

NMR relaxation study of molecular dynamics of liquid crystalline side-on organosiloxane tetrapodes

D. Filip^{a, b}, C. Cruz^{a, c}, P.J. Sebastião^{a, c}, M. Cardoso^a, A. C. Ribeiro^{a, c},
M. Vilfan^d, T. Meyer^e, P. H. J. Kouwer^e, G. H. Mehl^e

a CFMC-UL Av. Prof. Gama Pinto 2, 1649-003 Lisboa, Portugal

*b "Petru Poni" Institute of Macromolecular Chemistry, Alleea Gr. Ghica
Voda 41A, 700487-Iasi, Romania*

c IST-UTL, Av. Rovisco Pais 1049-001 Lisboa, Portugal

d Jozef Stefan Institute, Jamova 39, 1000 Ljubljana, Slovenia

*e Dep. Chemistry, University of Hull, Cottingham Road, Hull HU6 7RX,
United Kingdom*

Proton NMR relaxation measurements were carried out on two liquid crystalline organosiloxane tetrapodes with side-on mesogenic groups, exhibiting nematic and smectic C phases, and on a monomeric analogue.^{1,2} NMR relaxometry of the tetrapodes systems yields T_1^{-1} dispersions clearly different from those of conventional calamitics. The influence of molecular tendency to form interdigitated structures is evidenced by frequency dependent relaxation rate in the isotropic phase – indicating the presence of ordered clusters far above the phase transition – and by the diminished role of molecular self-diffusion in ordered phases. Nematic-like director fluctuations are the dominating relaxation mechanism whereas the translational displacements are strongly hindered by the interdigitation of dendrimer arms.

(1) K. Merkel, A. Kocot, J.K. Vij, R. Korlacki, G.H. Mehl, and T. Meyer, *Phys. Rev. Lett.* **2004**, 93, 237801

(2) D. Filip, C. Cruz, P.J. Sebastião, A. C. Ribeiro, T. Meyer, G. H. Mehl, *Mol. Cryst. Liq. Cryst.* **2005**, 436, 17