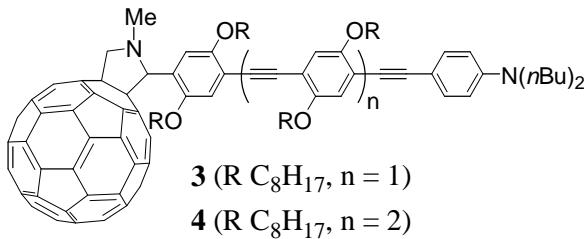
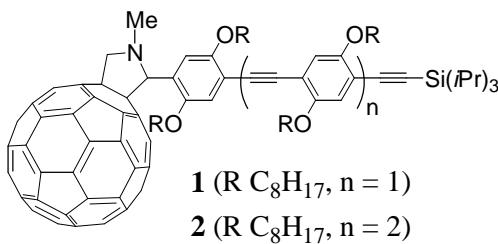


Photovoltaic Devices from Fullerene-Oligophenyleneethynylene Conjugates

Jean-François NIERENGARTEN

Institut de Physique et Chimie des Matériaux de Strasbourg, Groupe des Matériaux Organiques
23 rue du Loess, 67037 Strasbourg (France)

Exploitation of the particular electronic properties of fullerenes for solar energy conversion has become a field of intensive investigation.¹ The modification of semiconductor or metal electrodes with fullerene thin films has revealed promising photoelectrochemical properties.² Good solar energy conversion efficiencies have also been obtained with sandwich-type π -conjugated polymer/fullerene heterojunctions.³ As part of this research, we have recently shown that fullerene derivatives in which an oligophenylenevinylene (OPV) group is attached to C₆₀ can be incorporated into photovoltaic cells.⁴ This molecular approach for solar energy conversion appears to be particularly interesting since the bicontinuous network obtained by chemically linking the hole-conducting OPV moiety to the electron-conducting fullerene subunit prevents any problem arising from bad contacts at the junction, as observed for polymer/C₆₀ blends. Furthermore, this new synthetic approach also offers great versatility for design tuning the photovoltaic system. We now report the preparation of plastic solar cells from fullerene-oligophenyleneethynylene (OPE) derivatives **1-4**.⁵



For compounds **1** and **2**, the efficiency of the resulting photovoltaic devices is similar to those prepared from corresponding fullerene-OPV conjugates.⁴ Interestingly, by increasing the donating ability of the conjugated oligomer substituents in **3** and **4** owing to the presence of the aniline group, the efficiency and the sensitivity of the photovoltaic devices are increased by one order of magnitude.⁶ The latter observation shows clearly the advantage of our molecular approach allowing structure/activity relationships.

Acknowledgements

This work was supported by the CNRS and the French Ministry of Research.

References

- [1] J.-F. Nierengarten, G. Hadzioannou, N. Armaroli, *Mater. Today* **2001**, *4*, 16-18.
- [2] For recent examples, see: H. Imahori, T. Azuma, A. Ajavakom, H. Norieda, H. Yamada, Y. Sakata, *J. Phys. Chem. B* **1999**, *103*, 7233-7237; P. V. Kamat, S. Barazzouk, S. Hotchandani, K. G. Thomas, *Chem. Eur. J.* **2000**, *6*, 3914-3921; C. Luo, D. M. Guldi, M. Maggini, E. Menna, S. Mondini, N. A. Kotov, M. Prato, *Angew. Chem. Int. Ed.* **2000**, *39*, 3905-3909; P. V. Kamat, S. Barazzouk, K. G. Thomas, S. Hotchandani, *J. Phys. Chem. B* **2000**, *104*, 4014-4017; O. Enger, F. Nesch, M. Fibbioli, L. Echegoyen, E. Pretsch, F. Diederich, *J. Mater. Chem.* **2000**, *10*, 2231-2233; A. Ikeda, T. Hatano, S. Shinkai, T. Akiyama, S. Yamada, *J. Am. Chem. Soc.* **2001**, *123*, 4855-4856; S. Zhang, D. Dong, L. Gan, Z. Liu, C. Huang, *New J. Chem.* **2001**, *25*, 606-610; H. Gao, G. Luo, A. L. Ottava, H. T. Tien, *J. Electroanal. Chem.* **2001**, *496*, 158-161; H. Imahori, H. Norieda, Y. Nishimura, I. Yamazaki, Y. Sakata, S. Fukuzumi, *J. Am. Chem. Soc.* **2001**, *123*, 100-110.
- [3] G. Yu, J. Gao, J. C. Hummelen, F. Wudl, A. J. Heeger, *Science* **1995**, *270*, 1789-1791; L. S. Roman, M. R. Andersson, T. Yohannes, O. Inganäs, *Adv. Mater.* **1997**, *9*, 1164-1168; L. Ouali, V. K. Krasnikov, U. Stalmach, G. Hadzioannou, *Adv. Mater.* **1999**, *11*, 1515-1518.
- [4] J.-F. Nierengarten, J.-F. Eckert, J.-F. Nicoud, L. Ouali, V. Krasnikov, G. Hadzioannou, *Chem. Commun.* **1999**, 617-618; J.-F. Eckert, J.-F. Nicoud, J.-F. Nierengarten, S.-G. Liu, L. Echegoyen, F. Barigelletti, N. Armaroli, L. Ouali, V. Krasnikov, G. Hadzioannou, *J. Am. Chem. Soc.* **2000**, *122*, 7467-7479.
- [5] T. Gu, J.-F. Nierengarten, *Tetrahedron Lett.* **2001**, *42*, 3175-3178.
- [6] T. Gu, D. Tsamouras, C. Melzer, V. Krasnikov, J.-P. Gisselbrecht, M. Gross, G. Hadzioannou, J.-F. Nierengarten, *ChemPhysChem.*, in press.