

BODIPY-derivatives as solid state colour converters for visible light communications

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In this paper we study a family of solid-state, organic semiconductors for visible light communications. The star-shaped molecules have a boron-dipyrromethene (BODIPY) core with a range of side arm lengths which control the photophysical properties. The molecules emit red light with photoluminescence quantum yields ranging from 22-56 %. Thin films of the most promising BODIPY molecules were used as a red colour converter for visible light communications. The film enabled colour conversion with a modulation bandwidth of 73 MHz, which is 16 times higher than of a typical phosphor used in LED lighting systems. A data rate of 370 Mbit/s was demonstrated using On-Off keying modulation in a free space link with a distance of ~15 cm.

Organic semiconductors have been widely studied in the last few decades for optoelectronic devices such as transistors¹, lasers^{2,3,4} amplifiers⁵, solar cells^{6,7} explosive sensors⁸ and organic light emitting devices (OLEDs)⁹. They are broadband visible emitters whose emission can be tuned by changing the chemical structure, and can have high photoluminescent quantum yields (PLQY) of up to 90% in undiluted films^{5,9,10}. This combined with simple and low cost solution processing techniques make them attractive materials for optoelectronic devices.

An emerging application area for organic semiconductors is in the field of visible light communications (VLC). Increasing demand for wireless communications has driven research into improving data transmission concepts^{11,12,13,14,15,16,17,18,19-21,20,21}. The concept of VLC is to use solid state room lighting to transfer data. This idea can exploit lighting installations to their full potential by providing illumination and data communication simultaneously, in an efficient, safe and low cost method. White LED lighting commonly uses blue LEDs coated with a yellow-emitting phosphor which acts as a colour converter for some of the blue light. An important consideration is the excited state lifetime of the phosphor/colour converter. High data transmission rates require short excited state lifetimes. Phosphors typically have a microsecond lifetime which severely limits the data transmission rate, so there is a need for colour converters with much shorter lifetimes.

Organic semiconductors are attractive alternatives to using phosphors as colour converters for VLC because they have faster radiative lifetime and high PLQY. They can also be modified to emit at any visible wavelength, allowing tuneable colour converters. The potential of using conjugated polymers for VLC has recently been demonstrated, using the commercial polymer Super yellow (Merck) in solution. The experiment combined the yellow emission of the polymer with a blue LED to produce white light and a data rate of 1.68 Gbit/s over a distance of 3 cm²². Improved colour rendering or wavelength division multiplexing requires fast red colour converters. We have recently shown that boron-dipyrromethene (BODIPY) molecules in solution can give a saturated red emission and a bandwidth of 39 MHz and data rates of ~ 98 Mbit/s^{23,24}. However, for practical applications, a compact and solid state structure is required. In this paper we study a family of star-shaped BODIPY molecules in the solid state, and demonstrate that they have an increased bandwidth allowing a data transfer rate of 370 Mb/s at a distance of 15 cm with simple on-off keying.

The molecules (shown in figure 1, top) have a star shape with boron-dipyrromethene as the core (BODIPY) and oligofluorene arms in a ‘Y’ shaped arrangement. The size of the molecule is modified by increasing the number of fluorene units in each arm from 1 to 4, and the corresponding molecules are labelled as Y1, Y2, Y3 and Y4²⁵.

The oligofluorene arms have a substituent effect on the BODIPY core and therefore the increase in the number of arms influences the photophysics of the material.

Solid state colour converters were spin-coated from solution to make thin films on quartz substrates for the photophysical studies. Films of thickness ca. 100 nm were deposited from solutions at concentrations of 10 and 20 mg/ml for photophysical and communications measurements respectively. These were spin-coated at 1500 rpm for 60 seconds. Figure 1 (bottom) shows the absorption and PL spectra of the films for each molecule. The absorption spectra show peaks at 450 and 620 nm for Y1; 350, 473 and 626 nm for Y2; 350, 477 and 625nm for Y3; and 366, 480 and 625nm for Y4. The absorption band below 400 nm is attributed to the fluorene arms and increases in intensity with arm length. Y1 has a peak in absorption at 450 nm matching well the emission of the blue LEDs used for lighting, and this feature shifts slightly to longer wavelengths for longer arms as seen in Figure 1. The peak emission wavelengths were in the red region of the spectrum, 663, 679, 682, 682 nm for Y1, Y2, Y3 and Y4 respectively. The PL emission comes from a donor-acceptor state as has been described previously²⁴. The bathochromic shift with increasing side-arm length is also evident in solution^{23,24} and indicates that on exciting the core, the BODIPY and neighbouring fluorene units acts as a single emitting species and not as separate components. Time-resolved fluorescence measurements were conducted using the time correlated single photon counting (TCSPC) technique, exciting the materials at 375 nm and detecting at the corresponding peak PL wavelengths. Exciting at 375 nm directly excites the oligofluorene arms, resulting in a fast energy transfer to the core component which would be a very fast (femto-picosecond) process. Due to instrument limitation, we do not see this very fast component. The resulting decays are shown in figure 2 and have bi-exponential decays. The lifetime values are given in table 1 in the nanosecond range and show Y3 having the shortest average decay. The PLQY of the films are also different showing lower values compared to solutions. In films the PLQY values range from 22-56%, whereas in solution form they were 55-62%²³.

