

**Martin-Luther-Universität Halle-Wittenberg**  
**Naturwissenschaftliche Fakultät II**  
**Chemie und Physik**  
**SFB TRR 102**



**SFB/TRR 102-KOLLOQUIUM**

**am Donnerstag, dem 07.06.2012, 17.15 Uhr,**

Gustav Mie Hörsaal, Theodor – Lieser - Str. 9, 06120 Halle

Es spricht:

**Prof. George Floudas**

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zum Thema:

**“Discotic liquid crystals in motion”**

Discotic liquid crystals (DLCs) are materials where self-assembly is driven by non-covalent intermolecular interactions. During the self-organization process, the disk-shaped molecules organize into columns that further assemble into two-dimensional arrays whereas the alkyl chains fill the intercolumnar space. Highly ordered columnar structures of hexa-peri-hexabenzocoronenes (HBC) were found to be very promising as active semiconductors in organic FETs (molecular wires) and photovoltaic devices. The self-assembly and molecular dynamics in a series of dipole functionalized nanographenes were studied using thermodynamic (DSC, PVT), structural (WAXS) and dynamic (DS, site-specific NMR, Rheology) probes [1-5]. These DLCs undergo a phase transformation from a dipolar ordered but structurally disordered liquid crystalline (LC) phase at higher temperatures to a dipolar disordered but structurally ordered crystalline phase (Cr) at lower temperatures [4]. The effect of dipole substitution is to change the energetics and to stabilize the LC phase. Within the LC phase the intra- and inter-columnar thermal expansions were different but both positive. In contrast, the Cr phase shows a negative thermal expansion (NTE) [3]. With respect to the dynamics, these small molecules display a range of hierarchical molecular motions (i.e. polymer-like) ranging from 0.01 ps to hours as shown by FTIR, NMR, DS and Rheology [5].

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  - [4] C. Grigoriadis, N. Haase, H.-J. Butt, K. Müllen, G. Floudas Soft Matter, 7, 4680 (2011).
  - [5] M. R. Hansen, X. Feng, V. Macho, K. Müllen, H.W. Spiess, and G. Floudas Phys. Rev. Lett., 107, 257801 (2011).