

A Novel n-Channel Organic Semiconductor Based on Pyrene-phenazine Fused Monoimide and Bisimide

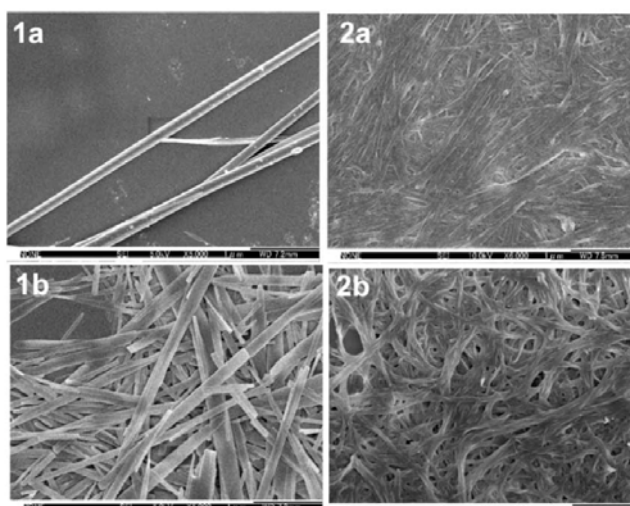
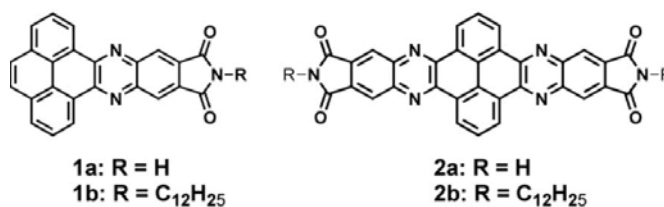
Long Chen, Tomoya Ishizuka and Donglin Jiang*

Department of Materials Molecular Science, Institute for Molecular Science,
5-1 Higashiyama, Myodaiji, Okazaki 444-8787, Japan

e-mail: longchen@ims.ac.jp

Synthesis of organic n-channel semiconductors is of increasing interest due to their utility in organic complementary inverters upon integration with p-channel semiconductors.^[1] However, there are only fewer n-channel semiconductors compared with p-channel counterparts. Typical examples of n-channel semiconductors reported up to date have been limited to naphthalene bisimides,^[2] perylene bisimides,^[3] fullerene and its derivatives,^[4] and perfluorinated conjugated molecules such as fluorinated copper phthalocyanines.^[5]

Here we report the synthesis of a new type n-channel semiconductor based on pyrene fused phenazine monoimide and bisimides (Figure on the right side; 1a and 2a). We also report their self-assembly to form well-defined nanostructures, as investigated by FE-SEM, TEM, AFM and XRD measurements. In order to tune self-assembly structure and improve solubility for solution-process device fabrication, dodecyl chains were anchored to the N-termini of the monoimide and bisimide derivatives (1b and 2b). Both the non-substituted and didodecyl-anchored bisimides self-assemble to give highly ordered nanobelts. Cyclic voltammetry measurement shows two redox peaks in negative potential region. The conductivity of the assembled nanobelts was investigated by a two-probe method on a 10- μm Pt gap. Upon doping with hydrazine,^[6] the nanobelt shows a largely enhanced electric current.



Reference:

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