Synthesis, structure, and opto-electronic properties of organic-based nanoscale heterojunctions

Bohuslav Rezek1*, Jan Čermák1, Alexander Kromka1, Martin Ledinský1, Pavel Hubík1, Jiří J Mareš1, Adam Purkrt1,2, Věra Cimrová2, Antonín Fejfar1, Jan Kočka1

Abstract

Enormous research effort has been put into optimizing organic-based opto-electronic systems for efficient generation of free charge carriers. This optimization is mainly due to typically high dissociation energy (0.1-1 eV) and short diffusion length (10 nm) of excitons in organic materials. Inherently, interplay of microscopic structural, chemical, and opto-electronic properties plays crucial role. We show that employing and combining advanced scanning probe techniques can provide us significant insight into the correlation of these properties. By adjusting parameters of contact- and tapping-mode atomic force microscopy (AFM), we perform morphologic and mechanical characterizations (nanoshaving) of organic layers, measure their electrical conductivity by current-sensing AFM, and deduce work functions and surface photovoltage (SPV) effects by Kelvin force microscopy using high spatial resolution. These data are further correlated with local material composition detected using micro-Raman spectroscopy and with other electronic transport data. We demonstrate benefits of this multi-dimensional characterizations on (i) bulk heterojunction of fully organic composite films, indicating differences in blend quality and component segregation leading to local shunts of photovoltaic cell, and (ii) thin-film heterojunction of polypyrrole (PPy) electropolymerized on hydrogen-terminated diamond, indicating covalent bonding and transfer of charge carriers from PPy to diamond.

Introduction

Electronic devices nowadays are commonly based on inorganic semiconductors (e.g., silicon or germanium) and, thus, face the problem of high-cost industrial processes (high-vacuum technology, clean rooms, or high-purity source materials). A possible way to reduce costs is the use of organic materials with semiconducting or metallic properties. Although the first report on the conductivity of doped polypyrrole (PPy) was published in 1963 [1], the breakthrough is attributed to the studies on doped polyacetylene since 1977 by Heeger, Shirakawa, and MacDiarmid [2], for which they were awarded the Nobel prize in 2000. Since then organic semiconductors have followed similar scientific evolution as inorganic ones (all-polymer field effect transistor in 1994 [3], organic integrated circuit by the Philips company (1998)) until recently, which have seen application of organic displays in cell phones, and the start of commercial production of organic photovoltaic (PV) cells.

PV effect in organic materials is different from inorganic ones. The binding energy between photo-excited electron-hole pair is strong due to the low dielectric constant, typically in the range of 0.1-1 eV. Therefore, the excitons are not dissociated by thermal energy, which is approximately 26 meV at room temperature. Additional driving force is needed, which can be supplied by introducing a layer of a second organic material (the so-called double-layer cell, Figure 1a). Typical PV power conversion efficiencies of such devices are not higher than 0.1% [4,5] because of the short exciton diffusion length (around 10 nm [6]) compared to the total film thickness needed for efficient light harvesting (100-200 nm).
Significant improvement emerged with the bulk-heterojunction design [7]. In the bulk-heterojunction design both materials (donor and acceptor) are prepared as a microscopic interpenetrating network (Figure 1b). Therefore, wherever the excitons are generated, an interface with the other material is always closer than the exciton diffusion length (in an ideal case). The remaining task is to ensure efficient charge carrier transport to electrodes. So far the best reported PV efficiency is the attainment of reaching 7.4% [8].

For a long time, electronics has been the domain of inorganic materials. Even though electronic behavior of organic materials has been known for several decades, their applications are still limited. So far, the research and development activity of organic and inorganic electronics has been strictly divided, although their combination might be fruitful. An example of promising organic-inorganic (hybrid) systems is the dye-sensitized PV cell. In this so-called Grätzel cell [9], a photoexcited organic dye gives electrons to the electrode via porous inorganic TiO2. The missing electron is returned from the surrounding electrolyte, which restores the original state at the counter electrode. Although the PV power conversion of this type of PV cells is relatively high (10-12%), their wider application is limited by the need of an electrolyte, which is commonly in liquid form.

Another example of an organic-inorganic system, which is extensively studied, is diamond in combination with organics, e.g., fullerene [10,11]. Such systems are highly promising for bio-sensoric or opto-electronic applications, as a charge transport between the two materials is observed. However, the basic electronic properties at their interface still need to be fully understood.

From the electronic point of view, diamond is a wide bandgap (5.5 eV) semiconductor, which in its intrinsic form is electrically insulating. Apart from the bulk conductivity induced by doping [12], intrinsic diamond exhibits a special phenomenon of surface conductivity when a thin (10-20 nm) electrically conductive ($p$-type) layer is formed at the surface [13-15]. It is observed under ambient conditions when the diamond surface is terminated by hydrogen atoms. Various electrically conductive areas or channels can be patterned on the surface by changing the surface termination [16,17]. Hydrogen termination can also be used as a starting surface for grafting of more complex organic molecules on diamond [10,18-20].

