

## **Formation of stimuli-responsive cyclophanes by self-assembly: the case of carbazole-based biradicals**

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Dynamic covalent bonds has recently received lot of attention because of their unique feature to become reversible under mild conditions.<sup>[1]</sup> In this context,  $\pi$ -conjugated biradical compounds has emerged as essential building blocks.<sup>[2]</sup> For instance, we have demonstrated that 2,7-dicyanomethylene-9-(2-ethylhexyl)carbazole biradical reversibly converts to a macrocycle cyclophane upon soft stimuli (temperature, pressure, light), showing strong chromic effects.<sup>[3]</sup> We now extent this study towards longer conjugated carbazole backbone (*i.e.*, indolocarbazole shown in Figure 1), aiming at investigating how the elongation of the conjugated backbone impacts on the formation of stimuli-responsive cyclophanes. The self-assembly process is investigated both in solution and solid state by linking theory and experiments.

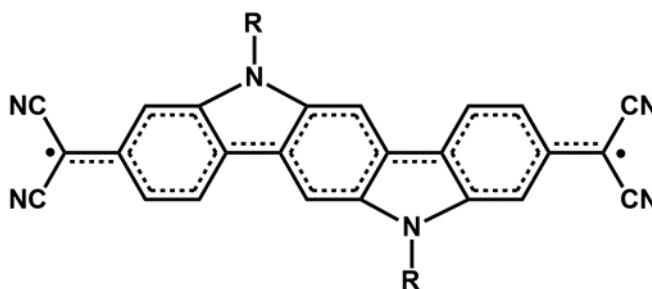


Figure 1. Indolocarbazole-based biradical systems under study.

### **References**

- [1] M. Abe, Chem. Rev. 2013, 113, 7011 – 7088.
- [2] O. Kohei, H. Shota, I. Yuki, S. Daisuke, S. Shu, Ang. Chem. Int. Ed. 2017, 56, 16597-16601.
- [3] D. Wang, C. C. Ferrón, J. Li, S. G. Valenzuela, R. P. Ortiz, J. T. L. Navarrete, V. H. Jolin, X. Yang, M. P. Álvarez, V. G. Baonza, F. Hartl, M. C. R. Delgado, H. Li, Chem. Eur. J. 2017, 23, 13776 13783.