

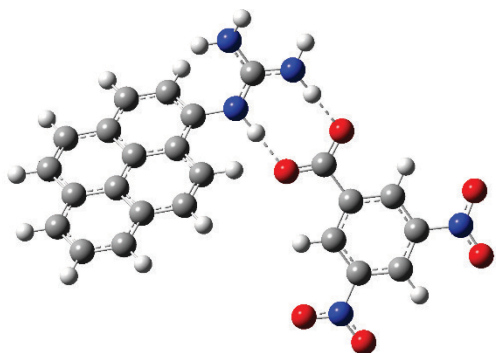
## GUANIDINE-CARBOXYLATE BONDING MOTIF IN SELF-ASSEMBLY OF AROMATIC DONOR-ACCEPTOR SYSTEMS

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Guanidine-carboxylate hydrogen bonding motif<sup>[1]</sup> was exploited in self-assembly of aromatic electron donor-acceptor (D-A) pairs. These artificial systems are models for study of photo-induced electron transfer processes through guanidine-carboxylate bonds in more complex natural photosynthetic



systems.<sup>[2]</sup> As typical electron donor systems, aromatic monosubstituted guanidines such as 1-pyrenylguanidine and 5-(4-phenyl guanidinyl)-10,15,20-triphenyl porphyrin were prepared. Synthetic procedure for conversion of aromatic amines to guanidines by *N,N'*-di-Boc-1H-pyrazole-1-carboxamide guanylation reagent was developed. Synthesis of aromatic guanidinium chromophores was optimized by using environmentally more friendly high-speed vibration milling technique.<sup>[3,4]</sup>

The fundamental physico-chemical properties of supramolecular D-A assemblies obtained via guanidine-carboxylate salt bridges were studied spectroscopically. Molecular modelling (DFT method) was employed in design of D-A systems and study of their molecular and electronic structures.

### REFERENCES

- [1] K. A. Schug, W. Lindner, *Chem. Rev.* **2005**, *105*, 67–114.
- [2] M. D. Archer, J. Barber, Photosynthesis and photoconversion. In *Molecular to global photosynthesis*, M. D. Archer, J. Barber, eds., Imperial College Press, London, **2004**, 1–41
- [3] D. Margetić, V. Štrukil, *Mechanochemical Organic Synthesis*, Elsevier, Amsterdam **2016**.
- [4] M. Đud, D. Margetić, *Internat. J. Org. Chem.* **2017**, *7*, 140–144.