In this study, we have chosen PPy because of its wide universality as a model of a chemically and optically sensitive organic dye. Its polymerization can be achieved by electrooxidation from a solvent [21], chemical vapor deposition [22], UV irradiation [23], or chemical polymerization [24,25]. PPy is a well known yet still remains under intensive study in many fields of applications, like sensors [26-28], biosensors [29,30], fuel cells [31,32], corrosion protection [33], or rechargeable batteries [34,35].

Organic-based electronic as well as optoelectronic devices are commonly made from several compounds and their mutual electronic cooperation at microscale is of key importance for the device properties. Therefore, revealing and understanding of microscopic structural, chemical, electronic, and optoelectronic properties is crucial for their further improvements. Such challenging task may be fulfilled by techniques based on local probe scanning. One of the prominent methods employing scanning probe is atomic force microscopy (AFM) with its diverse regimes of operation [36]. In AFM, a sharp tip mounted on a flexible lever (the so-called cantilever) can detect both morphologic and electronic information with high lateral resolution.

When the tip is in contact with the studied surface, a bias voltage can be applied between the sample and the AFM tip, and the induced electric current is dependent on the local conductivity at the position of the tip. As a result, maps of both topography and local conductivity are obtained at once in this so-called current-sensing AFM regime (CS-AFM) [37]. Applying this mode on organic materials is rather difficult, however, as the soft organic materials can be easily modified by the sharp AFM tip (radius typically 10-30 nm) under the applied force of typically several nN or more.

Application of an AC voltage between the sample and the AFM tip when the tip is at a certain distance from the surface creates a modulated electrostatic force that makes the cantilever oscillate because of contact potential difference. These oscillations can be minimized to zero by applying an additional DC voltage to compensate the potential difference. If the tip surface potential is calibrated, then the absolute values of the surface potential (and then the work function) can be obtained.
This is the principle of the so-called Kelvin force microscopy (KFM) [16].

In spite of being a powerful tool for resolving microscopic structural, mechanical, and electronic properties, AFM had not been able to provide direct chemical contrast until very recently [38]. Such chemical detection in AFM makes use of adhesion differences on atomic scale and was, thus so far, applicable only to atomically flat surfaces. Spatially resolved chemical information can be obtained in more straightforward way by micro-Raman spectroscopy mapping, where the focused laser beam plays the role of the scanning probe. Although it is an optical method, the resolution can be satisfactory at sub-micrometer scale [39]. The resolution can be further enhanced using special metal tip in the so-called tip-enhanced Raman spectroscopy [40]. This method works well on resonant systems such as in the case of carbon nanotubes [41], but its general application is still highly challenging.

In this study, we employ and combine advanced scanning probe techniques as well as macroscopic characteristics to provide us significant insights into the correlation of microscopic structural, chemical, and opto-electronic properties of organic-based heterojunctions. We demonstrate benefits of such multi-dimensional characterizations on (i) bulk heterojunctions of fully organic composite films using fullerene as electron acceptors, indicating differences in blend quality and component segregation leading to local shunts of PV cell [42,43], and (ii) thin-film heterojunction of PPy electro-polymerized on hydrogen-terminated diamond, indicating covalent bonding and enhanced exciton dissociation in such systems [44-47].

**Basic organic bulk heterojunction**

Basic type of organic bulk heterojunction was based on fullerene C60 as electron acceptor. Composite blends made of poly[(2,7-(9,9-dihexa)fluorence)-co-(1,4-(2,5-didecylaminoketo) phenylene)] (VYP-120, developed at the Institute of Macro-molecular Chemistry, ASCR, v.v.i.) and fullerene C60 (Sigma Aldrich) were spincoated on indium tin oxide (ITO)-covered glass substrates and dried at 50°C under vacuum for 4 h. For characterizing the PV performance of the thin film, top Al electrodes were evaporated through a shadow mask.

In spite of observed quenching of photoluminescence compared to polymer layer without the C60, the composite layer exhibited low PV power conversion efficiency (Jsc ~ 2 nA, Voc ~ 5 mV, η ~ 0.06%) [42]. AFM morphology (Figure 2a) revealed a relatively flat and smooth surface (RMS roughness: 4 nm) which was covered with two types of clusters (lateral size either 100 nm or several μm) and dendrites (more than 10 μm). Electrical potential map obtained by KFM at the same area is shown in Figure 2b. KFM detected the highest surface potential (up to 50 mV) in the central part of the dendrite and the lowest surface potential (as low as -150 mV) in its immediate surrounding. Farther surroundings exhibit the potential between these two values. Typical value is around ~30 mV. Also fluctuations related with small clusters are seen in topography. As the layer is made of two materials, the observed variations in surface potential most likely correspond to variations in local chemical composition. The two extreme potential levels may correspond to individual materials while the intermediate potential is related to the blend.

