

*Erratum***Apparent slip at polymer-interface**J.L. Goveas^{1,a} and G.H. Fredrickson²¹ Materials Research Laboratory, University of California, Santa Barbara CA 93106-5130, USA² Department of Chemical Engineering, University of California, Santa Barbara CA 93106-5130, USAEur. Phys. J. B **2**, 79–92 (1998)**Section 5**

One of the goals of this paper was to derive an approximate stress constitutive equation for application to inhomogeneous polymer melts. While convenient, our approach was to force an N -mode model into a dynamical equation (Eq. (5.3)) for a stress field with a single relaxation time, which is an inexact description of melt dynamics, even for the homogeneous case.

We note a misprint in equation (5.3), where the term $6\Sigma_{ij}/(\zeta R_g^2)$ should appear with a negative sign.

Perhaps more importantly, the stress relaxation term $-6\Sigma_{ij}/(\zeta R_g^2)$ was chosen to reproduce the bulk Rouse viscosity, but this term incorrectly describes the normal stress relaxation. It should be replaced by

$$-\left\{ \frac{6}{\zeta R_g^2} (1 - \theta_{ij}) + \frac{45b^2}{36^2 \zeta R_g^4} \theta_{ij} \right\} \Sigma_{ij} \quad (1)$$

where θ_{ij} is a scalar function such that

$$\begin{aligned} \theta_{ij} &= 0, \quad i \neq j \\ &= 1, \quad i = j. \end{aligned}$$

While this change is immaterial for the purposes of the present paper, it can be significant for other applications in which first normal stresses come into play.

Appendix B

The mixing free energy given by equation (B.8) is missing terms that describe the entropy of mixing between A and B polymers. Such terms must be added carefully to be consistent with diagonal terms appearing in the stress constitutive equation, and to generate appropriate diffusion terms in equation (B.11). Details may be found in a subsequent paper (J.L. Goveas, G.H. Fredrickson, *J. Rheol.* **43** 1261 (1999)).

^a e-mail: jlgoivas@mrl.ucsb.edu