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Open-circuit voltage shifted by the bending effect for flexible organic solar cells

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The shift of the open-circuit voltage \( (V_{oc}) \) of flexible organic solar cells (OSCs) under bending conditions was investigated by fabricating bi-layer heterojunction and polymer-based OSCs on flexible polyethylene terephthalate (PET) substrates. To realize the performance variations of flexible solar cells characterized by important parameters, \( V_{oc} \) was measured when the substrate was bent under various curvatures. The \( V_{oc} \) was increased and decreased by using tensile and compressive stresses, respectively. The ratio of increase for \( V_{oc} \) is larger than the ratio of reduction, thus indicating that the intermolecular distance of an organic semiconductor is difficult to change because of the strong electrostatic repulsive force. A quantitative analysis of energy level by photoluminescence spectrum, UV-visible absorption spectrum, and quantum chemical calculation at various bending states was used to explain the \( V_{oc} \) as a function of bending curvature. The peak shifts of UV-visible absorption and photoluminescence spectra provide direct evidence on the variation in energy levels when devices are bent, which causes \( V_{oc} \) shifts. For bended organic semiconductor films, the bending curvature-dependent intermolecular distance was studied by Raman spectroscopy by analyzing the intermolecular couple energy. This study shows that the change of \( V_{oc} \) cannot be neglected in the application of flexible OSCs on a flexible loading circuit.

Introduction

Organic solar cell (OSC) devices have been rapidly developed in the past few years. Both small molecule and polymer organic materials have significant potential in photovoltaic cell fabrication because of their broad absorption band, which coincides with the solar spectrum, and simple fabrication technologies, such as spin-coating, inject printing, thermal evaporation, or roll-to-roll process. Furthermore, the power conversion efficiency (PCE) of OSC devices has been rapidly increasing. The plasticity and low fabrication cost of OSCs are the main factors that give these materials significant potential in next-generation solar power. The intramolecular characteristics of organic materials benefit OSCs that are applied on flexible electronics, such as OSC formations on polyethylene terephthalate (PET) substrates, which can bend and stretch without breaking. Flexible solar cells have been developed to power satellites. Flexible solar cells are lightweight, can be rolled for launch, and are easily deployable. Thus, flexible solar cells are suitable for various applications. Moreover, flexible solar cells can also be sewn into backpacks or outerwear. Flexible devices offer advantages to the development of portable merchandise. Flexible electronic devices frequently encounter bending. Thus, the influence of bending on the electronic functionality of flexible devices is an important issue. Previous studies have shown that the most obvious bending effect on OSC devices is the change of short-circuit current density (\( J_{sc} \)). The \( J_{sc} \) of a flexible device will change when the device is distorted by an external bending force. However, few reports have focused on the variation of open-circuit voltage (\( V_{oc} \)).

The physical mechanism of \( V_{oc} \) when a flexible device is under bending states is also unclear.

When a flexible device is bent, the microstructure of its organic films will be altered. Microstructure properties influence charge dissociation, charge transport, charge collection, and device performance. Obvious changes can be observed in many physical and electronic features while a flexible device is bent, for example, changes in \( J_{sc} \) and \( V_{oc} \). This study investigated the physical mechanisms of \( V_{oc} \) variation under bending. The results of this study will elucidate the issue of the bending effect on flexible OSC devices. Moreover, \( V_{oc} \) is an important parameter for loading devices on a flexible circuit because the loading process is a decisive factor in the performance of loading devices.

Pentacene and N,N’-diheptyl-3,4,9,10-perylenetetracarboxylic diimide (PTCDI-C6H13) were used as the donor and acceptor of active layer materials, respectively, for flexible OSCs fabricated on flexible PET substrates. Apparent shifts in \( V_{oc} \) were obtained for flexible OSCs bent under tensile and compressive stresses. The \( V_{oc} \) of OSCs increased and decreased when the device was under tensile bending and compressive bending, respectively. To illustrate the mechanism of \( V_{oc} \) under a bending state and identify the variations in the energy gap, photoluminescence (PL) and UV-visible absorption spectra were used to analyze the optical property variations of organic thin films during bending. During bending, the microstructure variation of the active layer was investigated by using Raman spectroscopy. The results of Raman...
and optical measurements support the effect of microstructure changes on $V_{oc}$ during bending states. Quantum mechanical calculations were used to verify the relationship between the change in molecular structure and the variation of $V_{oc}$. The analysis results for the effect of the bending effect on flexible OSCs can be potentially applied to photovoltaic devices that are used under winding to determine real power output.

