Polymer collapse and crystallization in bond fluctuation models

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While the $\Theta$-collapse of single long polymers in bad solvents is usually a continuous (tri-critical) phase transition, there are exceptions where it is preempted by a discontinuous crystallization (liquid $\leftrightarrow$ solid) transition. For a version of the bond-fluctuation model (a model where monomers are represented as $2 \times 2 \times 2$ cubes, and bonds can have lengths between 2 and $\sqrt{10}$) it was recently shown by F. Rampf et al. that there exist distinct collapse and crystallization transitions for long but finite chains. But as the chain length goes to infinity, both transition temperatures converge to the same $T^*$, i.e. infinitely long polymers collapse immediately into a solid state. We explain this by the observation that polymers crystallize in the Rampf et al. model into a non-trivial cubic crystal structure (the ‘A15’ or ‘Cr$_3$Si’ Frank-Kasper structure) which has many degenerate ground states and, as a consequence, Bloch walls. If one controls the polymer growth such that only one ground state is populated and Bloch walls are completely avoided, the liquid-solid transition is a smooth cross-over without any sharp transition at all.

In spite of having been studied for several decades, the phase transitions of a single long polymer in a bad solvent are still a subject of active research with occasional big surprises. When temperature is lowered, most polymers undergo a $\Theta$-collapse, which is typically a continuous (tri-critical) phase transition. At the transition point, repulsive entropic effects and attractive energetic forces would cancel exactly for an infinitely long chain. For finite chain length $N$ theory \cite{1,2} predicts logarithmic corrections which are also seen in simulations \cite{2–4} and in experiments \cite{5}, and which also affect the ummixin transition of long polymers \cite{6}. Qualitatively, the $\Theta$-transition resembles a gas-liquid transition, with the open coil resembling the gas and the dense globule being analogous to a liquid. At even worse solvent conditions, in many models (e.g. in off-lattice polymers with Lennard-Jones interactions between monomers \cite{7}) there occurs a second ‘liquid-solid’ transition that is similar to crystallization. Since it is a discontinuous transition, its properties are usually not universal and are not described by perturbative renormalization group methods.

But there are well known cases where this $\Theta$-collapse is preempted by a discontinuous ‘freezing’ or crystallization transition. One example is semi-stiff polymers \cite{8}, another is provided by (off-lattice) monomers with hard core repulsion and attractive interactions which extend only very little beyond the core \cite{9,10}. Still another is the cooperative one-step collapse of some proteins \cite{11}

A very surprising behavior which falls between these two possibilities – $\Theta$-collapse with subsequent liquid-solid transition, and immediate gas-solid freezing – was observed by Rampf et al. \cite{12–16}. In their model, the specific heats of finite chains show rounded peaks at distinct temperatures $T_{\text{crys}}(N) < T_{\Theta}(N)$, but as the chain length $N$ diverges,

$$\lim_{N \to \infty} T_{\text{crys}}(N) = \lim_{N \to \infty} T_{\Theta}(N) = T^*.$$

Thus, while finite chains show the expected two distinct transitions, infinitely long polymers collapse in a single “hybrid” transition into their ground state \cite{17}.

In \cite{9,10} it was conjectured that this is connected to the range of interactions in this model, since the phenomenon disappeared when also more distant monomer pairs are included in the interaction. But it is not clear what would be the detailed mechanism for that. In the present paper, we propose a different explanation which is based on the fact that discontinuous phase transitions arise typically if there is a large gap between the (few) low energy states that dominate the low-$T$ phase and the bulk of states at higher energies. This is of course also the reason why there is a first order collapse in hard-core polymer models with very short range attraction: The shorter the range of the attractive potential, be bigger will be the effect of a single displaced monomer, as it will cause many monomer distances to be out of the interaction range. In the Rampf et al. model, the energetic distance between ground and excited states is, in contrast, dominated by topological defects similar to Bloch walls in magnetic systems that arise from a spontaneously broken symmetry.

We verify this conjecture by studying also a modified model where all such topological defects are completely eliminated, by constraining all configurations to be in the same ergodic component. This modified model has of course the same ground state energy, but it has a dramatically changed phase diagram: There is a standard $\Theta$ transition, but no crystallization transition at all. Instead, the transition from a disordered “liquid” to the densely packed ground state is a continuous cross-over. A similar behavior seems to prevail for interacting self-avoiding walks on the simple cubic lattice, although the interpretation there is controversial \cite{18}. In the very careful studies of \cite{18}, clear peaks in the specific heat were observed which obviously are related to some crystallization phenomena. But all indications point to the fact that they are not associated with bulk crystallization,
but with changes in the geometric shapes and surface properties of clusters which are already completely compact and ordered in their bulk.

The model used by Rampf et al. is the bond fluctuation model (BFM) of Carmesin et al. [10], augmented by an ansatz for the interactions. The BFM lives on a simple cubic lattice. Monomers are represented by $2 \times 2 \times 2$ cubes, i.e., if site $x = (x, y, z)$ is occupied by a monomer, all sites with Euclidian distance $< 2$ are excluded. These monomers are linked by bonds with integer coordinate components and length $2 \leq l \leq \sqrt{10}$. Thus the set of all bond vectors is made up by the vectors $(2, 0, 0), (2, 1, 0), (2, 1, 1), (2, 2, 0), (2, 2, 1), (3, 0, 0), (3, 1, 0)$ and all their rotations and reflections (altogether 108 vectors).

The solvent is not treated explicitly, but implicitly by assuming an attractive (negative) contact energy for any pair of monomers that touches each other along part of their surface (line and corner contacts do not contribute to this energy). Notice that two monomers can touch on their surface (line and corner contacts do not contribute pair of monomers that touches each other along part of vectors).

