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Block copolymers confined in a nanopore: Pathfinding in a curving and frustrating flatland

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We have studied structure formation in a confined block copolymer melt by means of dynamic density functional theory. The confinement is two dimensional, and the confined geometry is that of a cylindrical nanopore. Although the results of this study are general, our coarse-grained molecular model is inspired by an experimental lamella-forming polystyrene-polybutadiene diblock copolymer system [K. Shin *et al.*, *Science* **306**, 76 (2004)], in which an exotic toroidal structure was observed upon confinement in alumina nanopores. Our computational study shows that a zoo of exotic structures can be formed, although the majority, including the catenoid, helix, and double helix that were also found in Monte Carlo nanopore studies, are metastable states. We introduce a general classification scheme and consider the role of kinetics and elongational pressure on stability and formation pathway of both equilibrium and metastable structures in detail. We find that helicity and threefold connections mediate structural transitions on a larger scale. Moreover, by matching the remaining parameter in our mesoscopic method, the Flory-Huggins parameter χ , to the experimental system, we obtain a structure that resembles the experimental toroidal structure in great detail. Here, the most important factor seems to be the roughness of the pore, i.e., small variations of the pore radius on a scale that is larger than the characteristic size in the system. © 2008 American Institute of Physics. [DOI: [10.1063/1.2829406](https://doi.org/10.1063/1.2829406)]

I. INTRODUCTION

Pattern formation of block copolymers in constraint situations or *confinement* is an important topic in polymer research, since the meso- or microstructure can be much better controlled when compared to the bulk. In general, molecular conformations and assembly are strongly influenced by confinement. In the absence of external constraints, the microstructure is dictated by the interaction between segments comprising the copolymer, the volume fraction of the blocks, and the molecular architecture. In a confined system, however, interfacial interactions, symmetry breaking, structural frustration, and confinement-induced entropy loss play a determining role and may lead to structures that differ from the ones found in bulk. As a direct result phase separation in confinement has been the subject of extensive theoretical and experimental studies.¹ The intriguing prospects from a technological viewpoint are the novel structures that can be achieved and that may serve as scaffolds for other nanostructures.

The simplest example of a block copolymer is a linear *AB* diblock copolymer. In the bulk several stable periodic microstructures can be formed, among which are lamellar, hexagonal, and body-centered cubic phases.² The equilibrium behavior for *AB* diblock copolymers in the bulk has been mapped out both experimentally and theoretically; the situation for more complex multiblock and/or branched

block copolymers is much less clear. Here, we focus on symmetric or nearly symmetric *AB* diblock copolymers. In the bulk, these copolymers microphase separate into lamellar microdomains with a characteristic equilibrium period L_0 , where grains of ordered lamellar microdomains are randomly oriented. Global orientation of the microdomains can be induced by confinement of the block copolymer in thin supported films or slits (one-dimensional confinement).^{3–6} Additionally, the interplay between surface fields (interaction with confining surfaces) and confinement effects (commensurability) can affect the phase behavior, and lead to the formation of surface reconstructions or hybrid structures.^{7–9} Both statics and dynamics^{10,11} of thin film phase behavior have been well studied.

An obvious next step in experimental and theoretical research is to consider systems where the confinement is effectively two dimensional. This is the case when the melt is confined inside a cylindrical nanopore of radius R . From a conceptual point of view, this type of confinement differs from the one-dimensional (1D) confinement since (1) there is only one confining surface, so there is only one surface field; (2) the nanopore solid surface is curved. By definition, surface reconstructions (analogs of the uni- or multilayer parallel lamellae $L_{\parallel}$ in slits) are therefore curved as well, which has an effect on the entropy contribution of the structure in the free energy. This can either result in a breakup of the nanopore in two relatively independent regions (close and away from the pore surface), or in curved multilayer structures that experience lamellar bending throughout the pore, with an

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increased bending towards the center of the pore, (3) just like in slits, surface reconstructions may be able to adapt their layer spacing to some extent in order to suit the cylinder radius R . One can expect the influence of this type of frustration to be less than in slits, as there is no block preference in the center of the pore. However, for small R , structures can experience substantial frustration due to incommensurability. To resolve this unfavorable situation, the lamellae can adapt an orientation perpendicular to the nanopore wall or the chains may find alternative packing and form hybrid structures, like in a slit, and (4) in addition to the commensurability issue for parallel structures or surface reconstructions [see (2)] the length of a nanopore can also affect the formation of perpendicular structures (incommensurability along the pore). In slits, these perpendicular structures are found for strong surface fields at incommensurable film thickness, and for weak surface fields independent of the film thickness. In nanopores, a mismatch of the pore length and the natural spacing L_0 can lead to extensional forces, and may stabilize hybrid structures as well.

Based on rather small data set of surface interactions and nanopore radii R , two pioneering computational studies (by Monte Carlo¹² and dynamic self-consistent field theory¹³) identified two structures: A slab and a multiwall tube morphology.¹³ In analogy with the classification scheme for structures in a slit,^{14,15} slabs can be associated with lamellae perpendicular to the confining surface ($L_\perp$) and multiwall tubes with surface reconstructions: Lamellae that are parallel to the confining surface ($L_\parallel$). One should note that this remark implies that the nature of these structures is the same as the lamellar structure found in bulk. It was concluded that two mechanisms control the structure formation in a nanopore: (1) In the case of weak surface interactions the lamellae orient perpendicular to the pore wall to form lamellar slabs and (2) for strong surface interactions, one of the blocks segregates preferentially to the pore wall, and the lamellae line up and bend to form concentric cylinders, the number of which is determined by the cylinder radius. The effect of commensurability was found to be less significant in this type of confinement, since no perpendicular structures were observed for incommensurate R and higher surface interactions.

Very recently, the significance of these theoretical studies increased dramatically by the appearance of a number of intricate experimental studies of polystyrene-*block*-polybutadiene^{16–19} (PS-*b*-PB) and polystyrene-*block*-poly(methyl methacrylate)²⁰ (PS-*b*-PMMA) diblock copolymers confined in nanopores. In these studies, bulk lamella-, cylinder-, and sphere-forming block copolymers were introduced into nanoscopic cylindrical pores in alumina membranes. For bulk lamella-forming block copolymers the predicted morphology^{12,13} of slabs was not found, most probably due to the strong disbalance of surface energies in the experimental setup (although these were not measured as such). The most frequent found structure was a concentric cylinder (multiwall tube¹³) morphology. However, also a new structure, a stacked-disk or toroidal-type¹⁶ structure, was found inside a nanopore with

an incommensurate pore diameter ($d/L_0 \sim 2.6$, with d the pore diameter).

As a result of these experimental findings, computational studies have considered the phase behavior of confined symmetric diblock copolymers in more detail, by means of self-consistent field theory²¹ (SCFT) and Monte Carlo^{22–25} (MC) methods. Based on two-dimensional (2D) calculations, the work of Li *et al.*²¹ aimed at constructing phase diagrams (χN vs $f=f_A$), and found lamellae for $f_A=1/2$ (the diagram was calculated for a single radius $R=8.5R_g$). Although this work can be seen as a stepping stone for the understanding of the nature of phase transitions under varying conditions, their findings do not directly relate to the essentially three-dimensional (3D) structures found in experiments. For instance, the slab morphology is out of the scope of their calculations. Moreover, the influence of the nanopore radius and the surface energetics was not considered in detail. Based on lattice MC simulations, Chen *et al.*; Feng and Ruckenstein, and Wang indeed found several new structures in three dimensions: A single helix, catenoid cylinder, gyroidal, stacked circle, and disordered structure in Ref. 22; a mesh, lamellae parallel to the pore axis, single helix, and double helix structure in Ref. 23; and catenoid cylinder in Ref. 25. A later study of Feng and Ruckenstein concentrated on the stability of the helical structure.²⁴ From a visual comparison we conclude that the mesh structure²³ is the same as catenoid cylinder.²² Similarly, one can claim that the gyroidal structure is, in fact, a defected structure, and comprises a coexistence of a mesh and a double helix structure.²² We adapt a common notation, and conclude that this reduces the number of computed nanopore structures to seven: Stacked disk, concentric cylinder, lamellae parallel to the pore axis, catenoid cylinder, disordered, single helix, and double helix.

Although these new structures were found, a fundamental understanding of the underlying mechanisms such as achieved for thin films (effective 1D confinement) is absent, except for the MC study of Wang.²⁵ However, the latter study is restricted to strong segregation, and most remarkably several of the other structures (for instance, the interesting helix and double helix) were not found at all. Moreover, the origin of the experimental stacked-disk or toroidal-type¹⁶ structure is still unexplained. More fundamental studies, including scans of a larger parameter space, are clearly needed for a deeper understanding.

In principle, one can anticipate several regimes. Equilibrium morphologies are minima of the free energy, containing both energetic and entropic contributions. In the absence of any surface field and for a strong surface field, either one of these contributions is dominating, leading to the stacked-disk and concentric cylindrical structures. Between these regions, for weak surface fields, the chains have more flexibility to adapt their packing and the system can and apparently will adapt other morphologies. Incommensurability may play a subtle role here. The details of this interplay remain to be determined, for instance, by the calculation of a structure diagram in three dimensions, depending on R and the surface interaction strength. Since we consider lamella-forming sys-

tems, f_A , the volume fraction of the A -block, is fixed (in contrary to Ref. 21) and the diagram is conceptually similar to the one on Ref. 14 for confined films.

