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Citation for published version (APA):

Document license:
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DOI:
10.1002/anie.201710487

Document status and date:
Published: 22/01/2018

Document Version:
Publisher’s PDF, also known as Version of Record (includes final page, issue and volume numbers)

Please check the document version of this publication:

• A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher's website.
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Temperature-Responsive Luminescent Solar Concentrators: Tuning Energy Transfer in a Liquid Crystalline Matrix

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Abstract: Temperature-responsive luminescent solar concentrators (LSCs) have been fabricated in which the Förster resonance energy transfer (FRET) between a donor–acceptor pair in a liquid crystalline solvent can be tuned. At room temperatures, the perylene bisimide (PBI) acceptor is aggregated and FRET is inactive; while after heating to a temperature above the isotropic phase of the liquid crystal solvent, the acceptor PBI completely dissolves and FRET is activated. This unusual temperature control over FRET was used to design a color-tunable LSC. The device has been shown to be highly stable towards consecutive heating and cooling cycles, making it an appealing device for harvesting otherwise unused solar energy.

Materials that can change color in response to external stimuli, which can be found both in nature[1,2] and artificial devices, are currently receiving much attention.[3–5] Alongside stimuli-responsive optics, a particularly heavily studied topic in contemporary chemistry is Förster resonance energy transfer (FRET), the through-space migration of energy from an excited donor to an acceptor molecule that subsequently emits this energy as light.[6,7] FRET plays an important role in energy harvesting systems, for instance in collecting panchromatic light in a narrow wavelength[8] or directing energy to specific reaction centers in the case of photosynthesis.[9] Artificial temperature-responsive FRET systems in supramolecular assemblies have been reported, in which aggregation of the donor and acceptor monomers takes place at lower temperatures resulting in FRET.[10–12] while at higher temperatures, emission is blue-shifted as the thermal disruption of the co-assemblies results in spatial separation of the donor/acceptor pair, causing emission from the donor to be observable.[11,13,14] However, considerable research into supramolecular FRET for energy-harvesting purposes is still needed[9] because controlling and directing light in stimuli-responsive optical materials and devices remains a challenge.

LSCs are translucent devices containing fluorescent dyes that absorb sunlight and re-emit this light at longer wavelengths.[15,16] A fraction of the emitted light is channelled by the high refractive index lightguide to the edges, on which thin photovoltaic (PV) cells are attached. FRET between a donor–acceptor fluorescent dye pair has been used to reduce one of the primary LSC loss mechanisms, reabsorption, if the acceptor molecules emit light outside the absorption band of the donor.[17]

In this work, we use a liquid crystal (LC) as host for a coumarin-derivative donor (Cou)/perylene bisimide acceptor (PBI) pair. By using an LC as solvent, the device performs differently from conventional solution-based supramolecular FRET systems. At low temperatures, directed emission from only the aligned donor is visible because of the aggregation of the acceptor dye in the LC solvent; while in the isotropic phase at higher temperatures, the acceptor is molecularly dissolved, generating a FRET pair with the embedded donor with a redshift in emission (see Scheme 1). This construction is interesting since it allows for simultaneous control over absorption and luminescence through a single trigger, temperature.