This assertion is proven by micro-Raman spectroscopy and mapping. The spectrum collected on the dendrite showed a sharp peak at 1468 cm⁻¹, which was attributed to fullerene C60, as well as a broad band at 1400 cm⁻¹ [42,48]. Similar spectra, although with lower peak intensity, were detected on the small clusters in the surroundings. The map and the spatial profile of C60 characteristic Raman scattering peak at 1468 cm⁻¹ are shown in Figure 2c. The image reveals higher concentration of C60 at the dendrite and low C60 Raman signal in its close surroundings.

Spectrum detected in the dendrite vicinity did not show the sharp peak, only a broad band around 1400 cm⁻¹ [42]. Similar spectra were obtained on the smooth surface in the farther surroundings. This band is not present in typical C60 Raman spectrum [48]. Hence, we attribute it to a photoluminescence (PL) background due to Raman laser excitation (note that it cannot be directly compared with usual PL spectra). Spatially resolved map and the profile of Raman intensity at the 1400 cm⁻¹ band are shown in Figure 2d. It is noteworthy that both Raman maps are qualitatively quite similar. Yet, quantitative comparison of the spatial profiles shows differences: the C60 signal is higher on the dendrite and small clusters, while the PL intensity prevails over C60 on the smooth surface and the very vicinity of the dendrite. We suggest that the map of sharp peak at 1458 cm⁻¹ is related with segregated crystallized C60 on the dendrite and on small clusters in the surroundings. The broad band at around 1400 cm⁻¹ may be attributed to C60 that is highly dissolved in the blend. This assertion is supported also by the AFM and KFM data.

In both Raman maps, there is a minimum in the vicinity of the dendrite. This is not effect of thickness as the AFM shows even surface compared to surroundings. On the other hand, there is clear difference in the work function detected by KFM. This indicates that this area consists predominantly of the conductive polymer. In such a case, the dendrite vicinity is not of heterostructural nature, and the conductive polymer may electrically shunt the device.
Figure 2 Multidimensional microscopic characteristics (images and spatial profiles) of the organic heterostructure blend made of fullerene C_{60} and poly[(2,7-(9,9-diHexa)fluorene)-co-(1,4-(2,5-didecylaminoketo) phenylene)]: (a) tapping-mode AFM surface morphology, (b) KFM surface potential, (c) micro-Raman intensity at 1468 cm\(^{-1}\), (d) micro-Raman intensity at 1400 cm\(^{-1}\), and (e) local conductivity as measured by CS-AFM (negative bias voltage applied on the sample). Positions of the profiles in the images are indicated by the dashed lines and arrows.
To resolve the above effect, we characterized local electronic transport properties of the organic film by CS-AFM, so that its integrity was preserved, and its surface morphology was the same as in tapping mode. The image of local conductivity (Figure 2e: note that the applied bias voltage is negative) shows that the dendrite and farther surroundings are electrically insulating. Conductivity in the close vicinity of the dendrite (area lacking $C_{\text{ion}}$) is an order of magnitude higher. This observation corroborates the conclusion that such features can cause electrical shunting of the organic film and reduce its power conversion efficiency.

**Advanced organic bulk heterojunction**

Based on the result with pure fullerene, more advanced bulk heterojunctions were prepared, with fullerene derivatives being used as electron acceptors: [6,6]-thienylC61 butyric acid methyl ester ([60]ThCBM), and [6,6]-thienylC71 butyric acid methyl ester([70]ThCBM). Poly(3-hexylthiophene-2,5-diyl) (P3HT) was used as electron donor. The blends were prepared in an inert atmosphere on ITO-covered glass substrates with a hole-conducting polymer (PEDOT:PSS) by spin coating from a 2.5% chlorobenzene solution [43]. Both blends were annealed at 125°C (the annealing improved the PV efficiency more than twice compared with non-annealed blends). Resulting thickness of the films was 140 nm, as measured by KLA-Tencor P-10 profiler. LiF (2 nm) and Al (60 nm) were thermally evaporated to make the second contact. Active area of the prepared PV cells was 6.5 × 2.5 mm².

Figure 3a, b shows macroscopic (with both top and bottom planar contacts) $I(V)$ characteristics of the two blend films measured under white light illumination on the inert nitrogen atmosphere (0 h) and after exposure to ambient air. The PV device made of P3HT:[70]ThCBM blend exhibits initial power conversion efficiency($\eta$) of 0.71% which is reduced, after exposure to ambient air for 6 h, to $\eta = 0.37\%$. The reduction is due to the decrease in short-circuit current ($I_{\text{sc}}$) from 4.02 to 2.08 mA/cm² while open-circuit voltage ($V_{\text{oc}}$) remains 0.55 V. Similar decreases in both power conversion efficiency and $I_{\text{sc}}$ were observed also on the device made of P3HT:[60]ThCBM blend (initial: $\eta = 0.86\%$, $I_{\text{sc}} = 4.05$ mA/cm²; after 6 h: $\eta = 0.42\%$, $I_{\text{sc}} = 1.8$ mA/cm²; $V_{\text{oc}}$ remains 0.48 V). In other words, both blends exhibit similar PV performance as well as its development in time.

