Revealing nanoscale optical properties and morphology in perfluoropentacene films by confocal and tip-enhanced near-field optical microscopy and spectroscopy

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Combining high resolution optical microscopy and spectroscopy, we propose a novel, generally applicable and highly sensitive method for determining the local morphology in organic semiconductor thin films (e.g. perfluoropentacene (PFP)). An azimuthally or radially polarized doughnut mode (APDM or RPDM) laser beam is focused by a high numerical aperture parabolic-mirror to excite a diffraction limited volume of the PFP film with an electric field polarized either exclusively in-plane or dominantly out-of-plane (relative to the substrate). We find two distinct morphologies of thin PFP films: molecular aggregates and crystalline terraces. The well-defined dipole emission patterns observed from the molecular aggregates strongly suggest the presence of localized excitations. For both laser modes, we observe that for the PFP aggregates, the photoluminescence (PL) emission from the main electronic transition is blue-shifted by about 10 meV, as compared to that from the molecular terraces. For the C–C bending modes, the Bg at 1581 cm−1 (ν3) and the Ag at 1316 cm−1 (ν0), we observe a decrease of the intensity ratio (I1/I0) from 0.6 (terrace) to 0.15 (aggregate). Furthermore, the intensity ratios (IAPDM/I RPDM) of ν1 excited by different polarizations increase from 0.12 (terrace) to 0.73 (aggregate). These results indicate that the PFP molecules orient rather parallel to the substrate in the aggregates, whilst more upright in the terraces. Benefiting from the nanometer scale optical resolution offered by the tip-enhanced near-field optical method, we observe clear optical contrasts between the molecular aggregate and the terrace as well as individual layers within a terrace. Tip-enhanced optical spectra locally taken from the molecular terrace and the aggregate show similar blue-shift of the main PL peak and change in the Raman intensity with different polarizations as from the far-field assembly-measurements, which further confirms the different molecular orientations in these two morphologies.

Introduction

Organic semiconductor thin films have attracted increasing attention for applications in flexible electronic devices.1 An important point is the chemical tunability by attaching or replacing certain functional groups. Fluorination is an important strategy, e.g., to make the compounds more stable against oxidation, to shift the relevant energy levels, or to create donor–acceptor pairs with the hydrogenated parent compound.2–7 Perfluoropentacene (PFP, C22F14), a derivative of the prototype system pentacene (PEN, C22H14) with exchange of the hydrogen by fluorine, is an obvious choice of study in this context.2,8–11 Compared with PEN which is a p-type semiconductor, PFP behaves as an n-type material due to the strong electronegativity of fluorine, providing the opportunity for bipolar device applications. For achieving high device performance, the fundamental properties, such as local morphology, orientation and intermolecular coupling of molecules in the films are of great interest.12–19

Although the crystal structure and the morphology of pure PFP thin films on various substrates can be investigated by X-ray diffraction (XRD) and atomic force microscopy (AFM),3,4,20 as well as other microscopic techniques,21 optical methods such as absorption,22–24 photoluminescence (PL)25 and infrared26 spectroscopy provide additional important information on the molecular energy levels and interaction in the films. Polarization-dependent micro-Raman spectroscopy provides valuable information of the...
molecular vibrations, and their modifications due to the local environment. For PEN, polarized Raman spectroscopy has been used to investigate the molecular orientation and the crystal phase of the films.\textsuperscript{27–29} Compared to linearly polarized laser beams, cylindrical vector beams, such as radially or azimuthally polarized doughnut modes (RPDM or APDM) provide distinct electric field distributions in the laser focus: a dominant out-of-plane electric field can be achieved with the RPDM whilst an exclusive in-plane electric field is produced with the APDM. Based on the polarization-dependent PL or Raman intensities, accurate and quantitative information about the molecular orientation can be obtained with diffraction-limited spatial resolution of typically half of the excitation laser wavelength.\textsuperscript{28,30}

To further correlate details of the nanoscale morphology with optical properties, an optical resolution below the diffraction-limit is required which can be achieved by tip-enhanced near-field optical microscopy and spectroscopy.\textsuperscript{31–33} This technique provides images with an optical resolution down to several nanometers\textsuperscript{33–36} and sensitivities allowing for the detection of single molecules and single surface states.\textsuperscript{37–40}

In this paper, we use a home-built parabolic-mirror assisted confocal and near-field optical microscope\textsuperscript{41–44} to study the optical properties and morphology of PFP films by PL and Raman microscopy and spectroscopy with high spatial resolution. We reveal the molecular orientations in different morphologies by analysing the intensity of different Raman vibrational modes with two distinct laser excitation polarizations.

