling and non-radiative charge-recombination suppression, the most intuitive method is to incorporate a third component to form a multi-component blend, e.g. the ternary structure. In fact, the ternary has long been proposed for different purposes, such as achieving complementary absorption, optimizing morphology, and facilitating charge transfer. Different working mechanisms of the ternary blend have also been proposed, e.g. the charge-transfer model, energy-transfer model, parallel model, and alloy model (Günther et al., 2023; Wang et al., 2023). Further, the uncertainty in terms of fundamental understanding of multi-component OPVs has hindered the development of a unified material selection rule to universally achieve higher device performance. In most ternary OPVs, a lower non-radiative charge recombination and blue-shift in absorption spectra have been generally observed, while the fundamental mechanism underlying these phenomena is less explored.

3.1 The proof-to-concept study

We took advantage of the ternary strategy for a novel approach to tailoring the electron-vibration coupling in OPVs via controlling the intermolecular interaction, and propose using the dilution effect for multi-component OPVs. By blending two non-fullerene acceptors (NFA), specifically ITIC and 4TIC, we found the two molecules formed a good mixture at the molecular level, as shown in Fig. 1 (Zuo et al., 2022). Interestingly, the blend film exhibited a steeper absorption tail, indicating the formation of “ordered” intermixing. Moreover, we identified that the photon-emission center was still localized on the low-band gap 4TIC, and the Forster energy transfer from ITIC and 4TIC occurred in the blend. However, ITIC provided a lower local polarizability surrounding 4TIC and interrupted the molecular packing of 4TIC. As a result, the emission properties of the blend were quite different from those of pristine 4TIC; the PL spectra exhibited a blue shift and PLQY increased. We also numerically analyzed the electron-phonon coupling, and calculated the Huang-Rhys parameter (S_H) for 4TIC in pristine and blend films, by decoupling the Barry-center and Stokes shift from the absorption and PL

spectra. We found that the S_H of 4TIC in the blend was reduced, and this unambiguously verified that electron-vibration coupling was suppressed via the intermolecular interactions. Further, from the analytical model based on the Marcus-transition theory and Frank-Condon equations, we identified that the degree of S_H suppression exactly accounted for the four-fold increase in PLQY (Jong et al., 2015). We then analyzed the electron-vibration coupling in the device via the EQE and electroluminescence (EL) spectra, and found that the properties of the blend film were passed on to the device, contributing to the increase in ELQE.

For an OPV device, charge generation and transport are also very important. Our comprehensive analysis showed that the dilution effect does not negatively affect the charge-generation process, but the incorporation of ITIC becomes a bridge for electron transfer from the PM6 donor to the 4TIC acceptor. Interestingly, we observed that the charge could spontaneously jump from 4TIC to ITIC, which is supposed to be a thermally-activated process (Liu et al., 2018; Shi et al., 2017). We identified this mechanism using several NFA pairs of different

LUMO offsets, and found the intensity of the higher LUMO NFAs to be in inverse proportion to LUMO offsets. This significantly affected the charge-transport process, because the microscopic mechanism is the hopping of the charges within the adjacent molecules. The LUMO offsets appear to affect composition-dependent carrier mobility.

With this understanding, we further outlined the material-selection rules for high-performance dilution-effect multi-component OPVs, which include: (1) good miscibility to facilitate molecular intermixing and enable the dilution effect to occur, (2) a larger band gap to ensure that the emission center is not changed and to form a lower local polarizability, (3) slightly higher LUMO levels to guarantee stability of the D:A charge-transfer center and efficient charge transport, and (4) lower local polarizability and/or formation of specific intermolecular packing structures to suppress the intramolecular bond vibrations that contribute to the molecular frontier orbitals. Then, we confirmed the generalizability of the dilution effect for enhancing the device performance of OPVs for a large range of NFA systems.

