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Self-Assembly of Lamellar Microphases in Linear Gradient Copolymer Melts¹ NICHOLAS B. TITO, Dartmouth College, SCOTT T. MIL-NER, The Pennsylvania State University, JANE E. G. LIPSON, Dartmouth College — The ability to create 'designer copolymers' with tunable properties by tailoring their monomer composition has garnered recent interest in their molecular self-assembly. Here we investigate lamellar microphases in linear gradient binary copolymer melts using a variety of techniques, including solutions of self-consistent field equations, scaling theory, and analysis of the strong-segregation limit. The Flory scaling theory predicts the scaling of the equilibrium lamellar width L_{eq} as a function of comonomer incompatibility as characterized by χ . From the strongly segregated limit there are conformational fluctuations, and it is the tradeoff between the entropic effect of these relative to repulsive comonomer interactions that determines L_{eq} . We discover that $L_{eq} / R_g \sim (\chi N)^{1/6}$; remarkably, this is the same result as for symmetric diblock copolymers, although for quite different physical reasons.

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