Results and discussion

The topography (Fig. 1(a) and (b)) of the PFP film shows that the film mostly forms ordered molecular terraces. The topographic line profile (Fig. 1(c)) reveals 1.5 nm high steps corresponding to the length of an upright-oriented single PFP molecule.\textsuperscript{3} We also observe large PFP ‘aggregates’, by which we mean large, anisotropic features. The aggregates are typically about several tens of nanometers high and hundreds of nanometers long, as marked by the circle in Fig. 1(b). As the orientation and stacking of molecules can be revealed from their Raman and PL spectra, we study the nature of these two distinct morphologies by polarization-dependent confocal and near-field optical microscopy and spectroscopy.

We first perform topography-correlated confocal PL and Raman imaging and spectroscopy from molecular terraces and aggregates in the PFP film. Fig. 2(a) and (b) show confocal PL images (10 \textmu m \times 10 \textmu m) of a PFP film excited with RPDM and APDM, respectively. In the focal volume of these polarization modes the PFP molecules are excited differently: mainly perpendicular to the substrate with RPDM and purely parallel to the substrate with APDM. In Fig. 2(a) and (b) we observe localized and diffraction-limited excitation patterns which have intensities which exceed the average PL by one order of magnitude. In general the PL excitation rate is proportional to |\nabla \cdot \mathbf{E}^2|, where \mathbf{E} is the excitation electric field and \mathbf{D} is the transition dipole moment. For the image obtained with RPDM we observe a bright and almost circular Airy disc pattern from the structure marked by the white arrow in Fig. 2(a). This pattern reflects the projection of the transition dipole moment parallel to the surface normal of the sample.\textsuperscript{45} When RPDM is switched to APDM, the pattern changes to two parallel lobes as marked by the white arrow in Fig. 2(b). In this case, the pattern reflects the direction of the transition dipole moment projected into the sample plane which orients along the two lobes as indicated by the white arrows (on both ends). Hence for a quasi-point-light source which could originate in the present case from structural defects or local excitations, we observe distinct coffee-bean shaped patterns which allow us to determine the orientation of a transition dipole moment as in the case of an isolated molecule.\textsuperscript{46,47}

Comparing the optical and the topography image (Fig. 2(c)), which are taken from the white dashed squares marked in Fig. 2(a) and (b), we clearly see that the localized excitations only occur at the aggregates and not at the terraces. In the topography images (Fig. 2(c)), both PFP aggregates and molecular terraces are observed. Fig. 2(d) shows the normalized optical intensities (upper panel, blue dots for RPDM, red dots for APDM) as well as the calculated electric field intensity distributions (E\textsuperscript{2}) (upper panel, solid lines) and the corresponding topography (lower panel) line profiles of a PFP aggregate, which is about 80 nm high, 160 nm wide and 800 nm long. Due to the far-field confocal optical resolution and the weak optical contrast, the PFP molecular terraces are not clearly resolved in Fig. 2(a) and (b).

To further understand the polarization-dependent PL and Raman spectra and their relations to the local morphologies of the PFP film, spectra were recorded from different terraces and aggregates (over the 10 \textmu m \times 10 \textmu m area in Fig. 2) with the two polarizations. Fig. 3(a) illustrates a stack of all spectra recorded

Fig. 1 (a and b) Topographic images recorded by raster scanning a sharp gold tip of the near-field optical microscope over the surface of a PFP film. (c) A respective topographic line profile that is taken from the region indicated by the line in (a). Single PFP molecular steps of about 1.5 nm are clearly visible.
with RPDM from the terraces to aggregates, showing increasing
intensities from the bottom to the top of the picture. Three
representative spectra for the high, moderate and low intensities
are shown in Fig. 3(b). All the spectra consist of sharp Raman peaks
and broad PL backgrounds. Clearly, the PL background dominates
the optical intensity that we observed in Fig. 2(a) and (b). As shown
in Fig. 3(b), the PL background of these spectra can be fitted by
three Gaussian curves: one main electronic transition ($S_1 \rightarrow S_0$) in
the range from 1.748 eV to 1.736 eV and two weak spectral shoulders
at 1.635 eV. The spectral shoulder at 1.635 eV has been
assigned to a vibronic band of the $S_1 \rightarrow S_0$ transition. The dashed
lines in Fig. 3(c) indicate the peak positions and show a blue-shift
from the terrace to the aggregate.