Films of Y3 were encapsulated for bandwidth and data transmission measurements as they have the shortest lifetime and high PLQY. Samples were encapsulated with glass using the UV cured epoxy, NORLAND 68. The epoxy was placed on top of the BODIPY compound Y3. UV curing was at 365 nm for 1.5 minutes. The encapsulation process was conducted in a glove box which had a controlled and inert atmosphere to avoid exposure to water and oxygen molecules, thus reducing photo-oxidation effects during measurements.

The capacity of the data link scales with its modulation bandwidth. To assess the bandwidth of Y3, the experimental setup presented in figure 3b was used. The fast (broad bandwidth) excitation source and detector

used were a diode laser emitting at 450 nm and an avalanche photodiode. To eliminate any residual excitation light at the detector, a reflectance geometry was adopted in combination with a suitable long-pass dichroic filter. The excitation source was driven by an amplified sinusoidal voltage generated by the network analyser, this was combined with a DC voltage from a power source using a bias-T. The received AC signal from the APD was recorded by the network analyser. To measure the bandwidth of the system the AC amplitude of the received signal was recorded as function of the modulation frequency. To factor out the bandwidth contribution of the different components of the experimental setup, first the bandwidth of the measuring system was evaluated by directing the excitation light into the detector. This bandwidth was subsequently subtracted by the network analyser from the recorded bandwidth spectrum for the material. Figure 3a shows the results of the bandwidth measurement of the Y3 BODIPY as well as, for comparison, the bandwidth of a commercial phosphor plate CL-840 (Intematix ChromaLit). It is evident in figure 3 that the BODIPY material has a much higher -3 dB bandwidth (75.5 MHz) than that of the phosphor (5 MHz). The Y3 film bandwidth is also twice that measured with the same material in solution.

The data transfer capabilities were also studied using the same setup as for bandwidth measurements. The excitation laser was modulated in intensity using a binary On-Off Keying (OOK)²⁶²⁶ modulation scheme i.e. a binary ‘one’ was represented by a light pulse of duration $1/R_b$, where R_b is the data rate and a ‘zero’ was represented by the absence of the pulse. At the receiver, a simple threshold decoding was used in which the received signal was compared against a fixed threshold level based on the expected signal level for ‘zero’ and ‘one’. If the received signal is above the threshold level, it was assumed to be ‘one’, otherwise ‘zero’. Figure 4 shows the measured bit error rate (BER) as a function of transmitted data rate for OOK modulation. The measurements were taken using a pseudorandom binary sequence (PRBS) of $2^{14}-1$, of which ~ 82000 bits were analysed at the receiver for errors. Hence, the minimum BER level is limited to 10^{-4} to have statistically high confidence level. By considering the forward error correction (FEC) floor of 3.8×10^{-3} (as recommended by the International Telecommunication Union (ITU) standard), a data rate of 370 Mb/s was achieved which is 4 times higher than previously reported for BODIPY colour convertors in solution.

To summarise, BODIPY cored-polyfluorene semiconducting polymers have been explored in the solid state to establish their feasibility as a red colour converter for VLC applications. The solid-state films exhibit PL lifetime shorter than their solution based counterparts and therefore result in a higher bandwidth of 73 MHz and a data transmission rate of 370 Mb/s, which are two and four times higher than that in solution respectively. This

bandwidth is 16 times higher than of common commercially available phosphors and shows that organic materials are promising materials as colour converters for VLC.