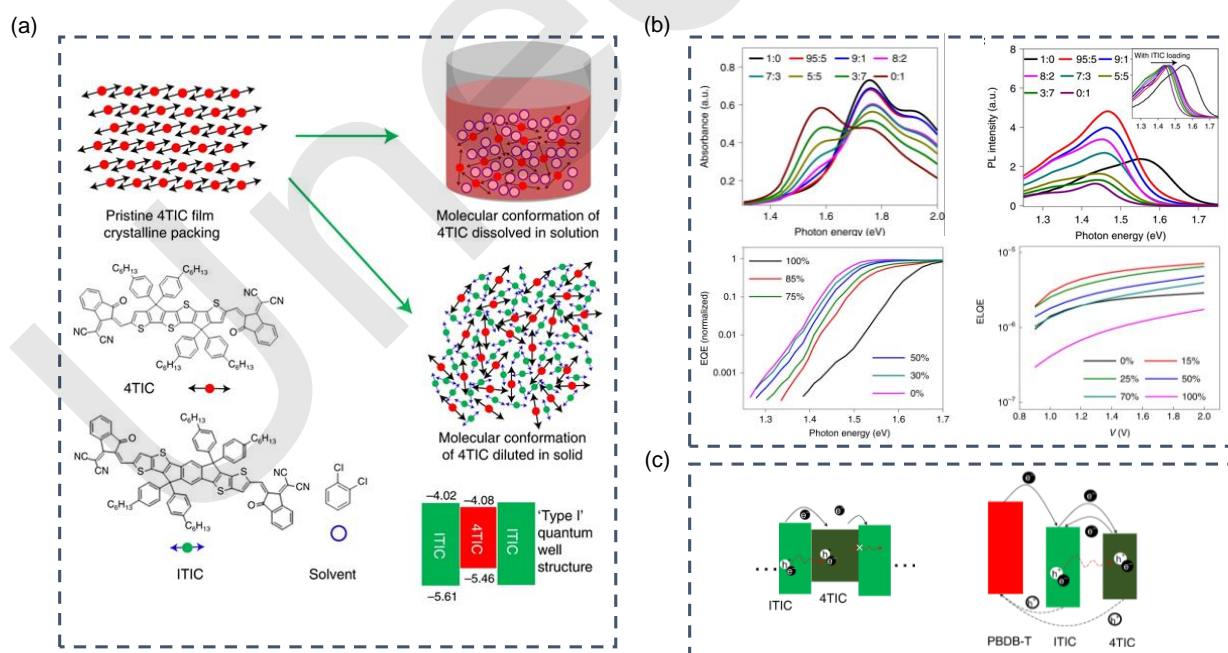


Fig. 1 (a) Schematic diagram of dilution effect, energy-level diagrams (unit, eV), molecular structures of ITIC and 4TIC, dilution with solvent or solid. (b) Absorption spectra, PL spectra, EQE spectra, and ELQE curves of ITIC-4TIC at different weight ratios. (c) Schematic diagrams for the charge and energy-transfer process between ITIC and 4TIC, and for charge and energy transfer in ternary blends (Zuo et al., 2022)

3.2 Effect of molecular packing

Although reduced electron-vibration coupling suppression has been confirmed in dilution-effect multi-component OPVs, it is still necessary to explore the microscopic mechanism in order to develop appropriate materials that would minimize non-radiative charge recombination and enhance device efficiency. It is clear that the use of a larger band gap tends to form a low polarization environment to reduce electron-vibration coupling, but it remains unclear whether the molecular packing effect also matters. Indeed, adequate characterization of the molecular packing is quite challenging, because the active layer of OPVs is amorphous and understanding of electron vibration requires very detailed molecular packing information. Therefore, we used a molecular dynamic (MD) simulation to study the molecular packing in detail (Li et al., 2022). Based on the MD results, we verified the formation of molecular intermixing between the two NFAs; moreover, we saw that the end-group cyano indanone (IC) packing became more ordered in the blend film. An enhancement in molecular packing after blending in a third component has also been observed in other studies (Chen et al., 2022). Considering that the IC and thieno[3,2-b]thiophene (TT) connection is the most rotatable chemical bond, and contributes significantly to the frontier molecular orbitals, enhancement of IC packing suppresses IC-TT-bond vibration. As a result, there is significant enhancement in PL intensity after blending BTP-eC9 with BTP-S2, and the ELQE of the dilution-effect MC-OPV increases dramatically (Li et al., 2022).