Moreover, two factors have not been considered in detail yet: The influence of the kinetic pathway and the value of $\chi_{AB}N$, where N is the total length of the block copolymer. Although we will not focus into detail in the latter factor, from Ref. 21 one can conclude that an increase of $\chi_{AB}N$ may lead to a decrease of the number of concentric lamellae. Here, we focus on calculating the structural diagram and the question of stability. Whether interesting structures such as double helix can actually be manufactured experimentally may be a very subtle issue. It is known that in strong confinement the dynamics may slow down and structures may be frozen into metastable states. This is particularly the case when the free energy difference between states is small or when there is a large energy barrier between different local minima of the free energy. In nanopores, this situation was actually observed in MC studies: In Ref. 22 for weak surface fields up to three different structures were found. Previously, a dynamic density functional theory (DDFT) study¹³ identified the single helix structure as a long-living intermediate structure between complete mixing and stacked disks for zero surface field. We will use this method here to consider the experimental system of Ref. 16 in detail. In contrast to traditional schemes of polymer phase separation dynamics where a Landau Hamiltonian is used with vertex functions calculated following the random phase approximation (see, e.g., Ref. 26), we numerically calculate the free energy F of polymer system consisting of Gaussian chains in a mean field environment using a path integral formalism.^{27,28} Our approach uses essentially the same free energy functional as in SCF calculations of equilibrium block copolymer morphologies by Matsen and Schick,²⁹ but complements the static SCF calculations by providing a dynamical picture of the system.

II. METHOD

Here, we shortly discuss the DDFT method^{27,30} for a bulk lamella-forming diblock copolymer melt. The diblock copolymers are modeled by a $A_{N_A}B_{N_B}$ Gaussian chain ($N = N_A + N_B$, $f_I = N_I/N$). The confined geometry is a cylindrical pore with varying diameter R . Periodic boundary conditions apply in the x -direction, along the pore. Calculations are carried out on a cubic $L_x \times L_y \times L_z$ grid with a spacing Δx that is related to the Gaussian bond length a via $a\Delta x^{-1} = 1.1543$.³¹ The spacing Δx is equal among different pore systems, making the free energy per volume element of Δx^3 easily comparable. Unless mentioned otherwise, all spacings are in units of this basic variable Δx . The pore is introduced into the simulation volume V by a masking technique; as a result the simulation volume contains both pore and mask points.³⁰ In mask points, elements of the subset $V^0 = \{\mathbf{r} = (x, y, z) \in V | \|(x, y, z) - (x_0, y_0, z_0)\| > R\}$ with $y_0 = L_y/2$, $z_0 = L_z/2$, all concentration and external potential field values are set to zero, except for the auxiliary field $\rho_M(\mathbf{r}) = 1$ that represents the mask itself.³⁰ The free energy for unconfined systems is given by (see Ref. 27 for details)

$$F[\rho] = -kT \ln \frac{\Phi^n}{n!} - \sum_I \int_V U_I(\mathbf{r}) \rho_I(\mathbf{r}) d\mathbf{r} + \frac{1}{2} \sum_{I,J} \int_{V^2} \epsilon_{IJ}(|\mathbf{r} - \mathbf{r}'|) \rho_I(\mathbf{r}) \rho_J(\mathbf{r}') d\mathbf{r} d\mathbf{r}' + \frac{\kappa}{2} \int_V \left(\sum_I \nu(\rho_I(\mathbf{r}) - \rho_I^0) \right)^2 d\mathbf{r}. \quad (1)$$

Here, k is the Boltzmann constant, T is the temperature, n is the number of polymer molecules in the volume V occupied by the system, and Φ is the intramolecular partition function for ideal polymer chains. The parameter κ determines the compressibility of the system (the dimensionless $\kappa' = \beta\kappa\nu = 20.65$, with ν the bead volume), and ρ_I^0 is the mean concentration of the I -block (where the average is taken over $V \setminus V^0$). For $\kappa \rightarrow \infty$ the system becomes incompressible. The external potentials U_I and the concentration fields ρ_I are related via the density functional.²⁸ The interchain interactions are incorporated via a mean field with interaction strength controlled by the Flory-Huggins parameters χ_{IJ} . In line with our earlier work the interactions are specified by the parameters ϵ_{IJ}^0 (in kJ/mol),^{27,28} which are directly related to the Flory-Huggins parameters by $\chi_{IJ} = 1000\epsilon_{IJ}^0/n_A kT$ (with n_A Avogadro's number and $T = 300$ the temperature in Kelvin). In case of nonzero surface interactions an extra cohesive term is added to the free energy (1) equal to³⁰

$$F^{\text{wall}} = \sum_I \int_{V^2} \epsilon_{IM}(|\mathbf{r} - \mathbf{r}'|) \rho_I(\mathbf{r}) \rho_M(\mathbf{r}') d\mathbf{r} d\mathbf{r}'. \quad (2)$$

In line with our earlier work, the interaction kernel is chosen Gaussian, and the important parameter ϵ_{IM}^0 denotes the scalar interaction strength²⁷ of bead I with the pore boundary. In the computations, $\epsilon_{BM}^0 = 0$ indicating that the B -blocks have no interaction with the wall, which is appropriate due to the fact that for an incompressible system of diblocks only the effective surface interaction $\xi = \epsilon_{AM}^0 - \epsilon_{BM}^0$ is of importance. Using this we obtain

$$F^{\text{wall}} = \sum_I \epsilon_{IM}^0 \mathcal{F}[\rho_I, \rho_M] = \xi \mathcal{F}[\rho_A, \rho_M], \quad (3)$$

where $\mathcal{F}$ is

$$\mathcal{F} = C \int_{V^2} e^{-3/2a^2(r-r')^2} \rho_A(\mathbf{r}) \rho_M(\mathbf{r}') d\mathbf{r} d\mathbf{r}' \quad (4)$$

with $C = (3/2\pi a^2)^{3/2}$ a normalization constant and a the Gaussian bond length. Since the bead-bead interaction ϵ_{AB}^0 is mostly considered a constant, we will use the notation (R, ξ) to denote points in the structural diagram.

For the simplest model, the evolution of the density fields is given by a Langevin equation³²

$$\frac{d\rho_I(\mathbf{r})}{dt} = M\Delta\mu_I(\mathbf{r}) + \eta_I(\mathbf{r}) \quad (5)$$

with M a constant mobility, Δ the Laplace operator, and η noise, distributed according to the fluctuation-dissipation theorem. Other transport coefficients, like the one for collective Rouse dynamics or reptation, exist,³² but are in general

too computationally demanding. We have recently found that our relatively simple model (5) can be appropriate to describe the experimental dynamics in detail. In Ref. 11 the experimental and calculated dynamics of cylinder- and sphere-forming diblock copolymers under an external electric field was compared, and good agreement was found based on DDFT with constant transport coefficients. In Ref. 10 the calculated dynamics of DDFT was shown to agree in full detail with scanning force microscopy (SFM) measurements of experimental dynamics in a thin film of a concentrated polystyrene-*block*-polybutadiene-*block*-polystyrene (SBS) solution. Finally, apart from the extra term in the free energy, confinement is accounted for by the boundary condition for the dynamic equations $n \cdot \nabla \mu_I = 0$, with n the normal pointing into the solid object.³⁰

III. RESULTS AND DISCUSSION

A. Nanopores: Data from literature

Since the nanopore simulation data in literature are rather scattered,^{12,13,22,23} we first present a short overview. A direct comparison of these different studies is complicated by the fact that important system parameters are different. A MC method was used to simulate the phase behavior for a A_5B_5 diblock copolymer²³ with $\varepsilon_{AB}=0.3, 1.0$, and 1.1 kT, and for a $A_{10}B_{10}$ diblock copolymer^{12,22} with $\varepsilon_{AB}=1.0$ kT. In both studies all other interactions are zero: Only the ε_{AS} (in kT), the interaction between A-blocks and the surface, is varied. In the DDFT calculations of Ref. 13, a A_8B_8 system was considered for $\varepsilon_{AB}=2.5$ kJ/mol and varying ε_{AS} (in kJ/mol) (in that article, S is actually called M or mask). Using the formula in Sec. II for $T=300$ K, we obtain $\chi=1.0$ and $\chi N=16$ (weak segregation). Using the expression from Ref. 33, $\chi \approx 5(\varepsilon/kT)$, we can recalculate the parameters used in the MC studies: $\chi N \approx 15, 50$, and 55 (weak to intermediate segregation) for Ref. 23 and $\chi N \approx 100$ in Refs. 12 and 22 (strong segregation), where it should be noted that Ref. 23 concentrates on $\chi N \approx 15$ (weak segregation). As a consequence, the results are distributed between weak, intermediate, and strong segregation regimes. In Fig. 1 we have combined the existing knowledge in schematic diagrams for different χN ; we have only differentiated between stacked disks, concentric cylinders, and alternative structures. It should be noted that the diagram for DDFT is essentially no *phase diagrams*, as the final structures are pathway dependent, and not minima of the free energy *per se*. This is a fundamental difference between static calculations aimed at deriving equilibrium morphologies, and our dynamic simulations aimed at mimicking experimental pathways, including visits to long-living metastable states. The important dimensionless spatial coordinate is the ratio of the pore diameter R and the lamellar domain distance L_0 ; this parameter is on the vertical axis. On the horizontal axis is ε_{AS} (in kT) of MC. Since the diblock is symmetric we can restrict ourselves to only positive values. We have used ε_{AS} (in kT) $\approx 1/2 \varepsilon_{AS}^0$ (in kJ/mol) for the conversion of the DDFT to the MC value.

We see that concentric cylindrical structures dominate all diagrams when ε_{AS} is large, irrespective of the value of χN . In this case, the surface field dominates and one block is

FIG. 1. (Color online) Structural diagram with existing simulation results. Different symbols denote systems in different segregation regimes: (triangles up) a A_5B_5 system with $N\chi \approx 15$, (triangles down) the same system with $N\chi \approx 50$ (Ref. 23), (circles) a A_8B_8 system with $N\chi \approx 16$ (Ref. 13), and (squares) a $A_{10}B_{10}$ system with $N\chi \approx 100$ (Ref. 22). Open symbols represent stacked disks, closed symbols concentric cylinders, and red symbols alternative structures. We have scaled the axes to make a comparison: The vertical axis shows the reduced pore radius R/L_0 , with R the pore diameter and L_0 the particular lamellar spacing in bulk, the horizontal axis $|\varepsilon|$, and the energetic interactions of A with the surface in the MC method. We have used the relation $\chi \approx 5(\varepsilon/kT)$ (Ref. 33) for the conversion of our energetic FH parameter.

energetically favored at the pore wall. For neutral pores, i.e., in the absence of a surface field ($\varepsilon_{AS}=0$), stacked disks dominate. In stacked disks the chains can adopt a packing

that is similar to the one in normal bulk lamellae. However, for the strongest segregated system stacked disk coexists with a single helix for this surface field. For small nonzero values of ϵ_{AS} also other structures can be found.

