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Materials for thermally activated delayed fluorescence and/or triplet fusion upconversion

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Two exciton harvesting mechanisms have shown great promise. Both result in delayed fluorescence but the photo-physical mechanisms are divergent.

Thermally activated delayed fluorescence (TADF) operates by thermal up-conversion of triplet excitons into singlets by reverse intersystem crossing

(RISC), which is possible in materials where the singlet-triplet energy gap, ΔE_{ST} , is very small. TADF, originally referred to as E-type fluorescence, was observed by Parker and Hatchard in the organic small molecule eosin in 1961.¹ This photophysical phenomenon remained relatively poorly explored until Adachi and his team reported the first example of high-efficiency OLEDs using organic TADF emitters in 2012;² this is because electroluminescence devices using TADF emitters can attain 100% IQE. Indeed, in the ten years since this seminal discovery, there has been an explosion of interest in the design of materials of this class, particularly as emitters in OLEDs, but also in photocatalysis, sensing and imaging applications. (Fig. 1).

Triplet-triplet annihilation-based photon upconversion (TTA-UC) (*aka* triplet fusion) materials, by contrast, produce singlet states

in a bimolecular fashion when two triplet excitons interact to form a singlet exciton with less than twice the energy of the triplet and a molecule in the ground state. First termed as P-type fluorescence by Parker and Hatchard as it was observed in pyrene, and sensitized upconverted fluorescence was observed in phenanthrene/naphthalene or proflavin hydrochloride/anthracene solutions.³ As with the history of TADF materials, research into TTA-UC materials remained sluggish until the early 2000s when TTA-UC by non-coherent, low-intensity light such as sunlight was first demonstrated.^{4,5} The ability of TTA-UC materials to convert low-energy photons into higher-energy photons has been exploited in a number of applications beyond OLEDs including photovoltaics, photocatalysis, and optogenetics.

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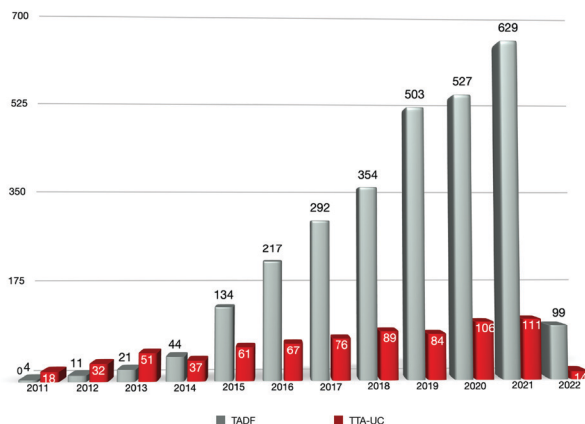


Fig. 1 Number of papers published on TADF and TTA-UC published since 2011. Data obtained from a Web of Science search of the terms “thermally activated delayed fluorescence” and or “TADF” and “triplet–triplet annihilation upconversion” or “TTA-UC” on 2/28/22.

Because of the recent remarkable progress in research related to these exciton harvesting mechanisms, we have organized a themed issue focusing on TADF and TTA-UC materials in *Journal of Materials Chemistry C*, which covers a wide range of topics from the development of new chromophores to the understanding of the mechanisms and their applications. Since TADF and TTA-UC are closely related topics albeit with different photophysical mechanisms, it is advantageous to combine them in one themed issue. For example, it has been recently reported that TADF materials show excellent performance as triplet sensitizers in TTA-UC because of the small energy loss in the intersystem crossing from the excited singlet state to the triplet state.

Though much chemistry has been recently explored in the development of TADF materials, both organic and metal-based, and the understanding of the underpinning photophysics has greatly improved, evidenced by the more than

thirty contributions to this themed issue, much remains to be addressed. For instance, the lifetime and performance of electroluminescence devices are correlated to the RISC rate; compounds that show very fast RISC remain both elusive and highly desired. In this issue, a large range of molecular design strategies, measurement techniques and applications are disclosed, complemented by insight provided by theoretical modeling.

Although numerous TTA-UC materials have been developed in the past 20 years, there are still many issues to be solved for practical applications. In particular, the efficiency of visible-to-ultraviolet TTA-UC and near-infrared-to-visible TTA-UC is still not sufficiently high, requiring the development of better sensitizers and emitters. In this themed issue, the latest TTA-UC materials with excellent performance and a detailed understanding of their exciton dynamics are reported. In addition, the quenching of the excited triplet by

oxygen molecules in air is a major problem in the application of TTA-UC in various fields. To solve this problem, rational material designs have been developed in recent years, and excellent examples are presented in this themed issue.

We hope that this themed issue will spark the interest of a wide range of researchers in luminophore design, spectroscopy, theoretical modeling, and device engineering of TADF and TTA-UC materials. We also hope that it will stimulate greater interaction between the TADF and TTA-UC fields and lead to an excellent fusion of research.

References

- 1 C. A. Parker and C. G. Hatchard, Triplet-singlet emission in fluid solutions. Phosphorescence of eosin, *Trans. Faraday Soc.*, 1961, **57**, 1894–1904.
- 2 H. Uoyama, K. Goushi, K. Shizu, H. Nomura and C. Adachi, Highly efficient organic light-emitting diodes from delayed fluorescence, *Nature*, 2012, **492**, 234–238.
- 3 C. A. Parker and C. G. Hatchard, Sensitized Anti-Stokes Delayed Fluorescence, *Proc. R. Soc. London, Ser. A*, 1962, **276**, 386–387.
- 4 D. V. Kozlov and F. N. Castellano, Anti-Stokes delayed fluorescence from metal–organic bichromophores, *Chem. Commun.*, 2004, 2860–2861.
- 5 S. Balushev, T. Miteva, V. Yakutkin, G. Nelles, A. Yasuda and G. Wegner, Up-Conversion Fluorescence: Noncoherent Excitation by Sunlight, *Phys. Rev. Lett.*, 2006, **97**, 143903.