To further confirm the molecular packing effect on electron-vibration coupling in the NFA-based OPVs, we proposed a design with mutual dilution, which involved selecting two NFAs with similar band gaps, i.e. BTP-eC9 and BTP-S9, as shown in Fig. 2a (Zhan et al., 2022). The two molecules had

almost identical optical properties and energetic structures, as well as polarizability. However, due to the use of extended conjugation at the end group, BTP-S9 exhibited better molecular packing behavior. Blending the two molecules together essentially excluded the effect of local polarizability on electron-vibration coupling, and allowed us to examine the molecular packing effect independently. We observed that the ternary OPV also exhibited improved ELQE and a lower V_{OC} loss. Significantly, the V_{OC} of the ternary OPV exhibited a convex composition-dependent behavior. This actually is hard to interpret with any existing working mechanism of ternary blends, but can be easily understood in terms of the dilution effect and molecular-packing-induced electron-vibration coupling suppression. Moreover, in the framework of the dilution effect, the variations in band gap and non-radiative charge recombination explain the composition-dependent V_{OC} behavior well for most of ternary blends, as demonstrated in our previous work. In addition, the blend film exhibited no blue shift in the absorption or PL spectra. On the contrary, the EQE response edge became sharper, suggesting that the blue shift in the blend mostly originated from the surrounding polarization effect (Liu et al., 2020).

Previous work has shown that substitution of a bromo group for a fluoro group can enhance intermolecular interaction. Inspired by this, we designed the BTP-H2 shown in Fig. 2b, in order to minimize electron-vibration coupling (He et al., 2022). We identified the strong molecular packing of BTP-H2 via the GIWAXS method, and this significantly induced spontaneous generation of delocalized excited species. A similar effect was observed when blending BTP-H2 with BTP-eC9. Due to the reduced non-radiative charge recombination, the V_{OC} loss of the ternary blend was significantly reduced, and the best efficiency reached was 19.4%.