Although the data are sparse, one can conclude that structures with alternative packing dominate the diagram for large χN and relatively small ϵ_{AS} . Incommensurability cannot be the only important factor since these alternative structures are even found for commensurate pore radii $R/L_0=1$,²² although there is some uncertainty since the bulk domain distance L_0 was not determined explicitly in this study. We can only conclude from these data that especially weak surface fields in combination with strong segregation (high χN) lead to structures with alternative packing.

B. Nanopores: System choice and boundary condition

The system was chosen to model the experimental polystyrene-*block*-polybutadiene (PS-*b*-PBD) diblock copolymer in Ref. 16, which has a volume fraction of 0.56 for the butadiene block. The molecular model considered here is a $S_{10}B_{12}$ ($N_S=10$, $N_B=12$) Gaussian chain. The interaction between S and B beads is chosen as $\epsilon_{SB}=2.0$ kJ/mol or $\chi=0.8$, and consequently $\chi N=17.6$. A diblock copolymer melt of this molecular composition $f_B=0.545$ forms lamellae in bulk, and the bulk lamellar distance was determined as $L_0=8.6$ (from here onwards, all distances are in units of Δx). Commensurability issues in the x direction, along the pore axis, may arise as a result of periodic boundary conditions. Since the effects of periodic boundary conditions have been considered in the literature,^{34,35} we refer to these works for a detailed discussion. Here, we only note that the studies mentioned in the previous paragraph have not considered this effect in detail. Only for one data set in Ref. 22, where both helices and stacked disks were found and no transition between the two different structures, it was shown that the number of MC simulations required to first find a helical structure peaks at particular cylinder lengths (our L_x). General conclusions from this study are not easy to make due to the absence of regularity.

C. Nanopores: Results and discussion

We have considered pattern formation in a $S_{10}B_{12}$ diblock copolymer melt confined in a nanopore of length $L_x=32=3.72L_0$, except for the larger pores where we have used $L_x=34=3.95L_0$ and $L_x=36=4.19L_0$. In order to give a unifying description for the structural behavior of slightly asymmetric diblock copolymers in nanopores, we first construct and discuss a diagram of simulated structures. Since these structures are the result of dynamic pathways and therefore may be metastable, we consider their stability by an interpolation procedure described below. Consequently, we focus on several factors that are important for structure formation: Two types of *incommensurability* issues, due to packing frustration along the pore (associated with perpendicular structures) and perpendicular to the pore (associated with parallel structures), and the kinetic pathway, that may lead to arrested structures. In the second part, we discuss

these issues in detail, and unravel the complex interplay of these factors and the surface field.

Packing frustrations along the pore, originating from the requirement of periodicity at the two pore boundaries, have not received much attention in the past, except for the unconfined situation.³⁵ In this study, it was shown that this frustration, quantized by L_x/L_0 , determines the (in)stability of several perpendicular structures. This effect, which can be related to *elongational stress or extensional force* in an applied shear field, will be discussed in detail for our results.

In the absence of extensional forces, the analogy to similar substrates in thin films suggests that the phase behavior in pores is due to an interplay of two factors: The strength of the surface field and confinement effects.^{1,14,36} For vanishing or weak surface fields, the elastic chain deformation associated with parallel structures will be avoided, and perpendicular structures are favored instead. Due to the selective block-surface interaction, parallel structures (surface reconstructions) will be promoted for surface fields above a certain threshold field strength. However, the available space in the confined geometry dictates the degree of chain compression or extension required for parallel structures. Large chain frustrations give rise to an entropically unfavorable situation and may even prevent the formation of parallel structures. Alternative (in thin films, perpendicular) structures are then formed instead. In thin films, the surface fields were shown to be additive and affect only the first layers of structure.³⁶ Consequently the available space, quantized by R/L_0 in pores, is important for both factors. The surface field and frustration due to confinement will strongly influence structure formation in small pores. In larger pores, the surface field has a limited range, and possible structure frustration can be distributed over more layers of structure.

We note that the diblock copolymer considered here is slightly asymmetric. As a consequence, the structure diagram is not completely symmetric as well. In the absence of surface interactions, the shortest (S) part of the chain is preferentially found close to the surface due to entropic effects. The value for which energetic and entropic contributions are balanced shifts to a small positive ξ .¹⁴ Moreover, for parallel structures in the curved geometry, this chain asymmetry leads to *curvature effects*, since the spontaneous curvature associated with each of the different blocks is slightly different, and *asymmetric packing frustrations* due to the different S and B domain sizes. In contrast to thin films, the majority component of the center layer in a parallel structure is not prescribed by surface energetics. Packing frustrations are therefore expected to be less significant than in thin films, where in most cases the block next to the surfaces is prescribed by the surface fields. However, the number of domains/layers for each of the blocks can *differ*. To focus on this effect we adapt two notations: If the majority component in a parallel structure is S - B - S - B - S or B - S - B - S - B , on a line perpendicular to the pore axis from wall to wall, we denote the structure as $L_{\parallel,1(1/2)}$ (for a concentric cylindrical structure, in line with the slit notation¹⁴), and by the order of the majority component in the layers from wall to center, SBS or BSB . Note that here the total number of S and B domains differs (2 or 3), as well as the composition of the central

FIG. 5. Free energy (vertical axis) vs surface field strength ξ (horizontal axis) for the reduced radii considered: (a) 0.58, (b) 0.70, (c) 0.81, (d) 0.93, (e) 1.05, (f) 1.28, (g) 1.74, and (h) 2.21. The free energies associated with the structures in Fig. 2 are denoted by open circles (for $L_x=32$) and open squares (for $L_x=34$ or 36). Similar structures are connected by straight lines to guide the eyes. The open squares in (a)–(f) are associated with extra calculations (in all cases $L_\perp$) for different L_x : (a) 34, [(b)–(e)] 36, and (f) 34. In (c) the free energy associated with the DH structure ($\xi=-0.5$) is calculated for other ξ (open diamonds).

defected structures are remarkably stable. Based on visual inspection, and knowledge about metastable intermediate states along the pathway of formation of the perfect structures, we assign a symmetry group preceded by the letter d (for instance, dDH is a defected double helix). Other new structures will be discussed and annotated as they appear.

Focusing on the kinetic pathway, we remark that all structures in the diagram of Fig. 2 were obtained following a diffusive pathway [Eq. (5)]. Earlier work showed that microphase separation in confined systems often starts close to the pore wall.³⁷ For the pore dimensions considered here, this effect will be small, but, especially for stronger surface fields, surface reconstructions will initially form, resulting in overall coverage of the pore wall by a single component. The transition to more stable structures with a different surface coverage (for instance, helices or stacked circles) requires considerable transport of material away from the wall, into the center of the nanopore. The fundamental mechanism of this transition is important, as it may hint when and where this process may be kinetically trapped. First, we consider which of the calculated structures is an equilibrium structure. For varying ξ , the free energy (see Fig. 5) associated with a particular structure (defined by ρ_S ; due to incompressibility, ρ_B is then also fixed) is given by [see also Eq. (1)] $F^{\text{tot}}=F+F^{\text{wall}}=F[\rho_S]+\xi\mathcal{F}[\rho_S,\rho_M]$. Since both $F[\rho_S]=a$ and

$\mathcal{F}[\rho_S,\rho_M]=b$ have a constant value for a particular ρ_S (ρ_M is the mask field, and fixed by definition) the free energy values for this structure and varying surface field can be found on a line defined by $a+\xi b$. We only need two data points (or the values of a and b) for the same structure to determine this line. However, we have to be careful using this procedure as structures with the same morphology *type* may differ in detail (and therefore in the values of a and b), such as geometrical quantities and the degree of segregation. For instances, interpolation suggests that the $L_\perp$ structures for $\xi=-0.4,-0.3$ and $\xi=0.1,0.2$ (both for $R=0.70L_0$) are different, although there is no structural difference in terms of easy computable geometrical quantities. Only when we consider the bare density values we find that the maximum of the concentration field $\theta_S(\mathbf{r})$ for $\xi=-0.4,-0.3$ is slightly higher than for the equivalent structures at $\xi=0.1,0.2$.

1. Structure diagram and stability

The structure diagram is shown in Fig. 2, and is indeed not symmetric. Equilibrium structures are distinguished by a gray background; this determination is based on the interpolation approach described above. Considering the general features of this diagram, one observes that several structures in this range are metastable. For large R/L_0 , only $L_\parallel$ and $L_\perp$

TABLE I. Radial distance between S - B or block-surface interfaces (denoted by D_i , $\sum_i D_i = R$, counted from the pore surface) in parallel structures for $\xi = -1.0$ and $\xi = 0.7$. The position $\bar{R}$ of the interfaces is determined by the condition $\theta_S(\bar{R}) = \theta_B(\bar{R})$.

R/L_0	Structure	D_1	D_2	D_3	D_4	D_5
0.58	$PL(SB)$	0.5	4.5			
0.70	$PL(SB)$	1.6	4.4			
1.74	$L_{\parallel}(SBSB)$	1.5	4.9	4.1	4.5	
0.81	$L_{\parallel}(SBS)$	1.2	3.4	2.4		
0.93	$dL_{\parallel}(SBS)$	1.4	3.6	3.0		
1.05	$L_{\parallel}(SBS)$	1.3	4.2	3.5		
1.28	$L_{\parallel}(SBS)$	1.5	5.0	4.5		
2.21	$L_{\parallel}(SBSBS)$	1.5	4.8	4.1	4.8	3.8
0.58	$L_{\parallel}(BS)$	1.3	3.7			
0.70	$PL(BS)$	1.8	4.2			
1.74	$L_{\parallel}(BSBS)$	1.8	4.2	4.9	4.1	
0.81	$L_{\parallel}(BSB)$	1.4	3.1	2.5		
0.93	$L_{\parallel}(BSB)$	1.4	3.3	3.3		
1.05	$L_{\parallel}(BSB)$	1.5	3.7	3.8		
1.28	$L_{\parallel}(BSB)$	1.8	4.4	4.8		
2.21	$L_{\parallel}(BSBSB)$	1.8	4.0	4.9	4.0	4.3

are stable; for small R/L_0 also other structures can be stable. The effect of incommensurability is limited, since we do not observe outliers of perpendicular structures for the larger radii considered (where the effect of the surface field is relatively small). However, for positive ξ (pore surface likes B) and $R/L_0 = 1.28$ or 2.21 , the $L_{\parallel}$ structure is only metastable and the transition to equilibrium structures trapped. Overall, this suggests that the pore radii $R/L_0 = 1.28$ and 2.21 are incommensurate, while $R/L_0 = 1.74$ is commensurate. Moreover, we see that for negative ξ (pore surface likes S), $L_{\parallel}$ is stable for $\xi \leq -0.4$, independent of the radius R/L_0 . This asymmetry is due to the curvature effects and asymmetric packing frustration mentioned above.

