

Backbone Stiffness of Cylindrical Brushes Consisting of Rodlike Side Chains. Influence of Side Chain Length on Main Chain Stiffness

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Introduction

A macromonomer is any polymer or an oligomer with a polymerizable functionality as an end group. Formally, the macromonomer homopolymerizes to afford a star- or comb-shaped polymer and copolymerizes with the conventional monomer to give graft copolymer. Recently, some interesting properties of poly-(macromonomers) have been explored as a simple model of brush polymers. However, most of macromonomer used in these studies is limited to that consisting from the flexible chains.[1,2]

In the present study, we report conformational properties of the polymacromonomers, consisting of a flexible polystyrene chain backbone and rod-like poly(*n*-hexyl isocyanate)(PHIC) side chains by SAXS, light scattering, $[\eta]$ and AFM measurements in THF.[3-5]

Experimental

SAXS Measurements. SAXS measurements of polymacromonomers were carried out in THF at 25 °C, using BL-10C with a synchrotron orbital radiation as an X-ray source set up in the Photon Factory of the High Energy Accelerator Organization at Tsukuba, Ibaraki, Japan. The wavelength of the X-ray was 1.488 Å. The scattered intensity was recorded by a position-sensitive proportional counter (PSPC) with 512 channels over a scattering vector range from 0.02 to 0.30 Å⁻¹. The scattered vector was calibrated using a sixth peak of dry collagen.

Results and Discussion

Figure 1 shows a double logarithmic plot of cross-sectional radius of gyration, $\langle R_c^2 \rangle_0^{1/2}$ of poly(VB-HIC- n_w) determined by SAXS in THF with weight averaged degree of polymerization of the side chain (n_w). The solid curve is a theoretical one given by the equation,

$$\langle R_c^2 \rangle_0 = \langle R_c^2 \rangle_m + (2L_w)^2/12$$

where $\langle R_c^2 \rangle_m$ is that of the main chain and L_w is the weight-averaged contour length of a rod. The solid curve quantitatively describes the experimental data, implying the brush-like conformation consisting of the rods as a side chain.

Figure 2 shows a plot of main chain stiffness parameter (λ^{-1}) with contour length of side chain. It is clearly seen that the main chain stiffness linearly increases with increasing side chain length.

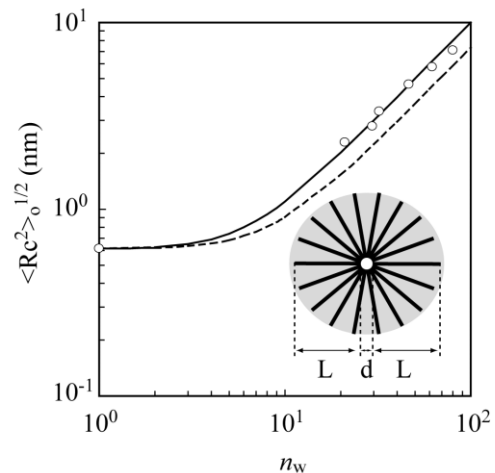


Figure 1. Dependence of the measured $\langle R_c^2 \rangle_0^{1/2}$ on degree of polymerization of side chain n_w for poly(VB-HIC- n_w) in THF.

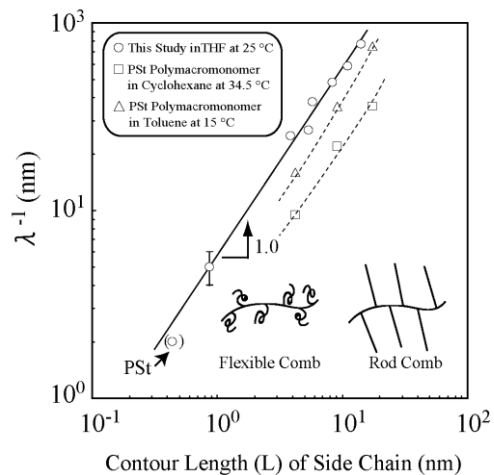


Figure 2. Dependence of Kuhn segment length λ^{-1} on side chain length L , compared with reference values of polystyrene macromonomer in cyclohexane and toluene.

References

- [1] K. Ito et al. *Adv. Polym. Sci.* 142, 129(1999).
- [2] S. Kawaguchi et al. *Adv. Polym. Sci.*, **175**, 317(2005).
- [3] M. Kikuchi et al. *Des. Monom. Polym.*, **7**, 603 (2004).
- [4] S. Kawaguchi et al. *Macromolecules*, **40**, 950(2007).
- [5] M. Kikuchi et al., *Polym. J.*, **39**, 330(2007).

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