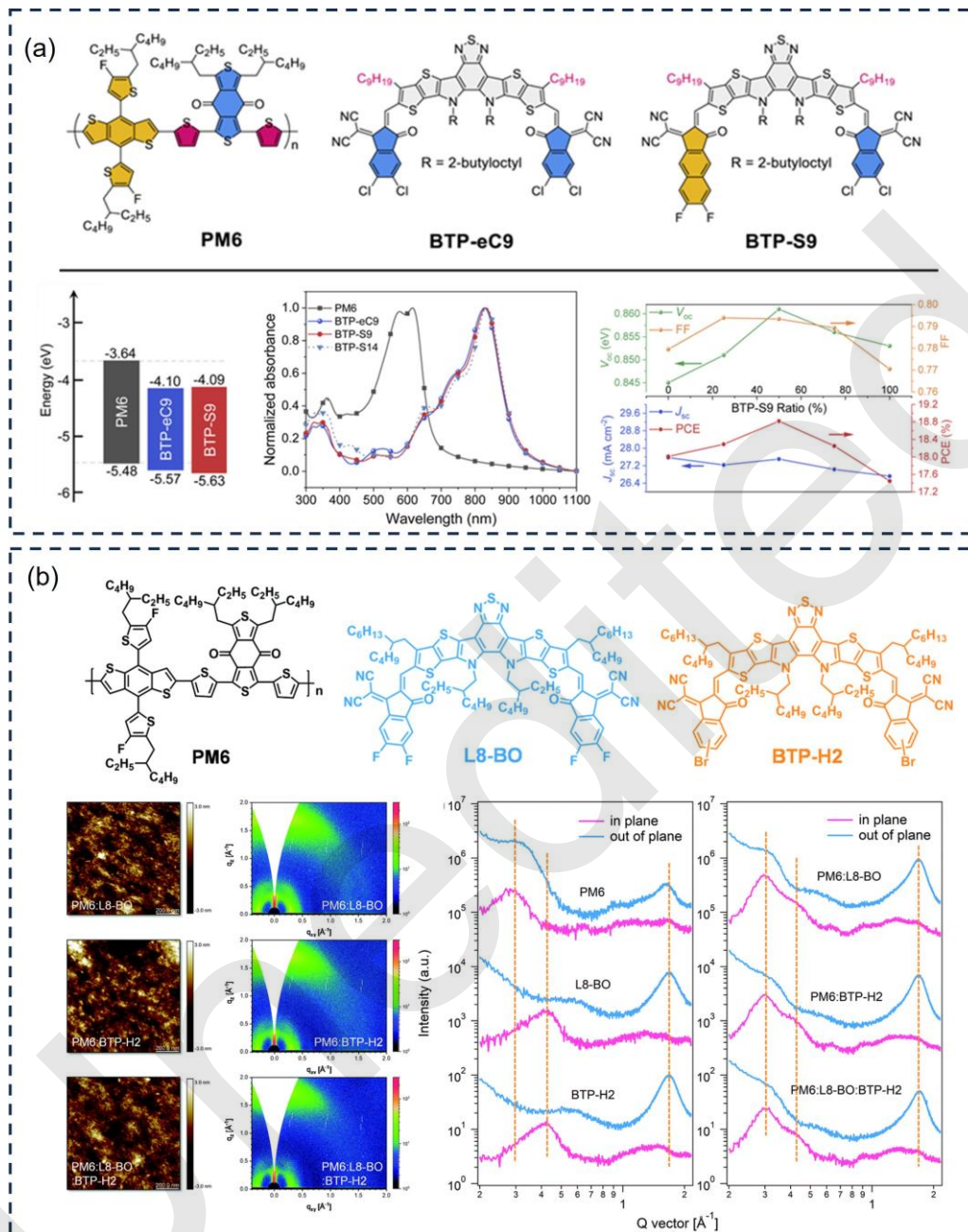


Fig. 2 (a) The chemical structures and properties of PM6, BTP-eC9, and BTP-S9, and the composition dependence of V_{OC} uplift in ternary devices (Zhan et al., 2022). (b) The chemical structures of PM6, L8-BO, and BTP-H2, and the results of GIWAXS testing (He et al., 2022)

3.3 Extended strategies

In dilution-effect MC-OPVs, the shrinkage of absorption spectra will cause a decrease in photo-current generation, primarily due to the surrounding polarization effect. To better balance photo-current generation and non-radiative V_{OC} loss, we propose a mixed diluent strategy, which involves one

large-band-gap diluent to tune the local polarizability and one low-band-gap diluent to further increase intermolecular interactions, specifically IC-IC packing. We selected the BTP-S16 and BTP-S17 as the low-band gap and large-band gap NFAs, respectively, to “dilute” the BTP-eC9 in the PM6: BTP-eC9 blend (Chen et al., 2023). With this design, a better compromise between charge generation and charge re-

combination was achieved, which was validated by the high PCE achieved (up to 19.4%), as certified by the National PV Industry Measurement and Testing Center (NPVM). To the best of our knowledge, this result represents a record for certified single-junction OPVs.

Unlike the charge-recombination center in a pristine acceptor blend, that in an OPV is in the CT state, which involves a coulombically bounded hole-electron pair on both the donor and acceptor. As a result, the electron-vibration coupling in the OPV is from both the donor and the acceptor. Therefore, there is still plenty room to suppress the non-radiative charge recombination via diluting the donor component.