For any perfect arrangement of atoms in such a structure there exist many other arrangements which are just shifted copies of it. Thus this structure, seen as a ground state of a statistical mechanics model, is highly degenerate. If we prepare a big system such that it is in one of these ground states in some region of space of characteristic size $L$ and in another ground state somewhere else, there must be “Bloch walls” between them. These are topological defects with energies $\propto L^2$. It is these Bloch walls which are responsible for the discontinuity and high temperature of the freezing transition.

In contrast to [12–16], where Markov chain Monte Carlo methods were used, we used PERM [4] and “flat PERM” [22]. PERM is a growth algorithm with importance sampling and re-sampling, i.e., chains are grown in a biased way, with the bias compensated by weight factors. On the basis of weights of partially grown chains, the “population” of chains is controlled, similarly as in genetic models, by pruning and cloning. In the original PERM algorithm the growth is controlled by comparing the weight of the current configuration with the estimated partition sum, i.e., with the average weight of all previous configurations with the same length. In flat PERM it is compared with the average weight of those previous configurations that have the same length and energy. Both versions of the algorithm work perfectly, if the current weights of partially grown chains are good indicators of their final weights when the growth is finished. This is true for polymers at temperatures above $T_\phi$, and even more so at $T_\phi$, where its success is spectacular [4]. But it deteriorates quickly, if growth conditions change much during the growth and configurations which finally would be very good are pruned away at early stages. This happens typically in first order phase transitions. While Markov chain Monte Carlo methods are based on the concept of changing complete configurations to increase their Boltzmann weight, in PERM no configuration can be changed once it has been created.

In addition, we tried also another version called “new PERM” (nPERM) [23]. All three methods gave very similar results. Typical “good” results are shown in Fig. 1. There we present estimates of the average squared end-to-end distances, plotted against $N$ for several temperatures near $T_\phi$. At $T = T_\phi$ one expects

$$R_{\text{end\textendash end}}^2 \propto N,$$

up to logarithmic corrections. Although the leading corrections are known theoretically [11, 2], it is well known that they do not quantitatively describe the behavior at presently reachable chain lengths [23–4]. Nevertheless we clearly see deviations from Eq. (3) which qualitatively
FIG. 1. (Color online) Semi-logarithmic plot of $N^{-1}$ times the average squared end-to-end distances against chain length $N$. Instead of temperature, we use the Botzmann factor $q = \exp(\epsilon/k_B T)$ to label each curve. At $T = T_\Theta$ one expects a horizontal curve up to logarithmic corrections.

FIG. 2. (Color online) Logarithms of the density of states $g(E)$, plotted against the energy per monomer, for the modified model restricted to the sites in a perfect crystal. States $g(E)$, where $E$ is the total energy. Using $\epsilon = 1$ in the following, $|E|$ is just the number of contacts. Indeed, these estimates are needed to control the population growth, so estimating them is an essential part of the algorithm. In Fig. 2 we show $\ln g(E) + 1.672 E$ at fixed $N = 512$ against $E$ for several runs. Notice that 1.672 is the value of the Botzmann factor per contact at the effective freezing temperature according to [12]. Thus we expect the curve to be double-humped with both maxima having roughly the same height. Each run used different random number generator seeds and different minor parameters in the algorithm, and ran for about one week of CPU time on a fast PC. We see very good results for large $E$ (i.e. for energies relevant at $T \geq T_\Theta$), but a complete failure at energies which, according to [12], dominate the crystallized phase. The only positive statement these simulations allow to draw is that there is presumably really a first order transition, since otherwise the failure would not be expected to be as dramatic. Similar plots were obtained for all $128 \leq N \leq 1024$.

Let us now finally look at simulations of the constrained model, where we allowed monomers to be placed only at one of the positions given in Eq. (2) (Fig. 3). This time we show data for different values of $N$, up to $N = 2048$. The data now look absolutely clean. Moreover, we obtained good data down to energies which are close to the expected ground state energy $E_{\text{min}} = -6.75 N + O(N^{2/3})$ (see Fig. 4). It could of course be that these results are spurious, and that the simulations just missed those configurations which would have created troubles. We consider this as highly unlikely and take Fig. 3 as an indication that there is really no crystallization transition. Instead, when temperature is lowered, holes in the configuration are gradually filled up until a compact configuration is reached. Indeed, Fig. 3 looks

\begin{equation}
T_\Theta = 2.103(10). \tag{4}
\end{equation}

One advantage of PERM is that it allows to estimate absolute values of partition sums and of the density of agreement with the theoretical predictions. We fully confirm the estimates for $T_\Theta$ given in [12, 14], but due to the much longer chains and higher statistics we can give much smaller error bars in spite of the theoretical uncertainty:

$T_\Theta = 2.103(10)$. \tag{4
much better than analogous plots for the simple cubic lattice. There we of course also have no topological defects, but the much lower coordination number implies that the growth is much more often blocked at low temperatures. For the fcc lattice, however, where coordination number is 12, we again found very clean results.

In summary, we have shown that the very strong discontinuous crystallization transition in the polymer model of [12] is due to spontaneously broken translational symmetry in its low energy state. Similar discontinuous transitions are expected (and found) in continuum models, if the polymer can freeze into a periodic monomer configuration. We do not expect any first order transition if the low energy state is glassy (such as in a Lennard-Jones homopolymer with bond lengths incommensurate with the Lennard-Jones radius) and in lattice models without spontaneous symmetry breaking, such as e.g. the simple cubic lattice. The reason why crystallization is qualitatively different in the bond fluctuation model of [11] is primarily not because of the increased average range of interactions, but because of the changed structure of the ground state and low energy excitations.

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