It is noticeable that the initial power conversion efficiency of the P3HT:[70]ThCBM blend is lower but its decrease in time is also slightly slower. Based on AFM topography (shown as inset images in Figure 3a, b), both blend films are smooth (RMS roughness around 2 nm) with shallow pits (10-20 nm deep, 100-200 nm in diameter). These pits do not affect the electronic homogeneity of the films. The films appear electronically uniform as observed by KFM [43]. Therefore, to understand the differences in PV performance, we characterized dynamic opto-electronic response by modifying standard KFM technique. We stopped the physical motion of the AFM tip in one direction so that the corresponding axis in the image represents time axis. During scanning in this modified regime we were switching on and off white light illumination repeatedly.

Resulting data (KFM images and their cross sections) are shown in Figure 3c, d. Under the white light illumination, the surface potentials of both blend films shifted. In the case of P3HT:[60]ThCBM blend, the shift of surface potential is slower. This indicates that, after the illumination is switched off, the film keeps negative charge. This behavior has already been observed also on BH PV cells containing PCBM [49] as well as all-polymer cells [50]. It was attributed to electron trapping in shallow traps [51]. Persisting negative charge may then limit the surface potential shift under repeated illumination being switched on. In the case of P3HT:[70]ThCBM, the surface potential comes repeatedly back almost to the starting level during the short term, which indicates that the electron trapping in this blend is minimized, and therefore it is beneficial for PV applications.

The absolute values of the overall surface potentials shifted over time independently on the illumination. This effect is attributed to degradation of the blend films, as the AFM/KFM characterization was performed under ambient conditions. Degradation of the blends was also observed as a decrease in photo-response. There was almost no photo-response observed after 250 min in the case of P3HT:[60]ThCBM blend, and the same state is reached after 1000 min in the case of P3HT:[70]ThCBM.

$I(V)$ and KFM characteristics indicate that the P3HT:[60]ThCBM degrades much faster than the other one. The difference between the macroscopic and microscopic performance is most likely caused by the presence of the metal electrode (in the macroscopic characterization), which partially seals the organic films. From this point of view, contact-less KFM characterization revealed different opto-electronic properties of the blends, not detected by the macroscopic $I(V)$ characteristics.

**PPy-diamond heterojunction**

For creating organic-inorganic junction, we synthesized and grafted the PPy on diamond electrochemically from pyrrole (0.24 M) and NaCl (0.1 M) aqueous solution by the application of a constant current (current density -0.3 mA/cm²) and the employment of a hydrogen-terminated intrinsic monocrystalline diamond (synthetic IIIa
CVD diamond) with conductive surface as a working electrode [44]. For the electronic transport measurements, we defined H-terminated conductive narrow channel (5 μm wide) on a monocrystalline diamond surface by selective oxygen plasma discharge treatment through a photolithographic mask. PPy was electrochemically synthesized on the channel. Synthesis and electronic measurement setups are shown schematically in Figure 4. Typical PPy growth curve is shown there as well.

Thickness of the PPy film was 25 ± 5 nm based on the height histogram of tapping-mode AFM image after local nanoshaving of the PPy film in contact mode AFM [44]. Then, we applied contact-mode AFM with increasing contact force. At certain threshold contact force, the sharp tip starts to penetrate and remove the organic film. This transition is evidenced in Figure 5. The threshold force deduced in this manner was about 40 nN. This value is comparable to the values observed on a system consisting of diamond with covalently grafted DNA molecules [20,52]. This indicates that the PPy molecules also establish covalent bonds with H-terminated diamond.

Surface potential measured by KFM on the place where the PPy film had been deposited and removed (Figure 6) is significantly lower (by about 0.1 V) than the potential on pristine H-terminated diamond surface. This change is similar to reports on DNA-diamond interfaces [53] and supports the assertion of PPy molecules being linked to diamond surface covalently while removing the H termination. We also observed that the surface conductivity disappeared after the synthesis and the removal of the PPy molecules from diamond surface, which is most likely caused by the missing H termination.

Covalent bonding between PPy and diamond was confirmed also theoretically. Calculated interaction energy of about 6 eV per bond pointed to a covalent character of the bonds formed at the one- and multi-bond contacts between PPy and diamond [54].

Figure 3 Macroscopic and microscopic opto-electronic characteristics of the organic blends (a, b) Macroscopically measured current-voltage characteristics of the blend films under illumination (white light, 80 mW) in the nitrogen atmosphere (0 h) and after exposure to ambient air for 2 and 6 h. The inset images in (a, b) show AFM topography images of the blend films (1 × 1 μm², height scale 30 nm). (c, d) Cross sections of the KFM images (shown as insets, 1 × 1 μm², potential scale 40 mV) under repeated on/off switching of illumination.

Rezek et al. Nanoscale Research Letters 2011, 6:238 http://www.nanoscalereslett.com/content/6/1/238 Page 6 of 12
Based on the above experimental and theoretical results, we propose a model that, during the electrochemical synthesis of PPy on diamond, the hydrogen atoms are removed and, instead, PPy molecules establish covalent bonds with carbon atoms of diamond.

