



Neutron Spin Echo Spectroscopy Viscoelasticity Rheology (Advances in Polymer Science) (Vol 134)

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Viscoelastic and transport properties of polymers in the liquid (solution, melt) or liquid-like (rubber) state determine their processing and application to a large extent and are of basic physical interest [1?3]. An understanding of these dynamic properties at a molecular level, therefore, is of great importance.

However, this understanding is complicated by the fact that different motional processes may occur on different length scales and that the dynamics are governed by universal chain properties as well as by the special chemical structure of the monomer units [4,5]. The earliest and simplest approach in this direction starts from Langevin equations with solutions comprising a spectrum of relaxation modes [1?4]. Special features are the incorporation of entropic forces (Rouse model, [6]) which relax fluctuations of reduced entropy, and of hydrodynamic interactions (Zimm model, [7]) which couple segmental motions via long-range back flow fields in polymer solutions, and the inclusion of topological constraints or entanglements (reptation or tube model, [8?10]) which are mutually imposed within a dense ensemble of chains. Another approach, neglecting the details of the chemical structure and

concentrating on the universal elements of chain relaxation, is based on dynamic scaling considerations [4,11]. In particular in polymer solutions, this approach offers an elegant tool to specify the general trends of polymer dynamics, although it suffers from the lack of a molecular interpretation. A real test of these theoretical approaches requires microscopic methods, which simultaneously give direct access to the space and time evolution of the segmental diffusion. Here, quasi-elastic scattering methods play a crucial role since they allow the measurement of the corresponding correlation functions. In particular, the high-resolution neutron spin echo (NSE) spectroscopy [12?15] is very suitable for such investigations since this method covers an appropriate range in time (0.005 to 40 ns) and space (r in nm [15]). Furthermore, the possibility of labelling by hydrogen-deuterium exchange allows the observation of single-chain behavior even in the melt.

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