**Experimental methods**

**2.1 Devices Fabrication**

A flexible PET film that was previously coated with an Al-doped ZnO (AZO) layer with a sheet resistance of 25 $\Omega/\square$ was used as the OSC substrate. The AZO layer served as the anode of OSCs and was patterned by using hydrochloric etching. The patterned substrates were cleaned in an ultrasonic bath with de-ionized water, acetone, and isopropanol for 15 min each. The substrates were then dried inside an oven at 80 °C. After drying, the substrates were treated by oxygen plasma for 5 min. Poly (3,4-ethylenedioxythiophene) (PEDOT:PSS, H.C. Starck) was spin-coated on PET substrates at 500 rpm/5 s and then spun at 2500 rpm/60 s to form a 30 nm-thick film. All samples were baked at 150 °C for 30 min to remove residual solvent from the film. A 45 nm-thick pentacene layer, a 45 nm-thick PTCDI-C$_{60}$H$_{15}$ layer, and an 8 nm BCP layer were deposited in sequence at a deposit rate of 0.5 Å/s under a pressure of 5 × 10$^{-6}$ Torr. The top electrode was constructed by using 10 nm-thick Ag and 60 nm-thick Al films. The top electrode was used as a cathode and was thermally deposited through a shadow mask to define the active area. The sizes of the active area of the small molecule-based OSCs are 0.5 mm × 0.5 mm. To extend our results, polymer-based OSCs consisted of poly(3-hexylthiophene) (P3HT) blends with indene-C$_{60}$bis-adduct as active materials were fabricated on the same flexible substrate. The sizes of the active area of the polymer-based OSCs are 3 mm × 2 mm.

**2.2 Measurements and Analysis**

Several acrylic holders with different curvatures were used in all electrical and optical measurements to bend the device into specific tensile or compressive conditions. Holder curvatures $K$ of 0, 0.33, 0.5, and 1 cm$^{-1}$ were used in this study. Holder curvatures were defined as the reciprocal of the curvature radius. The symbols “+” and “−” marked in front of the curvature represent that the device was bent under tensile and compressive states, respectively (Figure 1). The current density—voltage ($J$–$V$) characteristics of the devices were measured in a nitrogen-filled dark glove box. The $J$–$V$ curves were obtained by using a Keithely 2400 source meter, which were controlled via Labview. An Oriel solar simulator was used to illuminate the devices from the AZO side with a light intensity of 100 mW/cm$^2$ in AM 1.5 G condition. To realize the bending effect on OSCs, PL and Raman measurements (Jobin Yvon LabRAM HR system) were performed on active OSC layers to obtain the optical responses under bending states. The organic films were excited by 532 and 632.8 nm lasers for PL and Raman, respectively. The spectrometer resolution of this measuring system was 0.2 cm$^{-1}$. The UV-visible absorption spectra were measured by using a Cintra

![Fig. 1](image_url)  
**Fig. 1.** Schematic illustration of a flexible organic solar cell which is bent under various curvature states. The curvature $K$ of the solar cell is defined as the reciprocal of radius $R$ of the curvature.

![Fig. 2](image_url)  
**Fig. 2.** (a) Current density—voltage characteristic of a small molecule photovoltaic cell at the flat state. (b) Curvature dependence of open-circuit voltage ($V_{oc}$) for the flexible solar device. (c) The changing of $J_{sc}$ and $V_{oc}$ of unbent device which was under long time operating.

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The highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) of pentacene and PTCDI in the two-dimer system were calculated by DFT at the CAM-B3LYP/6-31G(d) level by using the counterpoise method and considered the dimer-centered basis set. The difference between the HOMO and LUMO of a molecule was defined as the band gap, and the difference between the HOMO of pentacene and the LUMO of PTCDI was denoted as the build-in energy ($V_{b0}$). All computations were performed by using the program suite of Gaussian 09.18

**Results and discussion**

The $J_{sc}$, $V_{oc}$, fill factor (FF), and PCE of an unbent OSC are 2 mA/cm$^2$, 333 mV, 0.44, and 0.294%, respectively [Figure 2(a)]. The $J−V$ characteristics of devices bent under different curvatures are shown in Figure 2(b). The $V_{oc}$ value shifts in different trends when the OSC device underwent tensile or compressive bending. In organic OSCs, the change of $V_{oc}$ is hardly obtained because the S value [in Eq. (1)] for organic heterojunctions is small; approximately 0.119.