4 Multi-functional OPV devices

The most prominent advantage of OPVs lies in their multi-functionality: the good semi-transparency, superior flexibility/stretchability, visual aesthetics, and other features that stem from the nature of organic semiconductors. Delicate design of the device structure and fundamental understanding of the optical and electrical processes are required to maximize overall performance.

4.1 Semitransparent OPVs

Solar energy consists of photons with different energies or wavelengths, and can be classified into the ultra-violet region, visible region, and infrared region. It can be used for different purposes such as lighting, electricity generation, and temperature control (Singh et al., 2020). The semitransparent (ST) OPV was developed to comprehensively deliver the above functionalities, which can be widely applied in buildings or vehicles. It is clear that ST-OPVs should selectively absorb in the invisible ultraviolet and near infrared regions, transmit the visible photons, and reflect the middle/far infrared photons. However, the absorption coefficient of inorganic semiconductors increases monotonously with the photon energy, and this means its visible absorption will be much stronger if it absorbs in the infrared region (Lee et al., 2020). The absorption spectrum of organic semiconductors features tunable absorption peaks and

valleys. It makes OPVs unique in their capacity for selective absorption, and especially suitable for ST-OPV applications (Chang et al., 2021; Liu et al., 2022; Xu et al., 2021; Zhang et al., 2022).

For ST-PVs, it is clear that selective absorption is a key factor; it is a qualitative concept that offers intuitive understanding of and guidance for material and device structural design. In a recent paper, we proposed the numerical definition of the absorption selectivity (S), by simultaneously taking the invisible absorption and visible transmission into consideration (Li et al., 2021). This numerical definition provides quantitative understanding and guidance for ST-OPV. This parameter was found to directly correlate to light-utilization efficiency, which is a comprehensive parameter to evaluate the overall performance of ST-OPVs. This essentially guides us in materials selection from a large pool of donor and acceptor materials, as well as in the optimization of device structures. For example, by analyzing the S values, we identified a combination of polymer donor PTB7-Th and NFA H3 as the most promising active layer, and selected it for ST-OPVs. We then optimized the interfacial properties of ultra-thin Ag and maximized the selective absorption via optical design. As a result, we developed a novel ST-OPV device structure with ultra-thin Ag capping and a TeO₂ anti-reflection layer, and achieved a high light-utilization efficiency of 4.0%, as shown in Fig. 3a.

An ST-OPV must convert enough photons into energy to qualify as a PV device. However, the above definition of S does not include the energy-conversion process. Therefore, we comprehensively incorporated energy conversion and selective absorption to optimize ST-OPVs. As shown in Fig. 3b, we used this logic to reappraise the strategies proposed to optimize the device performance of opaque OPVs in detail, including active-layer thickness, D:A ratios, and ultra-thin Ag film. As a result, we were able to further improve the LUE of ST-OPVs to over 5.0%, which is among the highest for ST-OPVs (Guan et al., 2022). We found that the device exhibited a good infrared-light rejection rate (IRR) of up to 89%, which is superior to commercial heat-insulation films such as 3M NV-25.

