

Understanding Photo-Oxidation in Ternary Organic Solar Cells

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Abstract

In recent years, organic solar cells (OSCs) have made significant progress achieving power conversion efficiencies close to 20% thanks to novel organic semiconductors [1]. However, beyond achieving high energy conversion efficiency, device stability remains a critical challenge for the large-scale commercialization of OSCs. In this context, a better understanding of the role of organic photovoltaic materials is essential, particularly for novel non-fullerene acceptors (NFAs) and their blends. These materials which constitute today the photoactive layer of OSCs play a crucial role in light absorption, charges generation, charges transport and overall device stability [2]. In this study, we focus on the photo-oxidation of the ternary blend PTQ10:Y12:PC₇₁BM, identified as one of the most promising candidates for industrial application [3]. The polymeric donor (PTQ10), the two acceptors (Y12, PC₇₁BM) and blends (PTQ10:Y12, PTQ10:PC₇₁BM, PTQ10:Y12:PC₇₁BM), deposited as thin films were analysed by UV-vis absorption spectroscopy. These films were then exposed for 100 hours in a solar chamber in ambient air (1 sun, 50°C), with spectral absorption measurements acquired periodically. Figure 1 shows the main absorption spectra obtained. The initial photo-oxidation results revealed unexpected behaviour. All three mixtures degraded faster than both individual materials. It was also observed that the addition of a small amount of fullerene slowed down the degradation of the mixture, acting as a radical trap. In addition, the initial ageing tests on the OSCs showed a 40% reduction in performance for the binary mixtures, compared to only 24% for the ternary active layer. Using complementary analysis with infrared spectroscopy measurements, this communication will elucidate the photochemical degradation pathways of electron acceptor and donor materials and help establish a correlation with the degradation of solar cells.

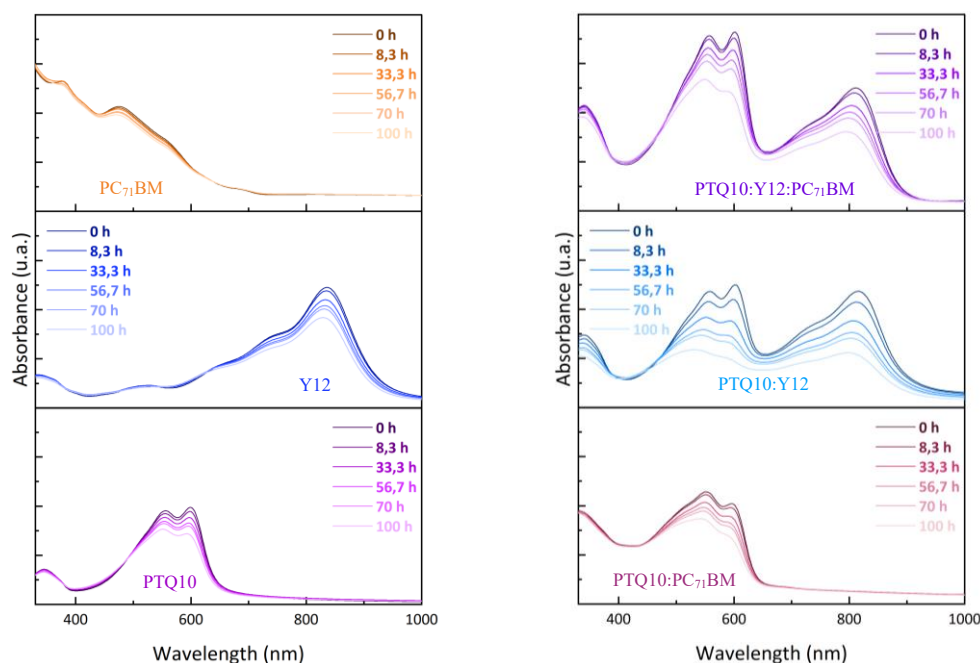


Fig. 1 – UV-vis spectra of photo-oxidized PTQ10, Y12, PC₇₁BM and blends films at different exposure times in a 1-sun chamber at 50°C in ambient air.

References

- [1] GUAN, Shitao, LI, Yaokai, XU, Chang, *et al.* Self-Assembled Interlayer Enables High-Performance Organic Photovoltaics with Power Conversion Efficiency Exceeding 20%. *Advanced Materials*, 2024, p. 2400342.
- [2] HAN, Jianhua, XU, Han, PALETI, Sri Harish Kumar, *et al.* Understanding photochemical degradation mechanisms in photoactive layer materials for organic solar cells. *Chemical Society Reviews*, 2024.
- [3] WACHSMUTH, Josua, DISTLER, Andreas, LIU, Chao, *et al.* Fully Printed and Industrially Scalable Semitransparent Organic Photovoltaic Modules: Navigating through Material and Processing Constraints. *Solar RRL*, 2023, vol. 7, no 21, p. 2300602.