Focusing on specific structures, we see that for large absolute values of ξ surface reconstructions, PL , dPL , $dL_{\parallel}$, $L_{\perp,\parallel}$, and $L_{\parallel}$, dominate. In particular, we find PL and $L_{\parallel,1}$ (SB or BS) for $R/L_0 \in \{0.58, 0.70\}$, $L_{\parallel,1(1/2)}$ (SBS or BSB) for $R/L_0 \in \{0.81, 0.93, 1.05, 1.28\}$, $L_{\parallel,2}$ ($SBSB$ or $BSBS$) for $R/L_0 = 1.74$, and $L_{\parallel,2(1/2)}$ ($SBSBS$ or $BSBSB$) for $R/L_0 = 2.21$. An analysis of domain spacing is given in Table I; the values are derived from the interface locations. Disconnected concentric cylinders $dL_{\parallel}$, where the cylinder close to the pore surface is broken up into four disconnected parallel lamellar patches, only appear for $R/L_0 = 0.93$. The distance between the patches increases with increasing ξ in this region. The $L_{\perp,\parallel}$ is the same as the structure in Fig. 8b in Feng Ref. 23, and related to $dL_{\parallel}$ (see discussion later). The number of perforations in PL can vary and show hexagonal ordering when considered in the 2D plane. Geometrically, the PL structure can be seen as an intermediate between $L_{\parallel}$ and H . Defected perforated lamellae dPL , distinguished from PL since the perforations do not show ordering on a larger scale, are found on the boundary between parallel and perpendicular structures. In response to the perforations the central cylinder sometimes adapts an oval cross section. Defected PL only appears for negative ξ (surface preference to S). The majority of surface reconstructions (PL or $L_{\parallel}$) for positive ξ (surface

preference to B) are not stable structures. They are examples of kinetically trapped structures, due to the presence of the pore surface, which gives rise to $L_{\parallel}$ related structures in the early stages of phase separation.

Perpendicular structures, tilted tacked disks $L_{\perp}^{\text{tilt}}$ and $L_{\perp}$ (four disks for all L_x considered), are found in the center of the diagram, for small ξ . For small relative radii R/L_0 , the disks are perpendicular to the pore surface, $L_{\perp}$. For larger R/L_0 there are only three disks, and they are tilted with respect to this surface, $L_{\perp}^{\text{tilt}}$. For the largest pore radii, $R/L_0 = 1.74$ and 2.21 , the pore length L_x ($L_x = 34 = 3.95L_0$ and $L_x = 36 = 4.19L_0$, respectively) is somewhat larger than the standard $L_x = 32$, and the structure is $L_{\perp}$. The $L_{\perp}^{\text{tilt}}/L_{\perp}^{\text{tilt}}$ structure are two coexisting perpendicular $L_{\perp}^{\text{tilt}}$ structures, with opposite tilt angles. The details of these mixed structures, not representing equilibrium structures for obvious reasons, will be considered later.

Helical structures, H , DH , and a new GH structure, are found at the rather broad boundary between perpendicular and parallel structures. These structures possess features relating them to both perpendicular and parallel structures: The axis of winding is parallel to the pore axis, and the wall coverage is nonuniform. The GH structure in (1.05,0.4) is related to the helical structures, but differs topologically since small helical patches are threefold connected with helical patches on the opposite side, and these connections join into two cylinders parallel to the pore axis. The GH for (1.16,0.5) and (1.40,0.4) share this property, but instead the connections form two lamellar patches and three cylinders, respectively. We call this structure gyroid-helical (GH) because of the threefold connectivity. Stable helices (left and right hand) and a double helix are formed for $R/L_0 < 1$ and negative ξ . The helical structures for positive ξ are all metastable. Metastable GH are only found for larger effective radii, $R/L_0 > 1$, adjacent to both parallel and perpendicular structures.

Several coexisting structures, $L_{\perp}/PL$, $L_{\perp}^{\text{tilt}}/H$, $L_{\perp}^{\text{tilt}}/GH$,

FIG. 6. (Color) The effect of extensional stress. Kinetics of nanopore structure formation for $R/L_0=0.70$, $\xi=-0.5$, and varying pore length L_x . (a) $L_x=32$ (stable structure: H) and (b) $L_x=36$ (stable structure: $L_\perp$). The images show isosurfaces for the S -block, for isosurface value $\bar{\phi}_S$. The numbers indicate the dimensionless time steps TMS.

$L_\perp^{\text{tilt}}/H$, and $L_\perp/L_\parallel$; and defected structures, $dL_\perp^{\text{tilt}}$, $dL_\perp$, dGH , and $dL_{\perp,\parallel}$ are found directly adjacent to their “perfect” counterparts, and stay defected after many simulation time steps (TMS). Apparently, the driving force for the removal of different types of defects in these structures is rather small, and as a result the structures are kinetically trapped. The only exceptions, coexisting H/PL and $PL/L_\parallel$ for positive ξ that are not adjacent to a PL structure, show that PL is associated with a free energy close to the one for H and $L_\parallel$, respectively. For $R/L_0=0.81$ and $-0.4 \leq \xi \leq 0.0$ the structures are very defected, and remain as such, even after a large number of extra timesteps (TMS).

2. The value of L_x/L_0 : Elongational stress and perpendicular structures

Earlier work for a “soft” confined system³⁵ concluded that the $L_\perp$, $L_\perp^{\text{tilt}}$, and H structures are related. Their stability depends on the extensional force on the system, originating either from an external field (experiments) or from boundary conditions (computations). Extensional forces may be present in experimental and computational studies dealing with structure formation in nanopores, but have not received much attention so far. The only exception is a computational study in the strong segregation limit (SSL),²⁵ where MC based on a grand canonical ensemble was employed. Especially in computational methods considering a canonical ensemble and a single L_x , this type of commensurability issues cannot always be avoided. We argue that a deeper understanding of this effect is relevant. In experiments, for instance, extensional forces play a role when the pore surface is very rough, the pore length is very small (often the case in applications considered in soft nanotechnology), and when shear fields are present, as is the case in a recently developed experimental technique for the fabrication of nanowires.³⁸ In the present study, we have considered this effect by varying L_x for a small set of selected parameters: $L_x=34=3.95L_0$ for $(0.58,-0.1)$ and $(1.28,0.1)$, and $L_x=36=4.19L_0$ for $(0.70,-0.5)$, $(0.81,0.2)$, $(0.93,0.2)$, and $(1.05,0.1)$. In all cases the microstructure evolves into $L_\perp$ (four disks), with an associated free energy that is lower than for the original structure for $L_x=32$. A detailed analysis of the structure evolution revealed that $L_\perp^{\text{tilt}}$ and H structures can also be kinetically related to $L_\perp$: In some cases, they mediate large-scale structural reorganization. This phenomenon can be observed from the helicity that appears during the evolution of the $L_\perp$ struc-

FIG. 7. Nanopore phase diagram for varying pore radius (in R/L_0) and surface field (in kJ/mol). Stable structures for each R/L_0 were determined using all free energy data in Fig. 5, irrespective of the pore lengths L_x . The absence of $L_\parallel$ for $\xi < 0$ ($R=0.58, 0.70$) and $\xi > 0$ ($R=0.70$) does not allow for the analysis of the boundaries of the PL region.

ture in a pore of $L_x=36$ and $(R/L_0, \xi)=(0.70, -0.5)$ (see Fig. 6 for the details of the formation pathway for different L_x). It is in agreement with earlier findings for a fully symmetric AB diblock in Ref. 13.

Apart from the single additional calculation for each of pores and $L_x \neq 32$, we used the $L_\perp$ structures to compute the free energy for an additional surface field strength ξ . Interpolation between these two values enables us to reconsider the stability of the structures in Fig. 2 with respect to $L_\perp$. We make two general remarks: (i) Although the interpolation technique is valuable for comparing the stability of different microstructures in pores of *equal* length (for varying ξ), in principle this procedure cannot be used to differentiate between microstructures in pores of *different* lengths. For instance, an instantaneous change of the pore length (or equal, extensional force) during the evolution gives rise to a deformation of the structure, and possibly a transition to a more stable structure. However, in contrast to SCF techniques, DDFT does not impose symmetry, and changes in free energy associated with a deformation of an existing structure, for instance, an affine deformation of a helical structure, cannot be quantized directly. Instead, additional calculations are necessary for many alternative structure types and are very time consuming in general.³⁵ We therefore anticipate on the results of the SSL study,²⁵ which considered nanopore structure formation in the absence of elongational stress. This study identified, besides lamellar structures, only stable PL . Consequently, we adapt a practical approach and use all data obtained by simulation (see Fig. 5), independent of L_x , to suggest a phase diagram of equilibrium structures (see Fig. 7), without considering the deformability of the nonlamellar structures. We note that the stability of the PL phase could not be determined indefinitely, since the alternative $L_\parallel$ phase was not formed for these small radii. Only for $R/L_0=0.58$ and positive ξ , concentric cylinders (BS) were formed, and the region where PL is equilibrium structure could be determined. (ii) The additional simulations for $L_x \neq 32$ were carried out for relatively weak surface fields, where perpendicular structures are likely to form. Interpolation indicates (see Fig. 7) that the equilibrium structure can be perpendicular even for relatively strong surface fields, for instance, due to incommensurability along the radial direction. However, the large structural rearrangements required for the transition to (stable) perpendicular structures, starting from the (unstable) parallel structures that are initially formed due to the strong