$$V_{oc} = S_0 \times V_{b1} - S_1 \times V_{b2} - S_2 \times V_{b3} + C$$  

(1)

where $C$ is a constant that describes the interface potential of the ideal ohmic contact, and the parameters $S_0$, $S_1$, and $S_2$ are the base on the index of the interface behavior.20 The $V_{b1}$, $V_{b2}$, and $V_{b3}$ in Equation 1 are the potential difference between acceptor LUMO and donor HOMO, the work function differences of donor/anode and acceptor/cathode, respectively. The $V_{oc}$ value decreases from 335 mV to 324 mV (a decrease of 2.9%) when the devices were bent from a flat state to a compressive state. Contrary to the compressive condition, the $V_{oc}$ increases by 3.42% when the flexible OSC was significantly bent to a tensile state with curvature of +1 cm$^{-1}$ (Figure 3). When the bending force applied on the OSC was released, the electrical characteristic returned to its original state, thus indicating that the band structure of an organic semiconductor could be affected by the bending effect. To investigate the effect of incident light intensity in the $V_{oc}$ of bent devices, Figure 2(c) shows that although the decay of $J_{sc}$ is observed under a long time interval, $V_{oc}$ remains constant. Therefore, no apparent correlation between $J_{sc}$ and $V_{oc}$ was observed in our flexible OSC. The active areas of small molecule- and polymer-based OSCs are 0.25 mm$^2$ and 6 mm$^2$, respectively, which result in corresponding light incident angles of approximately 1.5$^\circ$ and 8.5$^\circ$, respectively, at a bending curvature of 1 cm$^{-1}$. We believe that the $J_{sc}$ will be slightly affected by the effect of incident light intensity because of a small variation in the light incident angle. Despite this effect, the change in $V_{oc}$ remains hardly observable; accordingly, the effect of incident light intensity on the $V_{oc}$ shift at bending states can be ignored.

To realize the behavior of the $V_{oc}$ shift induced by the bending effect, the PL and UV-visible absorption spectroscopy measurements were performed on organic semiconductor films to study the fine structural changes under bending states. Figure 4a shows the PL spectra of a PTCDI-C$_{7}$H$_{15}$/pentacene thin film that was bent to various curvatures. Given that the luminescent signal of the PTCDI-C$_{7}$H$_{15}$ film was stronger than the luminescent signal of the pentacene film, the PL signals are dominated by PTCDI-C$_{7}$H$_{15}$. The main PL peaks of PTCDI-C$_{7}$H$_{15}$ film are located between 670 and 700 nm. Significant shifts of the peaks are observed within the PL spectra when various bending curvatures are conducted on the organic semiconductor film. A conspicuous blueshift of the PTCDI-C$_{7}$H$_{15}$ peaks appears on the PL spectra at the tensile region. An opposite redshift phenomenon is observed in the compressive region, thus indicating that the band gap of PTCDI-C$_{7}$H$_{15}$ increases in tensile bending and decreases in compressive bending. The gray curvy line in Figure 4a shows the level of the peak shift at different bending states. Figure 4b shows the UV-visible absorption spectra of the PTCDI-C$_{7}$H$_{15}$/pentacene film at various bending states. The
absorption bands of the PTCDI-C7H15 and pentacene films range from 500 nm to 600 nm and 600 nm to 700 nm, respectively\(^2,\)\(^2,\)\(^2\) The peak shift trend of the UV-visible spectra varies with the bending curvature and is the same as that of the PL spectra. Furthermore, the absorption peak shift of pentacene follows that of PTCDI-C7H15 (Figure 4c). Both optical results indicate that the band gaps of donor and acceptor molecules were reduced when the device was compressed. The band gap of both active layers increased during the tensile condition compared with the compressive condition. To clarify the mechanism of \(V_\text{oc}\) shift affected by the bending effect, the \(V_\text{oc}\) shift of the bent device and the peak shift of PL are compared in Figure 5. Interestingly, the variation of \(V_\text{oc}\) as a function of bending curvature closely approaches the behavior of the PL peak shift for the PTCDI film. This result indicates that the decrease in \(V_\text{oc}\) is caused by the reduction of the band gap for both active films during compressive states. When both active films are bent at tensile state, the enlargement of the band gap increases \(V_\text{oc}\). Thus, the \(V_\text{oc}\) shift of OSC can be attributed to the band gap variations of the active layers at the bending state. Previous researches have reported that band gaps are strongly dependent on the molecular distance of organic semiconductors\(^2,\)\(^2\) Moreover, the molecular distance within a thin film may be changed by external compressive or tensile stress. Therefore, further studies of the relationship between the changes of band gaps and molecular distance is needed to clarify the mechanism of \(V_\text{oc}\) shift at bending states.