**Scheme 1.** Proposed interaction mechanism of the C12-PBI (red)/Cou (yellow) molecules in the LC E7. At lower temperatures at which the LC is in its nematic phase (top left), the C12-PBI is in the non-active aggregated state, so all visible emissions (large green arrows) derive from the donor dye Cou, which does not aggregate in the host LC. At higher temperatures (top right), C12-PBI is more soluble, acting as acceptor for the donor in a FRET interaction, resulting primarily in visible emission from the acceptor (large orange arrows). The two molecules used in this FRET pair are the perylene bisimide C12-PBI and the coumarin derivative Cou, respectively (bottom).
In order to study FRET in a liquid crystalline host, a green fluorescent dye (Cou) with favorable optical characteristics to facilitate FRET ($\lambda_{	ext{abs}} = 450 \text{ nm}$, $\lambda_{	ext{em}} = 530 \text{ nm}$) was used as a dichroic donor dye. Nematic LC E7, which is often used in electro-optical devices, was chosen as the liquid crystalline host solvent (with a nematic-isotropic transition at 58°C, see Figures S1 and S2 in the Supporting Information for details). C12-PBI was synthesized as dichroic acceptor; because it is a tetraphenoxy-substituted perylene bisimide dye, it exhibits outstanding absorption and efficient emission ($\lambda_{\text{max}} = 573 \text{ nm}$, $\lambda_{\text{em}} = 606 \text{ nm}$; Supporting Information, Figure S3). The dodecyl chains utilized herein impart a high degree of molecular order in the LC matrix.

By placing the resulting solution in an LC alignment cell, a device is obtained that fluoresces bright green if left at room temperature (RT) for a few hours, displaying only emission from the donor Cou, indicating that C12-PBI luminescence has been deactivated; there is only a weak emission between 600 and 700 nm (see Figure 1). Remarkably, almost no absorbance by C12-PBI is measured at RT owing to dye aggregation. Upon heating, the green emission (from both the surface and edges of the cell) is gradually replaced by a red emission (see Figures 1 and the Supporting Information, Figure S4 and Video 1). The appearance of the absorption spectrum suggests that C12-PBI dissolves into the LC matrix. This most likely results in increased energy transfer from the Cou to C12-PBI, leading to the red emission.

To verify that energy transfer takes place at higher temperatures, fluorescence lifetime measurements were carried out. Time-correlated single-photon counting confirms that energy is transferred from Cou to the C12-PBI dye molecule. The fluorescence lifetime, $\tau_F$, of Cou in the hot, “red” state decreases by 29% (to 2.4 ns) compared to the “green” RT state. Concomitantly, when monitoring acceptor C12-PBI at elevated temperature, a delay of about 2.5 ns is seen before maximum counts are reached, strongly suggesting that energy is transferred to this molecule (see the Supporting Information, Figure S5 and Table S2 for further details). The photoluminescence excitation spectrum ($\lambda_{\text{em}} = 700 \text{ nm}$) supports this conclusion, as acceptor C12-PBI emission is detected at wavelengths in which normally only Cou is strongly excited (see the Supporting Information, Figure S6).

From comparing the absorbance and excitation spectra, we estimate a FRET efficiency of 54%.

To unravel the mechanism responsible for the disappearance of the absorption and emission of C12-PBI at low temperatures, the optical properties of 0.25 wt% of the dye in nematic LC host E7 (C12-PBI/LC) as a function of temperature were investigated. As in the donor/acceptor blend system, the C12-PBI/LC mixture displays a weak and rather featureless absorption band, accompanied by very weak, redshifted emission from the edge of the device at RT; see Figure 2. This featureless absorption band, together with the redshifted edge emission (likely a result of the absorption band broadening), is indicative of H-type aggregates (see the Supporting Information, Figure S7). Leaving the C12-PBI/LC mixture at RT for 2 to 3 h leads to the formation of amorphous aggregates, as confirmed by polarized optical microscopy. After heating to the isotropic phase of E7 at 58°C, the aggregates quickly dissolve and the cell adopts a more defined and stronger absorption typical of a molecularly dissolved state, and a strong red fluorescence is readily visible in minutes, reaching a maximum in about 15 min (see the Supporting Information, Figure S8). Therefore, both the transparency and luminescence of the C12-PBI/LC device are...
temperature-dependent. Concentration-dependent measurements show the absorbance and fluorescence increase always takes place around 58°C, independent of the dye concentration (tested from 0.12 up to 0.5 wt %).