To study optoelectronic properties at the PPy-diamond interface we first removed PPy from a small area (10 × 10 μm) using the AFM nanoshaving. Micro-Raman spectroscopy confirmed the removal of PPy [45]. This area and its surroundings were then studied by KFM in the dark and under the white light illumination. Similar to the experiments on advanced bulk heterojunctions, the KFM measurement was performed with one-scan direction disabled, and the illumination was repeatedly switched on and off (Figure 7a). Figure 7b shows temporal potential profiles that were obtained via KFM scanning across PPy-diamond and bare diamond surface (where PPy was removed) under repeated illumination switching. The positions of profiles are indicated by lines and arrows in the image. The changes of potential are relatively fast (<1 s, limited by KFM scan speed), reproducible, and well defined.

Based on the surface potentials combined with data from the literature for both materials, we were able to assemble the energy band configuration of the PPy-diamond system (Figure 8). Detailed description of the procedure for band diagram assembly is given in [45]. Theoretical calculations showed that, for the PPy-C:H interface, the charge neutrality level is located below the Fermi level, so the transfer of electrons occurs from an electrode to the PPy molecules [54]. This confirm downward band bending of PPy bands at the junction to diamond.

Under illumination, the observed positive shifts of surface potentials correspond to a decrease of work function in both PPy and diamond as indicated in Figure 8. These shifts of surface potentials were attributed to the SPV effects. Based on the SPV theory, a model of charge transfer between diamond and PPy was established [45]. This model suggests that under illumination strongly
bound excitons are generated in PPy. The excitons can be split at the interface with diamond, and the holes are trapped in diamond surface/interface states.

The manner in which these holes contribute also to sub-band gap optical excitation of free holes in diamond depends on many factors [45]. Therefore, we characterized electronic transport properties of the PPy-diamond system by $I(V)$ measurements both in the dark and under the white light illumination. In the dark, the diamond channel with grafted PPy on top was highly electrically insulating as expected due to the missing H termination. The conductivity does not recover even when we remove PPy and expose the diamond surface to the ambient air (adsorbates) again. Hence, the adsorbates play no role in this effect. Under the white light illumination, the channel with PPy turned electrically conductive (approximately 100 pA when 1 V is applied, see Figure 9a). Such changes of electrical current occur within 1 s and are reproducible [46]. This is in agreement with the fast changes in the surface potential as shown in Figure 7b.

We can explain this illumination-induced conductivity by the charge transfer of holes from PPy to diamond as suggested above. The electronic transport measurements indicate that the holes accumulated in diamond-trap states are excited to diamond valence band, where they support the in-plane conductivity [46]. Such concept is schematically illustrated in Figure 9b. Relatively quick response to the change of illumination can be explained by a quick re-trapping of photogenerated holes in diamond and their recombination with photo-electrons remaining in the vicinity of the PPy-diamond junction.
The observed enhancement of in-plane conductivity under illumination might be explained also by the bridging effect via the semi-conductive and photosensitive PPy. To resolve this issue, we have measured mobility of charge carriers using Hall effect. Mobility of holes in PPy and diamond differs by at least five orders of magnitude (H-terminated diamond: $10^0$-$10^2$ cm$^2$/Vs, PPy: $10^{-5}$-$10^{-10}$ cm$^2$/Vs). We prepared microscopic...
A conductive square (10 × 10 μm²) with electric leads at the corners by selective hydrogen and oxygen terminations. A resin layer with the opening at the position of the square has been prepared by UV lithography to restrict the active area for electrochemical deposition. PPy was electrochemically synthesized under similar conditions as described in the previous sections. The Hall voltage was detected by two electrometers, and the current of 2 pA was supplied by Keithley 220 source. The magnetic field applied during the experiment was ±0.2 T. The mobility was evaluated from the Hall voltages to 7 cm²/Vs [47]. However, the Hall voltages were slowly drifting in time, and thus the mobility was obtained with high error bar of 20 cm²/Vs.

To reduce the effect of Hall voltage drift, we re-designed electrical connection of the PPy-diamond structure as shown in Figure 10a and also increased the magnetic field. The obtained diagonal Hall voltage under white light illumination (cold light source, 40 klx) is plotted as a function of time in Figure 10b. The Hall voltage is still varying in

![Figure 10](image-url)
time, and the difference between values for positive and negative magnetic field, which should correspond to the true Hall voltage, is decreasing. We assign this effect to temperature gradients (up to several kelvins) arising in the measuring chamber because of heat flow from the magnet coils. These gradients create thermal and thermonagnetic electromotoric forces (EMFs) [55] of the order of 1 mV modifying the measured voltage. In more conductive samples, these EMFs can be partially suppressed by alternating the direction of the measuring current. Unfortunately, in our case, the necessary long times of RC relaxation and integration (tens of minutes) prevent a successful application of this procedure. However, after leaving the magnet to cool down to room temperature, the repeated measurement shows a similar dependence, confirming in this way that we really observed the Hall voltage.