To investigate the relationship between molecular distance changes and band gap shifts within a bending device, Raman spectroscopy was used to analyze the bending effect of organic semiconductor films. Figure 6 shows the normalized Raman spectra of pentacene with the vibrational band ranging from 1140 cm\(^{-1}\) to 1190 cm\(^{-1}\) is dedicated by the C-H in-plane bending of pentacene\(^3\). Both the 1158 and 1178 cm\(^{-1}\) bands are H-atoms with fully symmetric vibrations located at the end and on both sides of the pentacene, respectively. The vibrational band at 1155 cm\(^{-1}\) (denoted as \(v_1\) band) was split from the 1158 cm\(^{-1}\) band, which originates from the different interaction energy between the two adjacent pentacene molecules in a unit cell and is analogous to Davydov splitting. Thus, the \(v_1\) band appears in all spectra when the pentacene molecules are stacked to form a thin film. Given that the pentacene film is subjected to the compressive force, the \(v_1\) band has an obvious redshift away from the 1158 cm\(^{-1}\) band (Figure 6a). By contrast, the \(v_1\) band has a blueshift close to the 1158 cm\(^{-1}\) band when the pentacene film is bent at the tensile state (Figure 6b). The 1158 cm\(^{-1}\) is the vibration signal of a single pentacene molecule; therefore, no apparent shift can be observed in this vibrational band under bending states. According to the concept of Davydov splitting, the coupling energy \((\omega_1)\) of pentacene molecules can be calculated from the splitting level of the \(v_1\) and 1158 cm\(^{-1}\) bands, thus indicating that the coupling energy increases when the device is under compressive bending. In weak coupling, the vibrational coupling energy \(\omega_1\) can be calculated by the following equation:

\[
\Delta \omega \approx \frac{\omega_1}{2 \omega_0}
\]

where \(\Delta \omega\) and \(\omega_0\) are the amount of Davydov splitting and internal molecular band frequency \((\omega_0 = 1158 \text{ cm}^{-1}\) in this case), respectively. The \(\omega_1\) value of unbent pentacene is 11.38 meV. Compared with the \(\omega_1\) value of unbent pentacene, \(\omega_1\) increases 0.23 meV, (an increase of 2.04%) at a compressive bending of curvature \(-1\) and decreases 0.37 meV (a decrease of 3.22%) at a tensile bending of curvature of \(+1\) (Figure 3), thus indicating that the intermolecular distance of pentacene is reduced by compressive bending to result in the increase of coupling energy. The variation of coupling energy and \(V_\text{oc}\) shift has the same variation behavior as the bending curvature. It is noted that the vibrational band and \(V_\text{oc}\) shifts at tensile bending are more obvious than those at compressive bending because of the difficulty in changing the distance of pentacene molecules in a unit
cell. This difficulty is caused by the strong electrostatic repulsive force between pentacene molecules that are subjected to compressive force. Thus, the molecular spacing can be easily changed by tensile force compared with compressive force. Furthermore, the intensity of the 1178 cm⁻¹ band increases and decreases with the increasing curvatures of tensile bending and compressive bending (raw data not shown), respectively. This result indicates that the vibrational freedom degree of the H-atoms at both sides of the pentacene increases with the increasing molecular spacing in tensile bending. Thus, molecular spacing is affected by substrate bending, which leads to changes in electronic coupling energy and band gap.

The C–C aromatic stretching modes of pentacene are located between 1330 and 1390 cm⁻¹ (Figure 6). Three main vibrational bands at 1351, 1371, and 1379 cm⁻¹ can be easily observed in a Raman spectrum. Among these bands, 1351 and 1378 cm⁻¹ bands are affected by horizontal stress forces easily, 1351 cm⁻¹ band especially. The full width at half maximum (FWHM) of 1351 and 1378 cm⁻¹ bands increases with increasing tensile bending curvature. Nevertheless, FWHM decreases with increasing compressive bending curvature. This result shows that the degree of C–C vibration freedom increases when the pentacene film is under a tensile bending state, thus indicating that the intermolecular spacing of pentacene in a unit cell increases at tensile bending conditions again. Both Raman shifts of C–H and C–C aromatic stretching modes imply that the molecular spacing and energy level of pentacene film changes by applying compressive or tensile stress. In conclusion, for the variations of Raman bands, PL, and absorption characteristic peaks at bending conditions, the band gap and V_{oc} decrease with decreasing molecular distance under compressive bending. By contrast, the molecular distance was enlarged under the tensile bending to increase the band gap and V_{oc}. In OSC devices, the difference between the HOMO of the donor and the LUMO of the acceptor is considered a built-in potential, which is an important parameter of OSC. Earlier studies indicate that the V_{oc} of OSC devices is generally influenced by the built-in potential. However, how the bending curvature affects the HOMO and LUMO of active layers is still unclear.