Although the Raman and PL peaks from the PFP aggregates
are generally stronger than from the molecular terraces as
shown in Fig. 3(a), we find that the relative intensity ratios
for certain Raman peaks vary significantly with the locations.
To provide a clear comparison, we plot in Fig. 3(c) a series of
the normalized Raman peaks of the spectra shown in Fig. 3(a) after
the subtraction of the PL background. Red indicates the highest
intensity; whilst blue the lowest. Three typical spectra are shown
in Fig. 3(d) with an offset for clarity. The well-defined Raman
peaks are in good agreement with the published results$^{48}$ within
the spectral resolution. The intense Raman peaks observed in
the range from 1200 cm$^{-1}$ to 1600 cm$^{-1}$ are mainly C–C stretching
modes of the conjugated ring.$^{48}$ The most prominent peak labelled
as $\nu_1$ is located at 1316 cm$^{-1}$. The peak located at 1581 cm$^{-1}$ is
labelled as $\nu_2$. For the RPDM, the Raman peak $\nu_1$ from the PFP
terrace is more intense than that from the aggregate.

Similarly, spectra recorded with the APDM from terraces and
aggregates in the 10 $\mu$m $\times$ 10 $\mu$m area are shown as a series in
the sequence from low intensity to high intensity in Fig. 4(a). Compared
with the RPDM, in Fig. 4(b) the three representative spectra taken from the bright
($B_3$) moderate ($R_2$) and dark ($R_1$) areas by the excitation with RPDM. The Raman
peaks at about 520 cm$^{-1}$ and 950 cm$^{-1}$ are from the Si substrate. The PL
background are fitted using Gaussian function and the maxima of the fitted
spectra are indicated by the dashed lines. (c) A series of Raman spectra after the
subtraction of PL background in (a). The Raman spectra are normalized to the
1316 cm$^{-1}$ peak. (d) Three representative Raman spectra recorded with RPDM.
and its Raman tensor components are \( x^2, y^2 \) and \( z^2 \); however \( \nu_1 \) has a \( B_{3g} \) representation and its Raman tensor component is \( yz \). Therefore, the intensity of \( \nu_1 \) is sensitive to the molecular orientations. It is known that the PFP molecule in the terrace structures (on SiO\(_2\)/Si substrate) orients with the long axis almost perpendicular to the substrate surface. Since the dominant electric field in the focus of the RPDM is parallel to the substrate normal, an increase in the tilt angle of the PFP molecule with respect to the substrate normal leads to a weaker excitation, decreasing the intensity of the Raman mode. From the change of the intensity ratio of \( \nu_1 \) to \( \nu_2 \) modes, the orientation of the molecules with respect to the substrate normal can be deduced.

To obtain a comprehensive description of the molecular orientation, we analyse all spectra recorded from the sample. Fig. 5(a) shows the intensity ratios of the \( B_{3g} \) Raman mode \( \nu_1 \) against the \( A_1 \) mode \( \nu_0 \) \( (I_{\nu_1}/I_{\nu_0}) \) at different locations as recorded with RPDM and APDM. In these plots, the different locations are represented by their total PL intensities. In general, high PL intensities correspond to PFP aggregates while low PL intensities correspond to the molecular terraces. We differentiate the experimental results from the terraces and aggregates by red dashed lines in Fig. 5(a) and (b), according to the topography and the corresponding PL intensity. Due to the exclusive in-plane excitation with APDM, the \( \nu_1 \) peaks are weak and the \( (I_{\nu_1}/I_{\nu_2}) \) ratio is about 0.15 for all the locations (Fig. 5(a) right). For RPDM, the relative intensities of \( \nu_1 \) peaks vary systematically with the locations: for PFP aggregates, the \( (I_{\nu_1}/I_{\nu_2}) \) ratio is about 0.15.

For molecular terraces the PL intensity is generally lower than that from the aggregate, however, the \( (I_{\nu_1}/I_{\nu_2}) \) ratio of the Raman peak increases (up to about 0.6) with the decreasing PL intensity. The changing in the \( (I_{\nu_1}/I_{\nu_2}) \) ratio indicates that PFP molecule in the aggregate orients differently from it in the molecular terraces. A high \( (I_{\nu_1}/I_{\nu_2}) \) ratio indicates a small molecular tilting angle with respect to the surface normal. This can be further confirmed by comparing the intensity of the \( \nu_1 \) Raman peak excited with RPDM and APDM, respectively. Fig. 5(b) shows the intensity ratio \( (I_{\text{APDM}}/I_{\text{RPDM}}) \) of the Raman mode \( \nu_1 \) recorded from the terraces and aggregates. The small ratio indicates a small tilting angle which refers to the terraces; while the large ratio indicates a large tilting angle which refers to the aggregates. To demonstrate the above conclusion quantitatively, the following calculations are performed.