FIG. 8. (Color) The effect of extensional stress for the double helical structure ($R/L_0=0.81$, $\xi=-0.5$). Upon a variation of the pore length L_x a number of metastable structure classes [denoted by DH_i ($i=1-3$)] were identified. The images show isosurfaces for the S -block, for isosurface value $\bar{\theta}_S$. Numbers indicate the pore length L_x (in grid units).

surface interaction, may lead to arrested structures, i.e., microstructures that are kinetically trapped in metastable states. A good example is the DH structure in $(0.81, -0.5)$. Although the phase diagram of Fig. 7 shows that $L_\perp$ is the equilibrium structure, we challenged the stability of the DH structure by choosing a number of systems with different pore lengths L_x (each system was quenched from a homogeneous mixture at $TMS=0$, see Figs. 8 and 9). We never obtained the equilibrium structure as a result of the dynamic pathway. Instead, upon variation of the extensional pressure, we obtain structures with an overall parallel orientation that lack symmetry on a larger scale. Only for $L_x=32, 35$, and 38 we find an almost perfect DH structure. We rank these other structures in three different classes, labeled as DH_i . Structure $DH1$ can be seen as a highly interconnected and defected $L_{\perp,\parallel}$ structure, where the center lamella contains large holes and is connected to the other structure. Structure $DH2$ is completely symmetric; the axis of symmetry is $L_x/2$. Structure $DH3$ consists of two disconnected sheets of different structures, one similar to $PL(B)$ and the other similar to $PL(S)$. For $L_x=64$ the structure is very defected and a combination of other structures. Comparing the free energies in Fig. 9, we find that the one of the metastable DH structure ($L_x=32$) is the lowest, indicating that the extensional force is minimal for $L_x=32$ and the structure is trapped due to the diffusive kinetic pathway. The finding of DH for other L_x shows that the factor $3/L$ (with L the particular spacing of

FIG. 9. (Color) Evolution of the free energy associated with the structures in Fig. 8. Numbers indicate the pore length L_x .

the structure) plays an important role. In general, we see that an increase of the nanopore length slows down the separation dynamics. Close examination of the structures in diagram 2 reveals that the calculated structures for $-0.4 \leq \xi \leq 0.0$ are defected DH or resemble the intermediate $DH1$. In all cases, the kinetics disables the mass transport necessary for the $DH \rightarrow L_\perp$ transition.

3. The value of R/L_0 : Commensurability and parallel structures

From the phase diagram in Fig. 7 (see discussion for the derivation of this diagram above) we distinguish two specific values: $R=0.81L_0$ and $R=1.74L_0$. For $R=0.81L_0$, the pore radius $R < L_0$ and the surface field is therefore strong, but confinement effects apparently prevent the formation of parallel structures, in favor of the $L_\perp$ structure, up to large surface field strengths. Since this radius also marks the transition of SB ($0.70L_0$) to SBS (or BS to BSB), we conclude that $R=0.81L_0$ is incommensurate. For $R=1.74L_0 > L_0$, the surface field influence is much less. Apart from a symmetric $L_\parallel$ region for large surface fields (we note that the number of S and B domains is the same, independent of the wetting block), only a small region of perpendicular structures is observed for almost neutral pores, and we conclude that the pore size is commensurate. We consider the simulation results in Fig. 2 in more detail. Domain distances are shown in Table I and are inexact for the PL structure, which is not radial symmetric. For the smallest $R=0.58L_0$, parallel structures are found for strong surface fields. Most of these structures are PL [$PL(S)$ or $PL(B)$], and $L_\parallel$ is only found for $(0.58, 0.7)$. The interpolation procedure showed that, at the onset of the $L_\parallel$ region, the free energies associated with both parallel phases are relatively close (the value of b is almost equal, see discussion of the interpolation procedure before), which explains that $L_\parallel$ is found for $\xi=0.7$ due to kinetic trapping. Upon comparing the PL and $L_\parallel$ structures geometrically, we see that the formation of necks on the central cylinder in PL leads to an increased curvature of the S - B interface and contact area of the nonpreferred block with the pore surface. If we compare the position of this S - B interface for this incommensurate and the commensurate situation ($R/L_0=1.74$, S - B interface closest to the pore center, see Table I) we observe a reduced thickness for positive ξ and equal thickness for negative ξ . This illustrates the curvature effect mentioned before, meaning that the elastic chain deformation associated with the formation of $L_\parallel$ can much easier be facilitated when the interface is curved towards the shortest S -part of the chain, than to longest B -part. The same phenomenon can be observed for $R=0.70L_0$. Here, concentric cylinders ($L_\parallel$) are completely absent in the simulated range. Alternative PL structures, very similar to the ones for $R=0.58L_0$ apart from the number of perforations, can be found, but equilibrium $PL(S)$ structures are only found for rather low ξ values, again due to the curvature effect. In order to fill the pore, the domains are rather extended (Table I). The formation of parallel structures requires strong stretching of the part of the chain that contains the wetting blocks, giving rise to an entropic penalty that is only counterbalanced for stronger surface repulsion. For $R=0.81L_0$

FIG. 10. (Color) Kinetics of nanopore structure formation for selected pore radii and surface field strength (in kJ/mol): (a) $R/L_0=0.70$, $\xi=-0.4$, (b) $R/L_0=0.70$, $\xi=0.1$, (c) $R/L_0=0.81$, $\xi=-0.5$, (d) $R/L_0=0.93$, $\xi=-0.1$, (e) $R/L_0=1.28$, $\xi=0.2$, and (f) $R/L_0=1.74$, $\xi=0.3$. The images show isosurfaces for the S -block, for isosurface value $\bar{\phi}_S$. The numbers indicate the dimensionless time steps TMS.

surface reconstructions $L_{\parallel,1(1/2)}$ (SBS or BSB) are found and PL is absent, apart from the coexisting H/PL structure ($\xi=0.5$). However, from the phase diagram, we conclude that all surface reconstructions $L_{\parallel}$ are metastable and formed due to kinetic factors. The analysis of the domain distances in Table I shows rather small domain sizes, and therefore strong compression of the chains. Although $L_{\parallel,1(1/2)}$ is only found for positive ξ ($R=0.93L_0$), we observe many other parallel structures, suggesting that this radius R is somewhat commensurate. However, from the analysis in Table I we see that the domain spacing for this radius is still rather small (compared to bulk spacing, and the spacing of $D_3+D_4=9.0$ found for the inner two layers in the commensurate $BSBS$ structure for $\xi=0.7$ and $R/L_0=1.74$), especially towards the center. Moreover, the equilibrium structure in this range is $L_{\perp}$ (see Fig. 7). We focus on the disconnected lamellar patches ($dL_{\parallel}$) for $\xi \leq -0.2$. Because of the masking procedure, the cross section of the pore is not completely circular, and the curvature of the pore wall varies. Hence, one could expect that metastable concentric cylinders formed due to kinetic factors, and that the origin of these disconnections, appearing in regions of high curvature, is the chain-stretching required for the formation of undefected $L_{\parallel}$. Close examination of the

kinetic pathway towards the $L_{\perp,\parallel}$ structure [Fig. 10(d)] shows $dL_{\parallel}$ as an intermediate structure. In particular, along the pathway two of the lamellar patches and the central cylinder have merged to form the center lamellae. In the early stage, undulations lead to connections between the different structural elements in $dL_{\parallel}$ (very fast). Subsequently, these connections merge (and others are removed) by a complex process that includes the formation of holes (very slow). The structures along this pathway resemble the metastable $DH1$ structures that are found in $(0.81, -0.5)$ for varying L_x . Finally the $L_{\perp,\parallel}$ structure forms. Apparently the removal of defects depends on the surface field strength and is slowed down considerably by an increase of ξ from -0.1 to 0.0 . In MC calculations²³ the $L_{\perp,\parallel}$ structure was also found for a pore radius $R \approx L_0$. It was suggested that it represents an intermediate between the $L_{\parallel}$ and $L_{\perp}$ structures, but no further explanation was given. The stability analysis suggests that all non-perpendicular structures are metastable. We claim that this structure represents a kinetically arrested alternative for stable $L_{\perp}$. Both structures are formed for small to vanishing surface fields, in order to avoid the elastic chain deformation due to the curved S - B interfaces in concentric cylinders. Following the same reasoning, the $dL_{\parallel}$ structures (instead of $L_{\parallel}$) for $\xi < -0.1$ are metastable structures along the pathway to this $L_{\perp,\parallel}$ structure, and kinetically trapped since the undulations necessary for the formation of this structure are suppressed by the strong surface field.

For $R=1.05L_0$ the situation is weakly incommensurate, and $L_{\perp}$ is the equilibrium structure between $-0.4 \leq \xi \leq 0.8$. This can also be seen from the slightly perturbed spacing in Table I for $\xi=-1$. For $-0.5 \leq \xi \leq -0.3$ in Fig. 2 (dPL) holes are formed in the region of highest curvature, and we find that the density field in the center of the pore is slightly higher for $\xi=-0.5$ than for $\xi=-1$ (no holes). This dPL structure was also observed as the initial stage in the formation of DH , H , and $L_{\perp}$ structures for these ξ and smaller R , and for the slow transition to $dL_{\perp}^{\text{tilt}}$ for $\xi=-0.2, -0.1$, where we simulated up to 40 000 TMS. For positive ξ , the parallel structures are also metastable, but the free energy associated with the $PL(B)/L_{\parallel}$ coexisting structure ($\xi=0.6$) is slightly below the one extrapolated for perfect $L_{\parallel}$. This is once more an indication that the chain deformation associated with concentric cylinders can be relieved by the formation of necks, and therefore PL is more stable. Moreover, this transition from $L_{\parallel}$ to PL (and eventually the transition, via other intermediate structures, to $L_{\perp}$) is obviously very slow. The parallel structure $L_{\perp,\parallel}$ is absent, showing that the kinetic pathway leading to this metastable structure is rather unique.