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References

- ¹ H. Sirringhaus, Advanced Materials **26**, 1319 (2014).
- ² I. D. W. Samuel and G. A. Turnbull, Chemical Reviews **107**, 1272 (2007).
- ³ S. Chenais and S. Forget, Polymer International **61**, 390 (2012).
- ⁴ D. Moses, Applied Physics Letters **60**, 3215 (1992).
- ⁵ D. Amarasinghe, A. Ruseckas, A. E. Vasdekis, G. A. Turnbull, and I. D. W. Samuel, Proceedings of the IEEE **97**, 106 (2009).
- ⁶ N. Serdar Sariciftci, Current Opinion in Solid State and Materials Science **4**, 373 (1999).
- ⁷ Y. Infahsaeng V. Pranculis, Z. Tang, A. Devižis, D. a. Vithanage, C. S. Ponseca, O. Inganäs, A. P. Yartsev, V. Gulbinas, and V. Sundström, Journal of American Chemical Society **136**, 11331 (2014).
- ⁸ Y. Wang, P. O. Morawska, A. L. Kanibolotsky, P. J. Skabara, G. A. Turnbull, and I. D. W. Samuel, Laser Photonics Review **7**, L71–L76 (2013).
- ⁹ J. H. Burroughes, D. D. C. Bradley, A. R. Brown, R. N. Marks, K. Mackay, R. H. Friend, P. L. Burns, and A. B. Holmes, Nature **347**, 539 (1990).
- ¹⁰ Joseph R Lakowicz, *Principles of fluorescence spectroscopy*. (Springer, 2007).
- ¹¹ H. Elgala, R. Mesleh, and H. Haas, in *Communications Magazine, IEEE* (2011), Vol. 49, pp. 56.
- ¹² R. Mesleh, H. Elgala, and H. Haas, presented at the Wireless Communications and Networking Conference (WCNC), IEEE, 2012 (unpublished).
- ¹³ D. Tsonev, H. Chun, S. Rajbhandari, J. McKendry, S. Videv, E. Gu, M. Haji, S. Watson, A. Kelly, and G. Faulkner, Photonics Technology Letters, IEEE **26**, 637 (2014).
- ¹⁴ John Gancarz, Hany Elgala, and Thomas DC Little, Communications Magazine, IEEE **51** (12), 34 (2013).
- ¹⁵ D. O'Brien, L. Zeng, H. Le-Minh, G. Faulkner, J. W. Walewski, and S. Randel, presented at the Personal, Indoor and Mobile Radio Communications, 2008. PIMRC 2008. IEEE 2008 (unpublished).
- ¹⁶ Y. Wang, N. Chi, Y. Wang, R. Li, X. Huang, C. Yang, and Z. Zhang, Optics Express **21** (23), 27558 (2013).
- ¹⁷ R. Kraemer and M. Katz, in *Short-range wireless communications: Emerging technologies and applications* (Wiley, 2009).
- ¹⁸ P. A. Haigh, F. Bausi, Z. Ghassemlooy, I. Papakonstantinou, H. Le Minh, C. Fléchon, and F. Cacialli, Optics Express **22** (3), 2830 (2014).
- ¹⁹ G. Cossu, A. M. Khalid, P. Choudhury, R. Corsini, and E. Ciaramella, Optics express **20** (26), B501 (2012).

- 20 I. A. Barlow, T. Kreouzis, and D. G. Lidzey, *Applied Physics Letters* **94**, 243301 (2009).
- 21 I. A. Barlow, T. Kreouzis, and D. G. Lidzey, *Organic Electronics* **8**, 621 (2007).
- 22 H. Chun, P. Manousiadis, S. Rajbhandari, D. A. Vithanage, G. Faulkner, D. Tsonev, J.J. D. McKendry, S. Videv, X. Enyuan, G. Erdan, M. D. Dawson, H. Haas, G. A. Turnbull, I. D. W. Samuel, and D. C. O'Brien, *IEEE Photonics Technol.* **26**, 2035–2038 (2014).
- 23 M. T. Sajjad, P. Manousiadis, C. Orofino, D. Cortizo-Lacalle, A. L. Kanibolotsky, S. Rajbhandari, D. Amarasinghe, H. Chun, G. Faulkner, D. O'Brien, P. J. Skabara, G. A. Turnbull, I. D. W. Samuel, and *Advanced optical materials* **3**, 536 (2015).
- 24 C. Orofino-Pena, D. Cortizo-Lacalle, J. Cameron, M. T. Sajjad, P. P. Manousiadis, N. Findlay, A. L. Kanibolotsky, D. Amarasinghe, P. J. Skabara, T. Tuttle, G. A. Turnbull, and I. D. W. Samuel, *Beilstein Journal of Organic Chemistry* **10**, 2704 (2014).
- 25 A. L. Kanibolotsky, R. Berridge, P. J. Skabara, I. F. Perepichka, D. D. C. Bradley, and M. Koeberg, *Journal of the American Chemical Society* **126** (42), 13695 (2004).
- 26 Zabih Ghassemlooy, Wasiu Popoola, and Sujan Rajbhandari, *Optical wireless communications: system and channel modelling with Matlab®*. (CRC Press, 2012).