The same Hall voltages plotted as a function of magnetic field are shown in Figure 10c. If an arithmetic mean is performed for each applied magnetic field and these values are used for evaluation, then the resulting Hall mobility is $16 \pm 10 \text{cm}^2/\text{Vs}$. This revised value of mobility with much lower relative error supports the idea of charge transport via diamond and thus transfer of charge from PPy to diamond.

Conclusions

In this article, we showed that employing and combining advanced scanning probe techniques can provide significant insight into the correlation of structural, chemical, and opto-electronic properties of organic-based heterojunctions on the microscopic scale. We presented specific results obtained on fully organic bulk heterostructures and on heterojunction of diamond with electrochemically synthesized organic dye. Although these systems seem to be different, from the optoelectronic point of view, they behave in a similar way. In both systems, strongly bound excitons are created which dissociate at the interface with the other material. Correlation of advanced scanning probe techniques (AFM, KFM, and CS-AFM) with micro-Raman mapping identified the reasons for low PV efficiency and could resolve differences in optoelectronic quality and stability by contact-less monitoring of dynamic photo-response of the organic blends. We demonstrated the applicability of the above methodology also to a hybrid organic-inorganic system. By correlating AFM nanoshaving, KFM surface potentials, as well as $I(V)$ characteristics, we proved that during the electrochemical synthesis a covalent bond between diamond and PPy is formed. Such conclusion is difficult to make based on commonly applied techniques such as X-ray photoelectron spectroscopy because here we have all-carbon systems (polymer-diamond). Furthermore, based on microscopic KFM and SPV measurements, the model of charge transfer from PPy to diamond under illumination was proposed. The model was supported by in-plane $I(V)$ and Hall effect measurements.

Abbreviations

AFM: atomic force microscopy; CS-AFM: current-sensing AFM regime; EMFs: electromotoric forces; ITO: indium tin oxide; KFM: Kelvin force microscopy; PPy: polypyrrole; SPV: surface photovoltage.

Acknowledgements

The authors appreciate technical assistance of Z. Polášková, J. Potměšil, V. Jurka, K. Jurek, and J. Žemek. The authors also acknowledge the financial support of the Academy of Sciences of the Czech Republic, (project No. KAN40010701, and the Fellowship J.E. Purkině), the Ministry of Education, Youth and Sports of the Czech Republic (grants No. LC06040, LC510, 1M06031, AV0Z10100521), and the Grant Agency of the Czech Republic (grants No. 202/09/H041, P204/10/0212).

Author details

1Institute of Physics ASCR, v.v.i., Cukrovarnická 10, 16200 Prague 6, Czech Republic 2Institute of Macromolecular Chemistry ASCR, v.v.i., Heyrovského nám. 2, 16206 Prague 6, Czech Republic

Authors’ contributions

BR conceived the idea of PPy-diamond, designed, and coordinated the study, participated in scanning probe experiments and PPy synthesis, participated in the experimental setup development, participated in discussions and analysis, and prepared the manuscript. JC carried out scanning probe experiments, fabricated device structures, participated in the experimental setup development and discussions; helped in drafting the manuscript. AK provided diamond specimens and their surface terminations. ML carried out micro-Raman spectroscopy and participated in discussions. PH performed Hall measurements and data analysis. JM assisted with Hall measurement and discussions. AP prepared organic blends and measured $I(V)$ characteristics of the blends. VC participated in the design of the organic blends study and discussion; built up the experimental setup for preparation and macroscopic study of organic blends. AF conceived the scanning probe study of organic blends and the idea of comparing Hall mobility, participated in the experimental setup development. JK participated in discussions of electronic transport and overall results.

Competing interests

The authors declare that they have no competing interests.