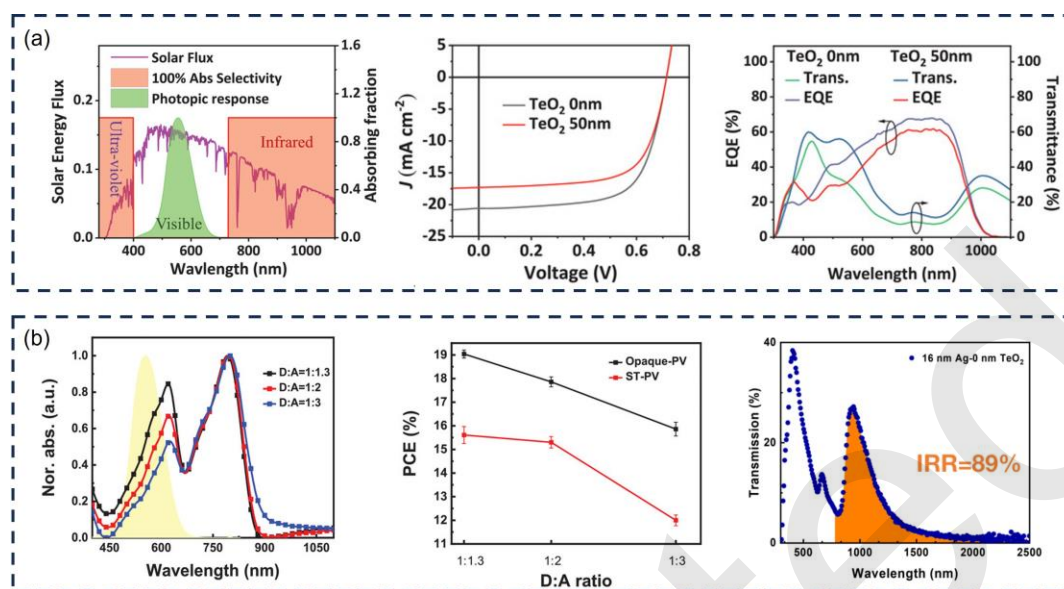


Fig. 3 (a) Solar flux, photopic response and absorption profile of absolute absorption selectivity (absorption selectivity is 100%); J-V curves, EQE curves, and transmittance of ST-OPV w/o and with 50 nm TeO_2 (Li et al., 2021). (b) Normalized absorption spectra of PM6:L8-BO:BTP-eC9 blend films with different D:A weight ratios of 1:1.3, 1:2, and 1:3, with the yellow region representing the human eye's visible response; PCE dependence on the changing D:A weight ratios and IRR of ST-OPV devices (Guan et al., 2022)

4.2 Flexible/stretchable OPVs

Organic semiconductors offer great advantages in terms of both optoelectronic properties and superior flexibility or stretchability. This is because they consist of intramolecular covalent bonds and intermolecular van der Waals interactions, which allow dislocation of molecules at no cost of defect formation under stress (Fukuda et al., 2020). In addition, organic molecules are soft, and can adapt to external stress by changing their conformation. However, advanced device structures are required to make OPVs flexible or stretchable. First, all layers. (the substrate, interfacial layer, transparent electrode, and back electrode) need to be flexible. Second, the interface between each adjacent layer must be robust enough for bending or stretching cycles (Huang et al., 2023; Yang et al., 2021; Ye et al., 2023). The two major requirements cause the most critical challenges for flexible OPVs.

We previously demonstrated that the ultra-Ag-based top-illumination (TUA) device structure is promising for flexible OPV application (Zuo et al., 2014). Different to previous designs in which the light penetrates from the substrate, the TUA structure consists of an electrode on the substrate, interfacial

layer, active layer, interfacial layer, ultra-thin Ag layer, and top optical capping layer, sequentially; and thus, it allows the light to reach the device directly without penetrating the substrate. This structure allows ease of integration with the substrate, good flexibility, low-cost production, and good scalability. Moreover, it enables a device performance of over 7% on both flexible and rigid substrates, which is quite good (Zhao et al., 2021). Taking advantage of recent advancements in NFA materials, we also fabricated high-performance flexible OPVs based on the TUA device structure, which exhibited a certified efficiency of 12% (Zhao et al., 2022). By tailoring the interfacial contact properties using the PCP-Li as the bridging layer, we successfully deposited aqueous PEDOT:PSS on the thick Ag layer. This allowed us to reach an even higher efficiency of 16% (Zheng et al., 2022). We improved the ultra-thin Ag film quality via the wetting method, and further enhanced device efficiency to over 17%, which represents the best among flexible OPVs (Zheng et al., 2023a). The device also showed hardly any change in efficiency after bending 100,000 times with a radius of 4 mm. In addition, we found that up-scaling the device area from 0.1 to 1 cm^2 caused no loss in efficiency, and device color could be easily tuned by simply adjusting the