Figures

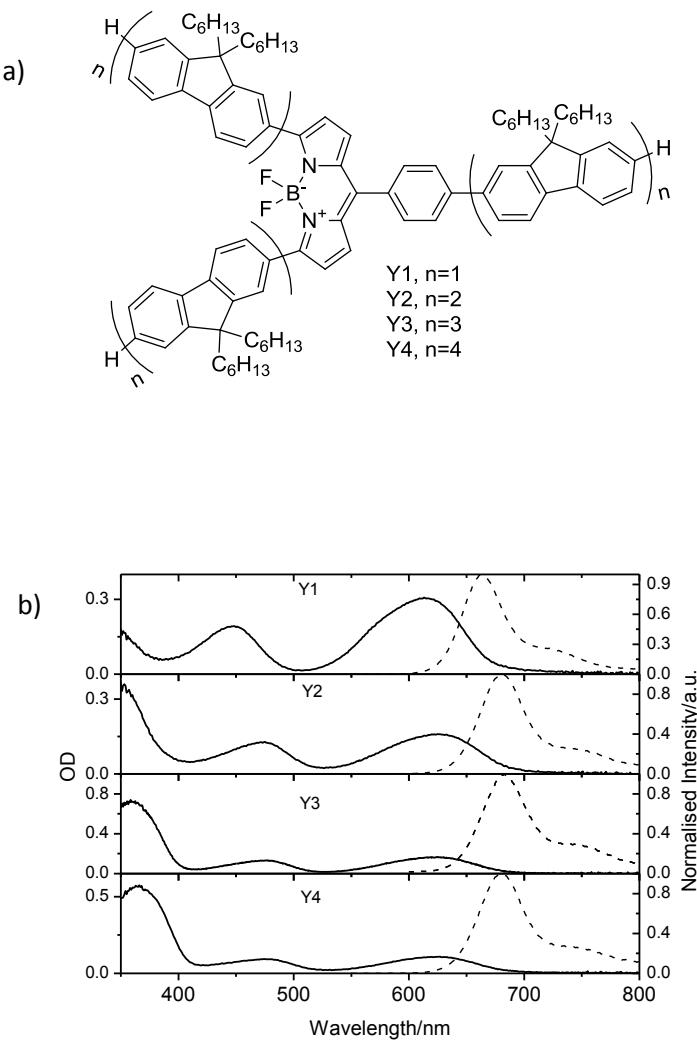


Figure 1: a) Molecular structure. b) Absorption and emission spectra of Y-BODIPY films

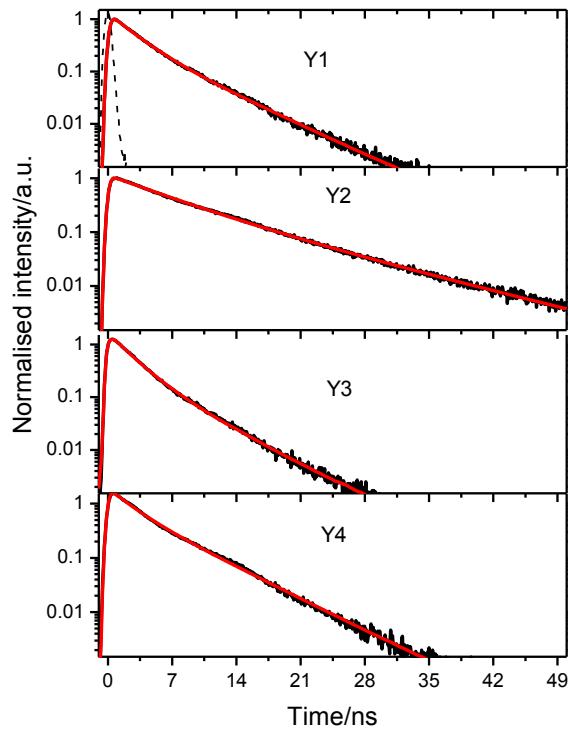
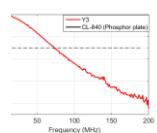


Figure 2: Time-resolved fluorescence measurements of Y-BODIPY films (black) and their fit (red) with the instrument response function (dashed line).