Received: 1 October 2010 Accepted: 18 March 2011

References

1. Botto B, McNeill R, Weiss D: Electronic conduction in polymers - iii. Electronic properties of polypyrrole. Aust J Chem 1963, 16:1090-1103.
2. Chiang C, Finchir C9, Park Y, Heeger J, Shirakawa H, Louis E, Gau S, MacDiarmid A: Electrical conductivity in doped polyaacetylene. Phys Rev Lett 1977, 39:1098-1101.
3. Garnier F, Hjalauir R, Yassar A, Srivastava P: All-polymer field-effect transistor realized by printing techniques. Science 1994, 265:1684-1686.
4. Lebens S, Tourillon G, Garnier F: Influence of the doping on the photovoltaic properties of thin films of poly-3-methylthiophene. Thin Solid Films 1968, 139:221-231.
5. Karg S, Riess W, Dyakonov V, Schwoerer M: Electrical and optical characterization of poly(phenylene-vinylene) light emitting diodes. Synth Met 1995, 54:427-433.
6. Stubinger T, Brütting W: Exciton diffusion and optical interference in organic donor-acceptor photovoltaic cells. J Appl Phys 2001, 90:3632-3641.
7. Yu G, Gao J, Hummelen J, Wud F, Heeger A: Polymer photovoltaic cells: enhanced efficiencies via a network of internal donor-acceptor heterojunctions. Science 1995, 270:1789-1791.
8. Liang Y, Xu Z, Xia J, Tsai ST, Wu Y, Li G, Ray C, Yu L: For the bright future - bulk heterojunction polymer solar cells with power conversion efficiency of 7.4%. Adv Mater 2010, 22:E135-E138.
9. O’Regan B, Gratzel M. A low-cost, high-efficiency solar cell based on dye-sensitized colloidal ti20 films. Nature 1991, 353:737.
10. Ouyang T, Loth K, Qi D, Wee A, Nieslédka M. Chemical bonding of fullerene and fluorinated fullerene on bare and hydrogenated diamond. Chem Phys Chem 2008, 9:1286-1293.
11. Strobel P, Riedel M, Risten J, Ley L. Surface transfer doping of diamond. Nature 2004, 430:439-441.
12. Remelz Z, Uzán-Saguy C, Baskin E, Kalish R, Avigal Y, Nieslédka M, Koizumi S. Photo-hall effect measurements in p, n and b-doped diamond at low temperatures. Diam Relat Mater 2004, 13:713-717.
13. Riedel M, Risten J, Ley L. Recovery of surface conductivity of h-terminated diamond after thermal annealing in vacuum. Phys Rev B 2004, 69:125338.
14. Takeuchi D, Kato H, R G, Yamada T, Vinod P, Hwang D, Nebel C, Okushi H, Yamazaki S. Direct observation of negative electron affinity in hydrogen-terminated diamond surfaces. Appl Phys Lett 2005, 15:52103.
15. Nebel CE, Rezek B, Shin D, Waranabe H, Yamamoto T. Electronic properties of h-terminated diamond in electrolyte solutions. J Appl Phys 2006, 99:033711.
16. Rezek B, Nebel C. Kelvin force microscopy on diamond surfaces and devices. Diam Relat Mater 2005, 14:466-469.
17. Rezek B, Watanabe H, Shin D, Yamamoto T, Nebel C. Ion-sensitive field effect transistor on hydrogenated diamond. Diam Relat Mater 2006, 15:673-677.
18. Hammers Rl, Butler JE, Lasseret T, Nichols BM, Russell JN, Tse KY, Yang W. Molecular and biomolecular monolayers on diamond as an interface to biology. Diam Relat Mater 2005, 14:661-668.
19. Hartí A, Schmirch G, Gardo J, Hernando J, Catharino SC, Walter S, Feulner P, Kromka A, Steinmüller D, Stutzmann M. Protein-modified nanocrystalline silicon thin films for biosensor applications. Nat Mater 2004, 3:758-742.
20. Rezek B, Shin D, Nebel C. Properties of hybridized dna arrays on single-crystalline undoped and boron-doped (100) diamonds studied by atomic force microscopy in electrolytes. Langmuir 2007, 23:7626-7633.
21. Kanazawa K, Diaz A, Gill W, Granti P, Street G, Gardini G, Cvek J. Polyppyrole: an electrochemically synthesized conducting organic polymer. Synth Met 1980, 1:329-336.
22. Mohammadi A, Hasan M, Liebdberg B, Lundström I, Salarék W. Chemical vapour deposition (cvd) of conducting polymers: polyppyrole. Synth Met 1986, 14:189-197.
23. Kaïsioriayula S, Qi X, Vetter C, Croes K, Pavlacky D. A structural and morphological comparative study between chemically chemisitised and polymerizedpolyppyrole. J Coat Technol Res 2010, 7:145-158.
24. Nakata M, Kise H. Preparation of polyppyrole-poly(vinyl chloride) composite films by interphase oxidative polymerization. Polym J 1993, 25:91-94.
25. Nakata M, Shiraishi Y, Taga M, Kise H. Synthesis of electrically conductive polyppyrole films by interphase oxidative polymerization. Makromolklules Chem - Macromol Chem Phys 1992, 193:765-771.
26. Hong L, Li Y, Yang M. Fabrication and ammonia gas sensing of palladium/polyppyrole nanocomposite. Sens Actuators B: Chem 2014, 205:25-31.
27. Yang X, Li L, Yan F. Polyppyrole/silver composite nanotubes for gas sensors. Sens Actuators B: Chem 2010, 145:495-500.
28. Sun A, Li Z, Wei T, Li Y, Cui F. Highly sensitive humidity sensor at low humidity based on the quaternized polyppyrole composite film. Sens Actuators B: Chem 2009, 142:197-203.