This remarkable behavior is different from common supramolecular systems in which the degree of aggregation depends on the concentration and reveals that the aggregation of C12-PBI is determined by the phase transition of the LC solvent. In the nematic LC state, E7 is a poor solvent for the dye, while in the isotropic state, it is a good solvent. The thermal stability of C12-PBI in the LC host and its ability to reversibly form emissive and non-emissive states was tested by placing the cell in an oven in which the temperature was cycled between 30°C and 75°C, each cycle taking 7 h to complete. After 44 consecutive cycles, the measured absorbance of the samples only changed by circa 10 %, indicating the system was quite robust. The integrated power output at the edge parallel to the alignment direction decreased by 8 % (see the Supporting Information, Figure S9).

From edge emission measurements, it was found that the two-dye mixture in a 5 × 5 cm² 20 μm cell gap LSC exposed to light from a solar simulator generates a total light power output of 25 mW (optical efficiency ηmax = 2.4 %) in the cold state and 34 mW (ηmax = 3.2 %) when warmed. Furthermore, as E7 is a positive dielectric, applying a voltage across the cell allows for electrical switching of the host LC between a planar (light absorbing) and homeotropic (light transmissive) orientations. This leads to a dual-responsive device, as temperature can be used to control the absorption and emissive colors, while the applied voltage affects the overall absorbance by the incorporated dichroic dyes (see Figure 3a and the Supporting Information, Figure S10).

Applied as a window, the C12-PBI dye alone could allow for a temperature-sensitive aperture. Adding a second dye such as Cou offers the advantages of tuning both absorption and emission colors and providing extensive aesthetic potential to the presented device, including local coloration of the window by selective area heating (with a soldering iron, for instance; see Figure 3b). The coloration effects can be expanded by adding different secondary dyes, yielding unique optical effects. As examples, we have combined the novel C12-PBI dye with an orange emitter, a blue UV dye, and a NIR-emitting pigment (see Figure 3 and the Supporting Information, Figures S11, S12, and S13). These dyes do not show liquid crystal-induced aggregation and the UV/Vis absorbance and edge emission spectra can be found in the Supporting Information (Figure S13).

To conclude, we report on a new mechanism for controlling and directing light in an optical material. A thermochromic system has been fabricated in which the overall absorbance and fluorescence as well as the degree of FRET can be tuned and controlled by a liquid crystalline host. The use of an LC in this system allows for additional benefits over conventional organic solvents, in that we not only obtain specific solubility and alignment of the fluorescent dye molecule but can also extend the device response to a dual-trigger system, utilizing either thermal and/or electric stimuli, affecting both the absorbance and emission spectra of the device. The color change is reversible with minimal loss in fluorescence. Such temperature responsive optical materials and devices will find applications as aesthetically appealing smart devices for harvesting otherwise unused solar energy, be it in the form of electricity-generating responsive shading materials for greenhouses or a photocatalytic microreactor, sending photons to designated reaction centers.

**Experimental Section**

The synthesis of the novel perylene dye N,N’-Bis(4-dodecylphenyl)-1,6,7,12-tetra(phenoxy)perylene-3,4,9,10-tetraacidobisimide (C12-PBI), is accomplished by imidization of 1,6,7,12-tetra(phenoxy)-3,4,9,10-tetraacidobisimide (C12-PBI) with 4-dodecylaniline. This procedure has been described in detail in the Supporting Information. C12-PBI was added at 0.25 wt% (ca. 2 × 10⁻³ m) to the host liquid crystal E7 (Merck KGaA). For the dual color systems, 0.25 wt% of C12-PBI and 0.24 wt% (ca. 8 × 10⁻³ m) of a coumarin derivative are both added to the same liquid crystal.

**Acknowledgements**

The authors would like to acknowledge the support of the TKI PPS Smart Materials for Greenhouses program for their financial support of this work. We are grateful to Stefan C. J. Meskers for his assistance with recording some photophysical characteristics of the described systems. The authors also wish to thank Gilles H. Timmermans for his suggestions on the manuscript.

**Conflict of interest**

The authors declare no conflict of interest.