capping layer, as shown in Fig. 4a. Aside from the intrinsic factors, OPV flexibility is also related to the geometry factor, with thinner devices delivering better flexibility. Without a substrate, the thickness of a TUA device is around 300 nm, which should be very flexible. However, fabricating “free-standing” high-performance OPVs or integrating them with ultra-thin substrates is quite challenging. An advantage of the TUA device is that it can be easily integrated with a variety of substrates, regardless of its optical transparency and film thickness. Therefore, we tried integrating the device on a 1.2-micro-meter polyimide substrate, and found that the device's performance was very similar to that on a glass substrate. However, this ultra-thin OPV had different mechanical properties. First, it exhibited a very high specific power density of 40 W/g. Second, such a thin device makes a small bump when compressed, which allows for pseudo-stretchability. Third, this device was adaptable to most non-planar surfaces, and easy to integrate with most surfaces, including human skin.

Bending the device exerts a combination of stretching and compressing, but with a low strain (~1%). While stretchable OPVs must undergo a large strain over 10%, which means more harsh requirements on materials and devices. Thereafter, the flexible OPV can be applied in more diverse scenarios compared to flexible ones. We took on this challenge by expanding the TUA structure of the substrate, back electrode, active layer, and top electrode. As shown in Fig. 4b, we used a polyurethane elastomer as the substrate; the back electrode consisted of PEDOT:PSS and Ag composites, and an elastic polymer (SEPS) was blended with the active layer to enhance its stretchability. Ag nanowires were used as a stretchable transparent electrode for ultra-thin Ag replacement (Zheng et al., 2023b). As a result, we achieved record-breaking efficiency of 16.23%, and after stretching for 200 cycles with a strain of 10%, the device retained 80% of the efficiency before stretching.