29. Guiseppe-Elie A. Electroconductive hydrogels: synthesis, characterization and biomedical applications. Biomaterials 2010, 31:2701-2716.
30. Singh M, Kathuria P, Jampana H. Polyppyrole based amperometric glucose biosensors. Sens Actuators B: Chem 2009, 143:430-443.
31. Rajapakse R, Murakami K, Bandara H, Rajapakse R, Velauthamurti K, Wijeratne S. Preparation and characterization of electroconducitively conducting polyppyrole-montmorillonite nanocomposite and its potential application as a cathode material for oxygen reduction. Electrochim Acta 2010, 55:2490-2497.
32. Yuan Y, Zhou S, Zhuang L. Polyppyrole-carbon black composite as a novel oxygen reduction catalyst microbial for fuel cells. J Power Sources 2010, 195:3490-3493.
33. Selvaraj M, Palraj S, Maruthan K, Rajagopal G, Venkatachari G. Synthesis and characterization of polyppyrole composites for corrosion protection of steel. J Appl Polym Sci 2010, 116:1524-1537.
34. Wang G, Yang L, Qu Q, Wang B, Wu Y, Holze R. An aqueous rechargeable lithium battery based on doping and intercalation mechanisms. J Solid State Electrochem 2010, 14:865-869.
35. Mihryana Y, Ahoyom L, Bennet A, Stromme M. Novel high specific surface area conducting paper material composed of polypyrrole and cladophora cellulose. J Phys Chem B 2008, 112:12249-12255.
36. Binning G, Quate C. Atomic force microscope. Phys Rev Lett 1986, 56:930-933.
37. Rezek B, Stuchlík J, Fejfar A, Kočka J. Microcrystalline silicon thin films studied by atomic force microscopy with electrical current detection. J Appl Phys 2002, 92:587-593.
38. Sugimoto Y, Abe M, Jelinek P, Pérez R, Morita S, Custance O. Chemical identification of individual surface atoms by atomic force microscopy. Nature 2007, 446:64-67.
39. Ledinsky M, Vetushka A, Stuchlík J, Fejfar A, Kočka J. Raman mapping of micro-crystalline silicon thin films with high spatial resolution. Phys Status Solidi C 2010, 7:704-707.
40. Yeo BS, Sadler J, Schmid T, Zenoib R, Zhang W. Tip-enhanced raman spectroscopy - its status, challenges and future directions. Chem Phys Lett 2009, 472:1-3.
41. Wang R, Wang J, Hao F, Zhang M, Tian Q. Tip-enhanced raman spectroscopy with silver-coated optical fiber probe in reflection mode for investigating multivariant carbon nanotubes. Opt Exp 2010, 19:1845-1848.
42. Čermák J, Rezek B, Cimrová V, Viprachtický D, Ledinsky M, Mates T, Fejfar A, Kočka J. Correlation of atomic force microscopy detecting local conductivity and micro-raman spectroscopy on polymer-fullerene composite films. Phys Status Solidi RRL 2007, 1:193-195.
43. Čermák J, Rezek B, Cimrová V, Fejfar A, Purna K, Varábková M, Kočka J. Time-resolved opto-electronic properties of poly(3-hexylthiophene-2,5-diyll-fullerene heterostructures detected by Kelvin force microscopy. Thin Solid Films 2010, 519:836-840.
44. Čermák J, Rezek B, Cimrová A, Ledinsky M, Kočka J. Electrochemical synthesis and electronic properties of polypyrrole on intrinsic diamond. Diam Relat Mater 2009, 18:1096-1101.
45. Rezek B, Cermák J, Kromka A, Ledinsky M, Kočka J. Photovoltage effects in polypyrrole-diamond nanosystem. Diam Relat Mater 2009, 18:249-252.
46. Čermák J, Kromka A, Ledinsky M, Rezek B. Illumination-induced charge transfer in polypyrrole-diamond nanosystem. Diam Relat Mater 2009, 18:800-803.
47. Čermák J, Rezek B, Hubák P, Mareš J, Kromka A, Fejfar A. Photo-conductivity and hall mobility of holes at polypyrrole-diamond interface. Diam Relat Mater 2010, 19:174-177.
48. Lo KL, Lee MC. Raman scattering of thin film c60/c70 fullerenes. Chin J Phys 1993, 31:653.
49. Maturukov K, Kemenrik M, Wienk M, Charter D, Janssen R. Scanning kelvin probe microscopy on bulk heterojunction polymer blends. Adv Funct Mater 2009, 19:1570-1573.
50. Chesa M, Birgù L, Kim J, Schlerk R, Friend R, Siringhaus H. Correlation between surface photovoltage and blend morphology in polylfluorocarbon-based photodiodes. Nano Lett 2005, 5:559-563.
51. Manduc M, de Boer B, Piausch G, Blom P. Trap-limited electron transport in dispersed semiconductor polymers. Phys Rev B 2007, 75:192302.
52. Rezek B, Shin D, Nakamura T, Nebel CE. Geometric properties of covalently bonded DNA on single-crystalline diamond. J Am Chem Soc 2006, 128:3884.
53. Rezek B, Shin D, Uetsuka H, Nebel CE. Microscopic diagnostics of DNA molecules on mono-crystalline diamond. Phys Status Solidi A 2007, 204:2888.
54. Kaminski W, Rozsíval V, Jelínek P. Theoretical study of electronic and transport properties of ppy-pt(111) and ppy-c(111)h interfaces. J Phys Condensed Matter 2010, 22:045003.
55. Popovic RS. Hall Effect Devices Bristol: Adam Hilger, 1991.