

# SYNTHESIS AND CHARACTERIZATION OF NON-PLANAR TETRAAZAPEROPYRENES (TAPPS) VIA BAY-SUBSTITUTION

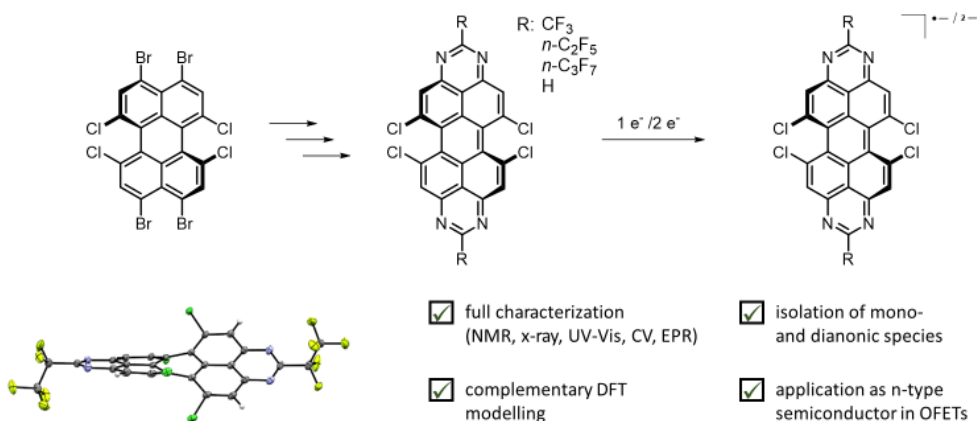
Benjamin A. R. Günther,<sup>a</sup> Sebastian Höfener,<sup>b</sup> Ute Zschieschang,<sup>c</sup> Hubert Wadepohl,<sup>a</sup> Hagen Klauk,<sup>c</sup> Lutz H. Gade<sup>a</sup>

<sup>a</sup> Heidelberg University, INF 270, 69120 Heidelberg, Germany

<sup>b</sup> Karlsruhe Institute of Technology (KIT), P.O. Box 6980, 76049 Karlsruhe, Germany

<sup>c</sup> MPI for Solid State Research, Heisenbergstr.1, 70569 Stuttgart, Germany

To this day, the research field of organic electronics is hampered due to the ongoing problematics in the fabrication of n-type semiconductors and their stability under ambient conditions.<sup>[1]</sup> Amongst different approaches facing this problem, the class of Tetraazaperopyrenes (TAPPs) has proven to circumvent many of these issues, however, has been limited in its variability as only the ortho-position could be functionalized.<sup>[2]</sup> In this context, we present an alternative TAPP-synthesis that enabled the isolation of a series of bay-chlorinated TAPPs. A detailed study revealed that the introduction of chloride substituents at this position not only leads to a stabilization of the LUMO levels and an increase of the electron affinity but also to an unprecedented twist of the peropyrene core of ca. 30°. Despite their non-planarity, the herein investigated TAPPs could be employed as n-type semiconductors in organic field effect transistors (OFETs) exhibiting electron mobilities of  $1\text{-}3 \times 10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ . The mono- and dianionic reduced species were isolated and characterized by UV-Vis spectroscopy and EPR or NMR, respectively. Finally, a comparison of bay- and ortho-chlorinated TAPPs was conducted to further elaborate the influence of the different substitution patterns.



[1] Y. Zhao, Y. Guo, and Y. Liu, *Adv. Mater.* **2013**, *25*, 5372–5391.

[2] S. C. Martens, U. Zschieschang, H. Wadepohl, H. Klauk, L. H. Gade, *Chem. Eur. J.* **2012**, *18*, 3498–3509; S. Geib, U. Zschieschang, M. Gsänger, M. Stolte, F. Würthner, H. Wadepohl, H. Klauk, L. H. Gade, *Adv. Funct. Mater.* **2013**, *23*, 3866–3874.