The Raman intensity of a certain mode is expressed by the Raman polarizability tensor \( (x_0yz) \), the electric field vector of the excitation \( (E_0) \) and scattered radiation \( (E_\alpha) \) as:

\[
I \propto |E_\alpha|^2 x_0yz E_0|^2
\]

With the expression of the Raman tensor as:

\[
\begin{bmatrix}
0 & 0 & 0 \\
0 & f & 0 \\
0 & 0 & f
\end{bmatrix}
\]

the excitation intensity \( I_x, I_y, I_z \) for the radial and azimuthal modes and the detection parameters \( [A, B] \) over the scattering cone, the Raman intensity of \( B_{3g} \) mode of PFP molecule which has a tilt angle \( \theta \) and twist angle \( \phi \) with respect to the substrate normal (Fig. 5(c)) can be calculated as in eqn (1):

\[
I_{B_{3g}} = f^2 \left[ A \sin^2 \theta (4 \cos^2 \theta \sin^2 \phi + \cos^2 \phi) + B \sin^2 \phi (2 \cos^2 \theta - \sin^2 \theta)^2 \right] I_x + f \left[ A \sin^2 \theta \cos^2 \phi + B \cos^2 \theta \cos^2 \phi \right] I_y + \left[ A \sin^2 \phi (\cos^2 \theta - \sin^2 \theta)^2 + \cos^2 \theta \cos^2 \phi \right] I_z + 4B \sin^2 \phi \cos^2 \theta \sin^2 \phi I_z
\]

(1)

In the calculation for the Raman intensity ratio (azimuthal against radial), the parameters are chosen as follows: \( A = 12.54 \), \( B = 1.89 \) for our experimental setup with an open angle of 86 degree, \( I_x = 6\% \), \( I_y = 88\% \) for the excitation intensity with RPDM; \( I_x = I_y = 50\% \), \( I_z = 0 \) for the excitation intensity with APDM. The calculated results are shown in Fig. 5(d). At small tilt angles (molecules standing upright) the ratio is much smaller, which is consistent with our experimental observations (about 0.12) at the PFP terrace. For PFP aggregates, the ratio is experimentally observed to be about 0.73, which agrees with the calculated intensity ratio \( I_x/I_y \) at larger tilting angles and a twisting angle in the range of 40 to 50 degree.
Besides the Raman intensity ratio changes, we observe also shifts of the main PL peak for different morphologies. To provide a complete description of the behaviour, Fig. 5(e) shows the main PL emission peak, which is obtained from the spectral fitting, against the intensity at different locations in the PFP film excited with both laser modes, respectively. The peak positions and spectral widths of the other two Gaussian curves for fitting the spectral shoulders were fixed during the fitting for all different locations. For the locations where the PL intensity increases from about 60k counts to 500k counts, we observe that the PL peak position shifts by about 10 meV to higher photon energy for both laser modes. Therefore, in general the PL emission from the PFP aggregates has higher photon energy than that from the PFP terrace. This blue-shift cannot be attributed to the film thickness-dependent HOMO–LUMO transition energies of the PFP film, where a red-shift of the absorption spectrum with increasing thickness has been reported. Other possibilities may be considered, for example that the change in the molecular orientation at the aggregates and the terraces induces different dipole moments coupling schemes, either a H- or J-aggregate like transition or PL emission from strained unit cells induced by the different local polymorph.

Confocal spectroscopy only allowed us to investigate how PL- and Raman-spectra vary at different locations with diffraction-limited optical resolution. To optically resolve molecular terraces and the details of the aggregates, we employ near-field optical microscopy and spectroscopy to push the optical resolution down to the nanometer regime. Hence, for all near-field optical measurements a sharp gold-tip is centered in the focal volume of a RPDM for an efficient plasmon excitation. Fig. 6(a) and (b) show the simultaneously recorded topography and near-field optical images of an area with a PFP aggregate. In the topographic image (Fig. 6(a)), we observe that the PFP aggregate has a width of about 100 nm and a height of about 120 nm. The corresponding near-field optical image (Fig. 6(b)) shows intense optical signals. According to the discussion based on the confocal Raman and PL spectra, the PFP molecules orient more parallel to the substrate plane in aggregates than those in the PFP terraces. As the main transition dipole moment lies along the short molecular axis, for PFP aggregates one would find a substantial component in the direction perpendicular to the substrate plane, which makes an efficient coupling to the tip plasmon and hence a bright near-field excited PL-signal possible.