For $R=1.28L_0$, the phase diagram suggests that the pore size is slightly too large for $L_{\parallel,1(1/2)}$. The detailed analysis in Table I shows that the chains in the center of the pore are stretched, compared to the bulk domain thicknesses ($D_B=4.7$ and $D_S=3.9$), and to the $L_{\parallel,1(1/2)}$ structures for other R . The boundary for the equilibrium $L_{\perp}$ phase shifts to stronger surface fields, but this shift is small compared to $R=1.05L_0$. The larger shift for positive ξ is due to the curvature of the S - B interface of the central cylinder, which is curved towards the shortest block for negative ξ .

For $R=1.74L_0$ surface reconstructions $L_{\parallel,2}$ are found for

positive and negative ξ . Moreover, only at the boundary with defected structures, these surface reconstructions are metastable. The stability region of $L_{\parallel}$ is almost symmetric around $\xi=0.1$, and we conclude that the situation is commensurate. The chains are slightly stretched (see Table I) compared to the spacing to the domain spacing in the bulk due to curvature effects. A comparison of the spacings in pores for $R/L_0=1.74$ and 2.21 suggests that especially the chains in the center of the pore are compressed for the largest pore. The driving force for the transition of the metastable surface reconstructions to $L_{\perp}$ is rather small (see Fig. 5).

4. Detailed kinetic pathways

Although some kinetic pathways have been considered in the previous subsections, we focus on a few more representative pathways here. For $R=0.70L_0$, different kinetic pathways were found leading to the final $L_{\perp}$ structures for $\xi=-0.4$ [Fig. 10(a)] and $\xi=0.1$ [Fig. 10(b)]. For the negative $\xi=-0.4$ the initial structure is $L_{\parallel}$. At later stages (200 TMS) the cylinder in the center of the pore has broken up to form short and disconnected cylinders, while part of the outer curved lamellae stays intact. Remarkably, parts of this outer curved lamellae remain intact during the transition and mediate the mass transfer from the wall into the pores interior. This is the source of the toroidal structures that appear as intermediate, at the positions where the stacked disks are later formed. For the positive $\xi=0.1$ the initial structure is bicontinuous [or very defected $PL(S)$] with many threefold connections in the center of the pore. These connections can be seen as defects, as very early in the evolution there is an excess of material at positions where stacked disks are later formed. Fourfold connections (small lamellar patches) and helical domains mediate the mass transfer needed for the formation of perfect stacked disks in this case.

The most interesting structure for $R=0.81L_0$ is the double helix (DH) for $\xi=-0.5$. Figure 10(c) shows that the formation of this structure is a rather slow process, and starts from $L_{\parallel}$ (100 TMS). In the early stages, equidistant perforations form in the outer curved lamellae, similar to the ones formed for $R/L_0=1.05$ and $-0.4 \leq \xi \leq -0.3$. These perforations grow in time while the cylinder in the center of the pore undulates and breaks up to form a bicontinuous network. In contrast to before this bicontinuous network itself forms the nucleus of the final structure, which is the double helix structure. Due to its helical nature, defect removal cannot be mediated by the formation and breakage of helical connections and thus the process is rather slow.

The metastable parallel structure GH is only formed for positive ξ , and either stabilized by an extensional force or kinetically trapped along the pathway of the $L_{\parallel} \rightarrow L_{\perp}$ transition. Its finding adjacent to perpendicular structures (deformed $L_{\perp}$) and $L_{\parallel}$ for $R=1.28L_0$, or perpendicular structures and PL , coexisting with $L_{\parallel}$ and H , for smaller pores suggests that it may evolve from PL . Close examination of the pathway for $R/L_0=1.28$ ($\xi=0.4$) and $R/L_0=0.93$ ($\xi=0.3$) shows that the initial structure $L_{\parallel}$ rapidly transforms (after 100 TMS) into a very defected structure, where the perforations in the outer B layer are uneven and randomly positioned.

Since PL only appears in coexistence with other structures for positive ξ , we conclude that the kinetic pathways for the formation of GH and PL must be different.

For $R=1.28L_0$, we concentrate on the effect of elongational stress on the evolution of the $L_{\perp}^{\text{tilt}}/L_{\perp}^{\text{tilt}}$ structure, where the equilibrium structure is $L_{\perp}$. In particular, we consider the structure evolution for $\xi=0.2$ in detail, since incommensurability issues with respect to the pore radius are absent. First, we note that the cylinder in the center of the pore for $\xi=-0.2$ shows some undulations, but the number of pores in the outer layer remains constant for longer simulation times, and no connections with the inner structures are formed within the limits of simulation time. These connections are required for the transition/nucleation of one phase (in)to the other. For $\xi=-0.1$ a defected structure without apparent symmetry at TMS=10 000 evolves into a helical phase at TMS=25 000. Since the axis of winding is away from the center, we conclude that the structure can be seen as a majority $L_{\perp}^{\text{tilt}}$ under some angle (with the pore wall), connected to a minority $L_{\perp}^{\text{tilt}}$ under a reverse angle. As a natural consequence, the connections between these two partial structures have a helical nature. We observe similar structures for $\xi=0.1$ and $\xi=0.3$. For $\xi=0.2$ the axis of winding coincides with the center of the pore, and we could classify this structure as analogous to the DH structure for smaller R . However, in line with the other structures for this radius this could be seen as a merger of two equally sized clusters of oppositely tilted and connected stacked disk morphologies. Moreover, in the evolution for $\xi=0.2$ [see Fig. 10(e)] we observe a fundamental mechanism: The orientation with respect to the pore wall of the total structure, as well as the change of chirality of local structure, are mediated by melting and reconnection of helical sections. In this way, unfavorable orientation and winding in part of the structure (for instance, giving rise to holes or increased curvature) can be avoided. We conclude that all these structures $-0.1 < \xi < 0.3$ are actually coexisting $L_{\perp}^{\text{tilt}}$ under two different angles. The helical connections between these two cluster structures are relatively stable in time, and analogous of the double periodic array of saddle surfaces found in lamella-forming thin films under an electric field.³⁹ Apparently, they do not cost much energy to the system. This is confirmed by the free energy plot in Fig. 5, where the free energies associated with these different helical structures fall almost on a single line.

To see whether the elongational stress is the cause for this fundamental mechanism, we also considered the structural evolution in the absence of incommensurability along the pore, $\xi=0.3$ and $R=1.74L_0$ [see Fig. 10(f)]. The detailed analysis confirms that the fundamental mechanism identified for formation of structure under elongational stress, i.e., reorientation on a higher structural level by helical connections, is also instrumental for the formation of equilibrium $L_{\perp}$ structures. One can clearly see that the tilted and interconnected disks that form in the initial stages (1000 TMS) slowly transform into perfectly stacked disks or $L_{\perp}$ (25 000 TMS). The mechanism is a moving front of helical connections, just like shown in Fig. 10(e). This effect can also be observed for $R=2.21L_0$ and $\xi=0.0$ or 0.2 (not shown), where the initial coexisting structure $L_{\perp}^{\text{tilt}}/L_{\perp}^{\text{tilt}}$ (TMS=10 000)

evolves into $L_{\perp}$ (TMS=20 000). From these observations we conclude that coexisting stacked tilted disks with helical domains are the preferred transient structure, independent of the elongational stress. The most probable cause is that interconnected tilted structures can much easier accommodate the necessary large-scale structural rearrangements that are necessary for the formation of $L_{\perp}$. Helical connections play an important role in the removal of defects, and slowly change the tilt of the intermediate structure, leading to the formation of equilibrium $L_{\perp}$ structures.

D. Discussion: Comparison to other studies

1. Experiments

Up to now, experimental groups have considered structure formation of (nearly) symmetric PS-*b*-PB (Refs. 16–19) and PS-*b*-PMMA (Ref. 20) confined in nanopores. Almost all observed experimental microstructures belong to the class of concentric cylinders. This does not come as a surprise, as for most diblock copolymers one of the blocks has a higher affinity for aluminum. We have previously concluded that, especially for the larger pore sizes considered, surface fields above a certain (relatively low) threshold value lead to so-called surface reconstructions: Concentric cylinders. In the experimental procedure, the block copolymers were introduced into the alumina nanopores at elevated temperatures via capillary action and subsequently cooled. After this procedure a weak base was used to dissolve the alumina and produced the freestanding rods of block copolymer that were imaged. The thus obtained rods show several features: The cross sections vary in shape (from ellipsoidal to spherical) and the thickness of the rod varies substantially along the rods (due to the roughness present in the alumina pores). As a result, the actual pore radius can only be determined approximately. Although some measurements of surface tension were performed, exact numbers quantifying the affinity of PS, PB, and PMMA for the aluminium pore material are missing.

We conclude that the stacked-disk or toroidal-type structure in nearly symmetric PS-*b*-PB block copolymer¹⁶ is the best candidate for a true comparison between our calculations and experiments. We therefore choose the simulation parameters as close as possible to the values for this experimental system. The alternative structure was found at a temperature of $T=398.15$ K, above the glass transition temperature,^{17,19} in a system with a molecular weight of $M=18\,400$ (the number of monomers $N\sim 250$). The experimentally measured repeat period $L_0=17.6$ nm, giving rise to an incommensurate pore radius $R/L_0\sim 1.3$. We note that the effective Flory-Huggins (FH) parameter χ depends to some extent on the details of the experimental system. Using earlier measurements by the same group we find $\chi=0.058$ (Ref. 40) and obtain $\chi N\approx 14.5$. For a coarse-grained chain with $N=22$ ($A_{10}B_{12}$) we therefore find an effective Flory-Huggins parameter $\chi_{SB}=0.66$. Here we consider $\chi_{SB}=0.6$ instead, and focus on the influence of reduced FH parameter in general (the calculations in the previous sections were carried out for $\chi_{SB}=0.8$). Although we use the bare FH parameters in the

FIG. 11. (Color) Nanopore structures for $S_{10}B_{12}$ ($\chi=0.6$, $L_x=32$), matching the experimental system of Shin *et al.* (Ref. 16), (a) for pore radii $R=19$ ($L_x=36$), (b) $R=9$ ($L_x=32$), and (c) surface field $-1.0\leq\xi\leq 0.7$ (in kJ/mol). Parallel structures for surface fields higher than the threshold surface field strength are not shown. The structures in (c) are for constant surface field ($\xi=0.4$) and pore radius varying between 5 and 11 ($L_x=32$). For each radius, R , structures from two different viewing angles are shown. The images show isosurfaces for the S-block, for isosurface value θ_S .

calculations, the referring $\epsilon_{SB}^0=1.5$ kJ/mol for the choice of the temperature ($T=300$ K) in the standard input in DDFT.