To realize the changes of the HOMO and LUMO of PTCDI-C_{6}H_{13}/pentacene films at various bending conditions, a simple face-to-face structure for a dimer in a unit cell was used to calculate the HOMO and LUMO at a bending state (Figure 7a). The set distance of the dimers at a flat state is 4 Å, and the two dimers are vertically separated by a gap of 2 Å. The quantum mechanical calculations display that the HOMO of the pentacene and PTCDI-C_{6}H_{13} dimers increases with increasing dimer distance and that LUMO decreases with increasing dimer distance (Figures 7b and 7c, respectively). Figures 7d and 7e show the band gaps of the pentacene and PTCDI-C_{6}H_{13} dimers as a function of the dimer distance, respectively. The dimer distance is obtained from the difference between HOMO and LUMO. The band gaps of pentacene and PTCDI-C_{6}H_{13} increase with increasing dimer distance, thus indicating that the band gap decreases in the compressive stress performed on organic films and increases in the tensile case. These simulation results of the band gap completely coincide with the results obtained from PL and the absorption measurements in banding states. Figure 7f shows V_{oc} as a function of the dimer distance that is defined as the difference between the HOMO of donor and the LUMO of acceptor. The V_{oc} also increases with increasing dimer distance; this result corresponds with the result of the V_{oc} shift at bending states because V_{oc} is proportional to V_{bi}. When the current-voltage curves in the dark intersect with the curves under illumination, the compensation voltage V_{oc} is obtained in the inset of Figure 8. A comparison between the curvature-dependent V_{oc} (Figure 8) and the curvature-dependent V_{bi} (Figure 3) shows that both voltage trends tend toward coincidence. Moreover, the value of V_{bi} is very close to the built-in potential V_{bi}. The trends of the curvature-dependent V_{bi} and V_{oc} are almost the same; that is, the simulated HOMO and LUMO levels for the donor and acceptor under bending states are reasonable.

We extended this investigation and showed that the V_{oc} of polymer-based flexible OSCs can also be shifted by the bending effect. Polymer-based OSC consists of poly(3-hexylthiophene) (P3HT) blends with indene-C_{6}bis-adduct, which was formed on the same polyethylene terephthalate substrate. The polymer-based OSC had a power conversion efficiency of 1.82%,
In summary, we performed a series of $V_{oc}$ measurements at different bending curvatures that play important roles in the performance of flexible OSCs at bending states. By increasing the bending curvature of the PET substrate, $V_{oc}$ is enhanced at tensile bending and is reduced at compressive bending. The asymmetry of the variation ratio for $V_{oc}$ at tensile and compressive states indicates that the intermolecular distance is difficult to change at compressive conditions because of the strong electrostatic repulsive force between semiconductor molecules, such as pentacene molecules and PTCDI-C$_2$H$_5$ molecules. PL spectroscopy, absorption spectroscopy, and Raman spectroscopy were used to investigate the intermolecular distance of organic semiconductor films at bending states. The intermolecular distance increases in the tensile condition and decreases in the compressive case, thus resulting in corresponding variations in the band gap and $V_{oc}$. We also predicted the HOMO and LUMO eigenvalues of pentacene and PTCDI-C$_2$H$_5$ dimers by employing DFT function at various bending conditions to simulate the variations of band gap and $V_{oc}$. The HOMO level increases with increasing intermolecular distance. On the contrary, the LUMO level decreases with increasing intermolecular distance. According to the proportion of $V_{oc}$ to $V_{oc}$, the swing of $V_{oc}$ at the bending states can be explained by quantum mechanical calculations. Thus, these experimental and quantum mechanical calculation methods can be powerful tools for analyzing molecular electronics and predicting molecular-orbital eigenvalues, respectively, for flexible devices at bending states.

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