Polymer	PLQY %	A ₁	τ ₁ (ns)	A ₂	τ ₂ (ns)	τ _{avg} (ns)
Y1	22.4	0.44	2.35	0.56	5.3	4
Y2	55.9	0.53	5.87	0.47	10.72	8.2
Y3	40.6	0.48	1.9	0.52	4.3	3.2
Y4	49.2	0.26	2.07	0.74	5.08	4.3

Table 1: Photoluminescence quantum yield excited at 375 nm and lifetime of Y-BODIPY films

a)



b)

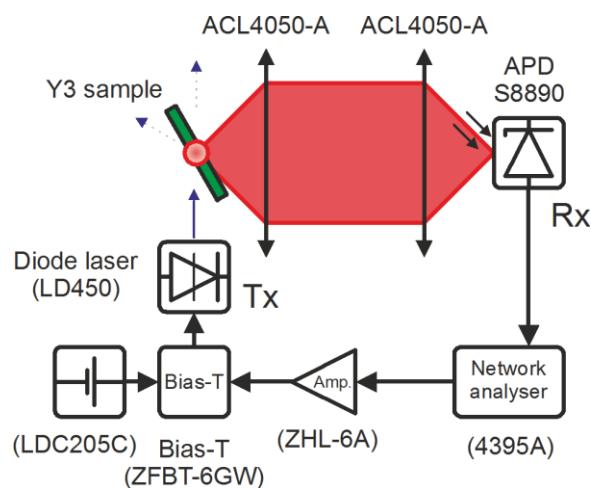


Figure 3: a) Result of small signal modulation bandwidth measurement for the Y3 BODIPY. The dotted line corresponds to -3 dB attenuation which defines the bandwidth of the system. b) Experimental setup used for the bandwidth and data transfer measurements.

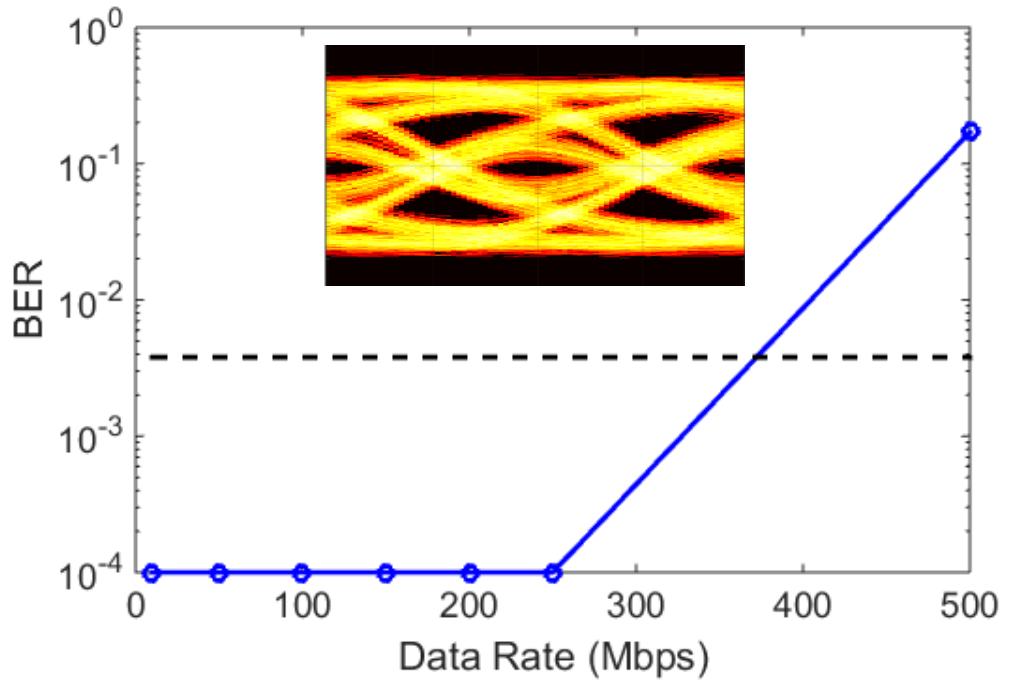


Figure 4: Plot of BER versus data rate for OOK. The dashed line corresponds to BER of 3.8×10^{-3} which is defined as the acceptable error floor for VLC. As inset, the eye diagrams at 250 Mb/s are presented.