The results of calculations for $\chi_{SB}=0.6$ and two radii R (and varying ξ) are shown in Figs. 11(a) and 11(b). We conclude that for the largest $R=19$ the effect of reduced χ is rather small. Comparing the results for $R=19$ in Fig. 11 and $R=2.21L_0$ in Fig. 4 (for the same cylinder length L_x) we see that the phase boundaries have not been shifted much. For low and high values of ξ concentric cylinders are found with an additional layer when compared to the results for higher block-block interaction. This finding is in well agreement with the 2D SCF calculations of Li *et al.*²¹ where a transition from L_3 to L_2 (notation of Ref. 21) is found for $\chi N>13$. The tilt of the $L_{\perp}$ for small ξ reflects the different bulk lamellar distance L_0 associated with the change in χ_{SB} and is due to an extensional force. For the smallest $R=9$, concentric cylinders ($L_{\parallel}$) are found for $\xi\leq -0.1$ and $\xi\geq 0.5$, with the same number of layers as for larger χ_{SB} . For $0.1\leq\xi\leq 0.3$, a perfect $L_{\perp}$ structure is obtained. Constructing the lines associated with different structures shows that $\xi=0.0$ and $\xi=0.4$ are indeed located on intersections of different lines. Based on the experimental knowledge that the B-block is preferentially found close to the pore surface, we further concentrate on $\xi=0.4$. For $R=19$ the structure contains perforations, which will not be visible in the experimental imaging procedure. This structure will appear as concentric cylinders ($L_{\parallel}$). For $R=9$, the structure is a mixture of $L_{\parallel}$ and $L_{\perp}$. The standard element of this structure are two opposite small lamellar patches oriented parallel to the pore surface. Along the pore, only the orientation of the elements changes and neighboring elements are orthogonal [see Fig. 12(a)]. The fourfold connections (instead of three) between the elements are short

FIG. 12. (Color) (a) High isodensity image of the cage-like structure for $R=9$ and $\xi=0.4$. (b) Structures for the system of Fig. 8 in a rough pore and $\xi=0.4$. The pore radius varies between $R=10$ (center region) and $R=9$ (outer regions). From left to right: Isosurface for the S -block (isosurface value $\bar{\theta}_S$) after 1000 TMS, the same for 20 000 TMS, isosurface of the B -block (isosurface value $\bar{\theta}_B$) after 20 000 TMS, crop of the isosurface at 20 000 TMS combined with an orthogonal slice of the B field in the center of the pore.

and curved cylinders. This mixed structure has interesting features that somewhat resemble the experimental toroidal-type structure of Shin *et al.*¹⁶ However, the structures still do not match in all details. Additionally, we varied the small pore radius for fixed $\xi=0.4$ [Fig. 11(c)]. This leads to $L_{\parallel}$ for $R=5, 8$ and $L_{\perp}$ for $R=6, 7, 10$, and 11 . In general, we conclude that the decrease of χN relieves some of the stresses present in the stronger phase segregated system, and leads to faster relaxation towards a stable structure.

Finally, the small scale of the experimental nanopores leads to constraints that are not effective in other systems. The results of the previous section show that the structure type can be very sensitive to the pore radius. Small variations of the pore radius (due to roughness) along the pore at relatively larger length scales may therefore affect the structure formation to some extent. This is, in particular, the case in small nanopores, where slightly different pore radii will promote different structure types and connectivity issues (of different types and periodicity) may appear. To study this effect, we prepared a pore with two pore radii, $R=9$ and 10 ($\xi=0.4$), equally distributed along the pore (the thickest part is located in the middle of the pore). Using $L_0=8.6$, the bulk characteristic lamellar spacing determined for the system with higher χ_{SB} , the relative pore sizes R/L_0 vary between 1.05 and 1.16 . Keeping in mind that L_0 will decrease for decreasing χ_{SB} , this range matches the experimental value of 1.3 reasonably well. Following the evolution of this structure [see Fig. 12(b)], we initially observe that simply the stable structures for the individual pore radii are formed and connect ($L_{\parallel}$ for $R=10$ and the cage structure for $R=9$). However, eventually the interface between the different structures acts as a driving force for a transition to another structure. The final and stable structure for the S -block is completely perpendicular, with stacked disk for $R=10$, but further away from this region the stacked disks become perforated and adapt the toroidal shape [see Fig. 12(b), second from left]. From a comparison between the experimental image and the simulated 3D structure we observe that cross section of the B -block in the center of the pore [Fig. 12(b), utter right] looks very similar to Figs. 1(C) and 1(D) in Ref. 16. Since

the structure for larger R and the same parameters is $L_{\parallel}$ we conclude that this computational system gives a very close match.

2. Other numerical studies

The differences in methodology and choices of the parameter range complicate the comparison to results of other computational studies in general. Although most of these studies were not aimed at modeling experimental systems, we note that none of them have reproduced the experimental toroidal-type structure.¹⁶ Moreover, the dynamics in our method is aimed at following the experimental pathway. Here, we focus on a qualitative comparison to the two most extensive MC studies for symmetric diblock copolymers in nanopores of three different radii: Feng and Ruckenstein²³ and Chen *et al.*²²

In Ref. 23 χN is relatively low and stacked disks ($L_{\perp}$) are found for neutral pores, as expected. The region where $L_{\perp}$ is stable shrinks for increasing pore radius. For large surface interactions, concentric cylinders ($L_{\parallel}$) are found. The only alternative structures for intermediate surface fields are PL ($R/L_0 \approx 0.8$) and $L_{\perp, \parallel}$ ($R/L_0 \approx 1.0$). These results are fully in line with the findings in this article (see Sec. III). Moreover, for $R/L_0 \approx 1.0$ and an increased χN , also helices and double helices were obtained for varying surface field strengths. We also find these structures but for somewhat smaller R . Previously, we identified these structures as kinetically trapped or stabilized by elongational forces. The particular surface field strength and pore radius at which this effect plays an important role may be expected to depend on the strength of the block-block interface, i.e., the value of χN . Moreover, also the slight asymmetry of our own block copolymer may lead to a shift of the parameter space where this effect is present.

In Ref. 22, the value of χN is considerably higher and the pore radii very small. For small to zero surface fields several structures are found to be stable: Stacked disks $L_{\perp}$, helices H , and perforated (concentric) cylinders PL . Strangely enough tilted stacked disks are absent, which may have to do with the absence of extensional stresses for the considered pore length. Previously (previous section) we found that for small radii and our choice of χN the free energy difference between the helical and perforated lamellar structures can be small. Our results also suggested that the helical structure can be formed via the merging of perforations in the perforated structure, and subsequently that PL is an intermediate structure in the $D \rightarrow PL \rightarrow H$ evolution. The fact that here PL is found for small surface fields is therefore not completely unexpected. For large surface fields, $L_{\parallel}$ is stable. The only true discrepancy between this study and ours is the stacked circle structure found for $D=26$ (Ref. 22) close to the stability region of the $L_{\parallel}$ structure. This structure, similar to $L_{\parallel}$ but with short bounded instead of an infinite cylinder in the pore center, could originate from packing problems in combination with a relatively large surface field. Although the bulk distance L_0 was not determined explicitly, we can take $L_0=16$ from counting the number of stacked disks in the cylinder, and find $R/L_0=0.81$. We refrain from further specu-

lations about the origin of this structure, and conclude that it is either an exotic structure or due to the higher χN .

Finally, we briefly consider the results of a nanopore study that employed dissipative particle dynamics (DPD). In Ref. 41 a A_5B_5 block copolymer system is simulated, both in neutral pores and pores with surface fields. The pore wall is not included as a geometrical constraint, but as soft core DPD beads with strong repulsion. The effect of extensional forces is not considered. The authors find that $L_\perp$ always forms in neutral pores, except for the smallest pore radius considered ($R=7$) (Ref. 41) where a double helix DH was found. For pores with strong surface fields only $L_\parallel$ forms, except for the smallest radius where $L_\perp$ is obtained. For surface field strength between these two extrema, several structures are obtained and hopping between different states of the system is observed. Although not mentioned as such, the structures in Fig. 5 of (Ref. 41) show that they can be arrested in alternative morphologies. In particular, helical structures were found to serve as intermediates. Although the parameter range in this study is different and limited, it confirms our findings that the system can rather easily adapt to incommensurability in larger nanopores. Moreover, for the intermediate parameter range (small, nonzero surface fields) there is a delicate balance between entropic and energetic contributions to the free energy, and structural transitions are mediated by helicity. In our more systematic study we show that structures can sometimes be arrested in these transition (helical) states. Moreover, we showed that for very small pore radii perpendicular structures can be found for relatively large surface fields.