Fig. 6(d) and (e) show the topography and the simultaneously recorded near-field optical images of an area where PFP terraces with single molecular steps are clearly resolved. The near-field optical images show optical contrast from different layers of the molecules. To demonstrate the correlation quantitatively, Fig. 6(f) shows the topographic height and optical intensity profile along one
line, the position of which is indicated in Fig. 6(d) and (e). Along this line the corresponding optical intensity increases from about 60 counts to 115 counts on average (130 counts at the maximum), while the height of the PFP terrace increases by only 5 nm.

The near-field optical spectra from the PFP aggregate and the molecular terraces are shown in Fig. 7. The near-field spectrum consists of sharp Raman peaks, a PL background from the molecules and also a contribution from the gold tip. In Fig. 7 the spectra from the gold tips alone are shown as grey lines. It should be noted that two different tips are used for the measurements. The PL spectra of the gold tips are fitted by single Lorentzian curves as indicated by the red lines in Fig. 7. Similar to the analysis for the confocal spectra, the PL background of the PFP molecules is fitted by three Gaussian curves (blue lines). After the subtraction of PL background, Raman peaks from the PFP aggregate and PFP molecular terrace are shown in Fig. 7(c) and (d), respectively. Again, we analyse the intensity ratio of Raman modes $I_n/I_0$ and the main PL peak position. From the PFP molecular terrace to the PFP aggregate, the ratio of the Raman modes $I_n$ against $I_0$ decreases from 0.69 to 0.15, while the spectral maximum of the main PL peak shifts from 1.74 eV to 1.75 eV. These near-field experimental results agree
well with the tendency demonstrated in Fig. 5(a) however with a much higher topographical and optical resolutions. In Fig. 7(e), the interactions between a gold tip and the different locations in a PFP film, at the terrace and at the aggregate, are shown. For PFP molecules in the terraces, the main transition dipole moment is relatively parallel to the substrate plane leading to a low PL signal.\textsuperscript{22}

In the molecular aggregate, the transition dipole moment orients more perpendicular to the substrate plane. The localized plasmon oscillation at the tip apex as excited by a RPDM is directed along the tip axis leading to an optical near-field under the tip apex which is mainly oriented perpendicular to the substrate plane. Hence it couples less efficiently with the molecular transition dipole moments in the molecular terraces as compared to the aggregates. Therefore, even though the PL intensity contrast is clearly visible from several molecular layers in the terraces (Fig. 6(d)–(f)), a large contrast can be observed in the near-field excited PL signal between the aggregates and the terraces (Fig. 6(a)–(c)).

**Experimental**

For the excitation a He–Ne laser at $\lambda = 632.8$ nm is used. The PL and Raman signals from the focal volume are collected by a parabolic mirror (numerical aperture of 0.998 in air) and are directed through two narrow band filters (in total 12 optical density) for blocking the laser line. The size (full width at the half maximum) of the diffraction limited spot (radial mode) on the sample is about 280 nm for the excitation with the 632.8 nm laser. The optical signals can be detected either by an avalanche photodiode for pixel by pixel photon counting as the sample is scanned through the focal field of an APDM or RPDM. We find that the PL peak position of the main electronic transition in the terraces is blue-shifted by about 10 meV with respect to the aggregates. With the tip-enhanced near-field optical approach, we are able to push the optical resolution to sub-diffraction limit. Clear optical contrast from a single or a few PFP molecule layers is observed. Near-field spectra further confirm the blue-shift of the PL peak and the decrease of the intensity ratio of the Raman modes from the PFP molecular terraces to the large aggregates. Combined with cylindrical vector beams, our confocal and near-field PL and Raman microscopy and spectroscopy provide sensitive methods to study the morphology, the orientation and the intermolecular interaction in thin molecular films.

**Acknowledgements**

The authors thank Reinhard Scholz for the scientific discussion and Kai Braun for improving the shear-force scanning tip runner. Financial support by the Projektförderung für Nachwuchswissenschaftler/innen University of Tübingen and the DFG (grant ME1600/5-3, BR4869/1-1 and SCHR700/20-1) is gratefully acknowledged.

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