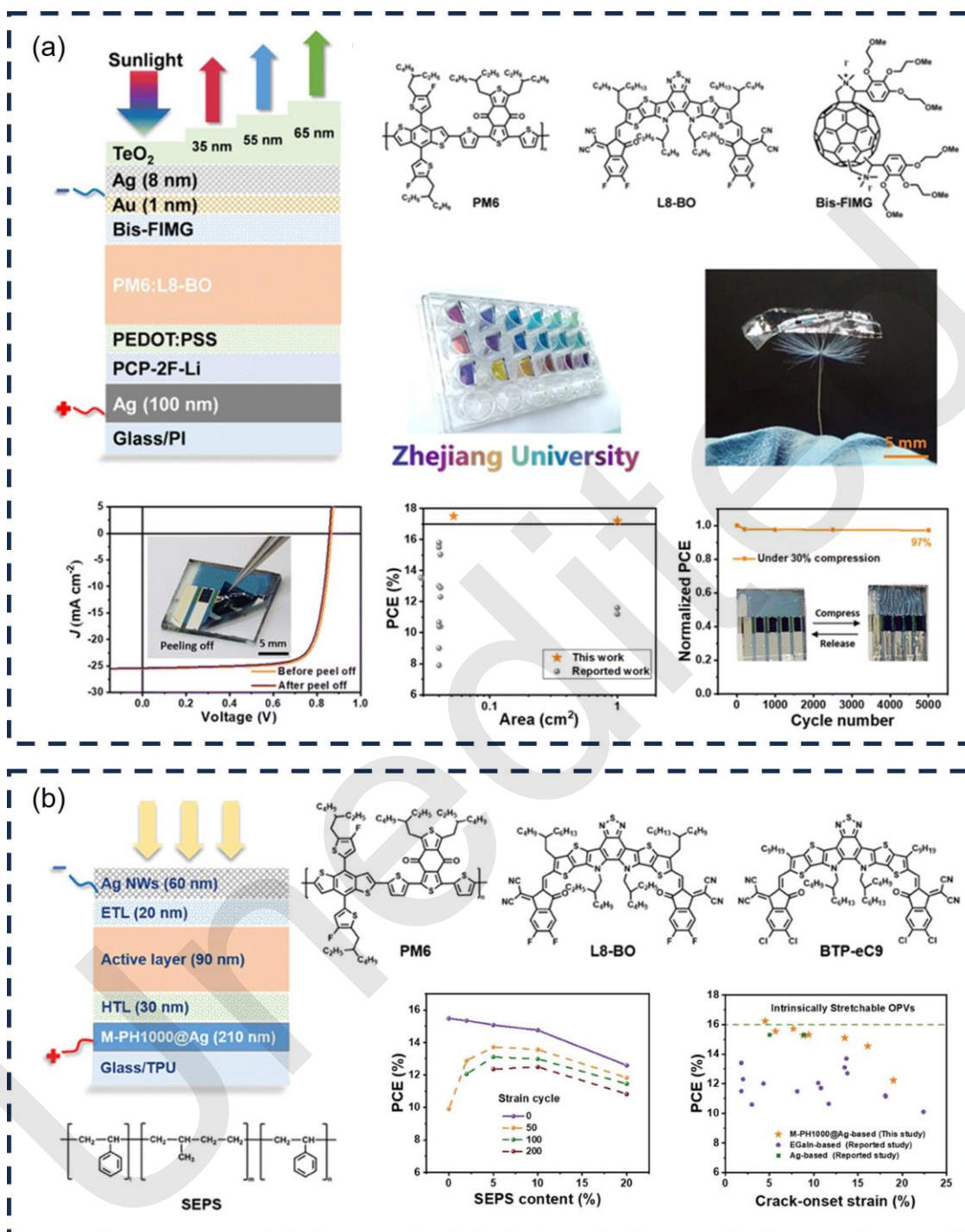


Fig. 4 (a) The structure and performance of an Ultra-thin OPV with a power density of nearly 40 W g⁻¹ (Zheng et al., 2023a). (b) The structure and performance of an intrinsically stretchable OPV (Zheng et al., 2023b)

5 Summary

In the past three years, our group has focused on addressing the V_{OC} loss issue in OPVs, and successfully reduced the non-radiative charge recombination

induced by electron-vibration coupling by tailoring intermolecular interaction. In particular, we propose using the dilution effect in producing multi-component OPVs, which inspires the general material-selection rule for high-performance OPVs. We also unveil the details underlying the local polariza-

bility effect and molecular packing effect. We have made efforts to improve the performance of multi-functional OPVs, including semitransparent devices and flexible/stretchable devices. Based on a fundamental understanding of the optical, electrical, thermal, and mechanical properties of OPVs, we propose the concept of absorption selectivity, and have designed novel devices that offer more and better functionality.

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Author contributions

Yiming WANG and Lijian ZUO wrote and revised the manuscript.

Conflict of interest

Yiming WANG and Lijian ZUO declare that they have no conflict of interest.

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概要: 由于半透明性、轻量化、优异的柔性和低成本等优点, 有机光伏器件被视为下一代可再生能源的有望来源, 使其在未来拥有更多样化的应用前景。与主流的硅基光伏技术相比, 有机光伏器件的效率仍需提高, 并且其在广泛场景中应用多功能性的潜力也需要进一步探索。本综述总结了我们在过去三年中取得的最新研究成果, 重点关注解决进一步提高器件效率的关键挑战, 并开发具备多功能性的有机光伏器件, 如半透明性、柔性、可拉伸性和美观性等。

关键词: 有机光伏; 多组分; 半透明; 柔性

中文概要

题目: 高性能多功能有机光伏器件