IV. CONCLUSIONS

We have used a DDFT method to study the behavior of a slightly asymmetric diblock copolymer $S_{10}B_{12}$ confined in a cylindrical nanopore. This block copolymer forms lamellae in the bulk, and was chosen to model the experimental system in Ref. 17. We find many structures that were also identified in nanopore studies for completely symmetric diblock copolymers, but also a few new structures. However, almost all exotic structures are metastable. Our aim here is to identify general rules for structure formation in nanopores. We have focused on the intermediate surface strength in the parameter space, where the effective surface field, determined by the difference of the two block-surface interactions ξ , is relatively small. Outside this region, curved concentric cylinders (in analogy with the thin film notation denoted by $L_\parallel$) are stable based on energetic arguments. The sign of the surface field determines the block that is preferentially found to wet the pore surface: For positive ξ the B -block is found close to the pore surface, and for negative ξ the S -block is preferential close to this surface. Moreover, for a small and vanishing surface field the elastic chain deformations present in the concentric cylinders are avoided by the formation of a stacked disk structure (denoted as $L_\perp$). The transition point from parallel structures ($L_\parallel$) to perpendicular structures ($L_\perp$) is located somewhere inside this region, and we find many alternative structures. Analysis of the free energy and the determination of a phase diagram of equilibrium structures

shows that all of these structures except $L_\parallel$, PL , and $L_\perp$ are metastable, and we have considered possible causes in detail. For stronger surface fields, that in general favor surface reconstructions (parallel structures), commensurability of the pore radius R and the natural lamellar domain distance L_0 is important to avoid large compression or extension of the chains in parallel structures. For weaker surface fields, favoring perpendicular structures, commensurability of the pore length L_x , and the natural lamellar domain distance L_0 is important to avoid large extensional forces. The commensurability issues, together with sometimes extremely slow transition dynamics, were found to cause the formation of alternative structures.

For stronger surface fields, phase separation is initiated by the presence of the interacting pore surface. As a result, the pathway from the homogeneous melt to more stable alternative structures includes a first stage of parallel structures or surface reconstructions, even when $L_\parallel$ is not equilibrium. Subsequently, when the driving force is relatively large, holes are formed (for larger R , due to undulations that form connections between different layers) and connect: A helical structure is formed at a larger scale. The most probable origin of this helicity is that helical structures can more easily accommodate structural reformations at a higher level by adapting their spacing and pitch than defected $L_\perp$ structures. At the final stages, the helical structures line up by defect movement: The formation and destruction of threefold connections. We find that these helical structures cost few energy to the system. Along this pathway, the structure can get kinetically trapped depending on the frustration that it experiences. We find metastable $L_\parallel$ structures with holes (denoted by PL , and similar to the mesh or catenoid structures in Refs. 23 and 22, respectively), helices, or double helices for decreasing surface field strength. In general, the transformation of the intermediate structures to other structures can be slowed down, or even be halted, by the presence of extensional forces due to incommensurability along the pore.

For weaker surface fields the influence of the confinement on the initial stages is less substantial. As a result the initial structure is very interconnected and defected. The mechanism for the removal of these defects, however, is very similar to the one described above: Small helical domains and threefold connections play an important role. Here, the extensional forces play a major role, and determine whether stacked disks, tilted stacked disks, or helical-like structures (which are actually clusters of tilted stacked disks under opposite angles, connected via helical domains) are formed.

We have also considered the experimental system of Shin *et al.*,¹⁶ in which a new toroidal-type structure was observed. Our results suggest that for weaker surface fields and small pore radius the structures are very sensitive to pore radius variations. Moreover, these pore radius variations give rise to interfaces between the stable structures for constant pore radii that exist in parts of the pore. These interfaces act as the driving force for the transition to other structures. For a certain surface field strength ($\xi=0.4$) and constant pore radius, we find a cagelike structure for a pore radius close to the experimental value, and perforated concentric cylinders (that appear as concentric cylinders in the experimental im-

aging procedure) for large pore radii. Moreover, by including the roughness of the experimental pore into our calculation in the form of a step profile, we were able to obtain a 3D structure that is very similar to the experimental structure.

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- ¹Q. Wang, in *Nanostructured Soft Matter: Experiment, Theory, Simulation and Perspectives*, edited by A. V. Zvelindovsky (Springer, Dordrecht, 2007), pp. 498–528.
- ²F. S. Bates and G. H. Fredrickson, *Phys. Today* **52**(2), 32 (1999).
- ³M. W. Matsen, *Curr. Opin. Colloid Interface Sci.* **3**, 40 (1998).
- ⁴K. Binder, *Adv. Polym. Sci.* **138**, 1 (1999).
- ⁵M. J. Fasolka and A. M. Mayes, *Annu. Rev. Mater. Res.* **31**, 323 (2001).
- ⁶T. Hashimoto, in *Nanostructured Soft Matter: Experiment, Theory, Simulation and Perspectives*, edited by A. V. Zvelindovsky (Springer, Dordrecht, 2007), pp. 45–98.
- ⁷L. H. Radzilowski, B. L. Carvalho, and E. L. Thomas, *J. Polym. Sci., Part B: Polym. Phys.* **34**, 3081 (1996).
- ⁸H. P. Huinink, J. C. M. Brokken-Zijp, M. A. van Dijk, and G. J. A. Sevink, *J. Chem. Phys.* **112**, 2452 (2000).
- ⁹Q. Wang, P. F. Nealy, and J. J. de Pablo, *Macromolecules* **34**, 3458 (2001).
- ¹⁰A. Knoll, K. S. Lyakhova, A. Horvat, G. Krausch, G. J. A. Sevink, A. V. Zvelindovsky, and R. Magerle, *Nat. Mater.* **3**, 886 (2004).
- ¹¹T. Xu, A. V. Zvelindovsky, G. J. A. Sevink, K. S. Lyakhova, H. Jinnai, and T. P. Russell, *Macromolecules* **38**, 10788 (2005).
- ¹²X. He, M. Song, H. Liang, and C. Pan, *J. Chem. Phys.* **114**, 10510 (2001).
- ¹³G. J. A. Sevink, A. V. Zvelindovsky, J. G. E. M. Fraaije, and H. P. Huinink, *J. Chem. Phys.* **115**, 8226 (2001).
- ¹⁴A. Knoll, A. Horvat, K. S. Lyakhova, G. Krausch, G. J. A. Sevink, A. V. Zvelindovsky, and R. Magerle, *Phys. Rev. Lett.* **89**, 035501 (2002).
- ¹⁵M. J. Fasolka, P. Banerjee, A. M. Mayes, G. Pickett, and A. C. Balazs, *Macromolecules* **33**, 5702 (2000).
- ¹⁶K. Shin, H. Q. Xiang, S. I. Moon, T. Kim, T. J. McCarthy, and T. P. Russell, *Science* **306**, 76 (2004).
- ¹⁷H. Q. Xiang, K. Shin, T. Kim, S. I. Moon, T. J. McCarthy, and T. P. Russell, *Macromolecules* **37**, 5660 (2004).
- ¹⁸Y. Y. Wu, G. S. Cheng, K. Katsov, S. W. Sides, J. F. Wang, J. Tang, G. H. Fredrickson, M. Moskovits, and G. D. Stucky, *Nat. Mater.* **3**, 816 (2004).
- ¹⁹H. Q. Xiang, K. Shin, T. Kim, S. I. Moon, T. J. McCarthy, and T. P. Russell, *J. Polym. Sci., Part B: Polym. Phys.* **43**, 3377 (2005).
- ²⁰Y. Sun, M. Steinhart, D. Zschech, R. Adhikari, G. H. Michler, and U. Gösele, *Macromol. Rapid Commun.* **26**, 369 (2005).
- ²¹W. Li, R. A. Wickham, and R. A. Garbary, *Macromolecules* **39**, 806 (2006).
- ²²P. Chen, X. He, and H. Liang, *J. Chem. Phys.* **124**, 104906 (2006).
- ²³J. Feng and E. Ruckenstein, *Macromolecules* **39**, 4899 (2006).
- ²⁴J. Feng and E. Ruckenstein, *J. Chem. Phys.* **125**, 164911 (2006).
- ²⁵Q. Wang, *J. Chem. Phys.* **126**, 024903 (2007).
- ²⁶I. W. Hamley, *The Physics of Block Copolymers* (Oxford University Press, Oxford, 1998).
- ²⁷J. G. E. M. Fraaije, B. A. C. van Vlimmeren, N. M. Maurits, M. Postma, O. A. Evers, C. Hoffmann, P. Altevogt, and G. Goldbeck-Wood, *J. Chem. Phys.* **106**, 4260 (1996).
- ²⁸B. A. C. van Vlimmeren, N. M. Maurits, A. V. Zvelindovsky, G. J. A. Sevink, and J. G. E. M. Fraaije, *Macromolecules* **32**, 646 (1999).
- ²⁹M. W. Matsen, M. Schick, *Phys. Rev. Lett.* **72**, 2660 (1994).
- ³⁰G. J. A. Sevink, A. V. Zvelindovsky, B. A. C. van Vlimmeren, N. M. Maurits, and J. G. E. M. Fraaije, *J. Chem. Phys.* **110**, 2250 (1999).
- ³¹N. M. Maurits, J. G. E. M. Fraaije, P. Altevogt, and O. A. Evers, *Comput. Theor. Polym. Sci.* **6**, 1 (1996).
- ³²N. M. Maurits and J. G. E. M. Fraaije, *J. Chem. Phys.* **107**, 5879 (1997).
- ³³J. J. Freire and C. McBride, *Macromol. Theory Simul.* **12**, 237 (2003).
- ³⁴G. Wang, G. Yan, P. F. Naeley, and J. J. de Pablo, *J. Chem. Phys.* **112**, 450 (2000).
- ³⁵H. Morita, T. Kawakatsu, M. Doi, D. Yamaguchi, M. Takenaka, and T. Hashimoto, *J. Phys. Soc. Jpn.* **73**, 1371 (2004).
- ³⁶H. P. Huinink, J. C. M. Brokken-Zijp, M. A. van Dijk, and G. J. A. Sevink, *Macromolecules* **34**, 5325 (2001).
- ³⁷K. S. Lyakhova, A. V. Zvelindovsky, and G. J. A. Sevink, *Macromolecules* **39**, 3024 (2006).
- ³⁸M. L. Ma, V. Krikorian, J. H. Yu, E. L. Thomas, and G. C. Rutledge, *Nano Lett.* **6**, 2969 (2006).
- ³⁹A. V. Kyrilyuk and J. G. E. M. Fraaije, *J. Chem. Phys.* **125**, 164716 (2006).
- ⁴⁰J. N. Owens, I. S. Gancarz, J. T. Koberstein, and T. P. Russell, *Macromolecules* **22**, 3380 (1989).
- ⁴¹J. Feng, H. Liu, and Y. Hu, *Macromol. Theory Simul.* **15**, 674 (2006).