

Continuous melting of compact polymers

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The competition between chain entropy and bending rigidity in compact polymers can be addressed within a lattice model introduced by P.J. Flory in 1956. It exhibits a transition between an entropy dominated disordered phase and an energetically favored crystalline phase. The nature of this order-disorder transition has been debated ever since the introduction of the model. Here we present exact results for the Flory model in two dimensions relevant for polymers on surfaces, such as DNA adsorbed on a lipid bilayer. We predict a continuous melting transition, and compute exact values of critical exponents at the transition point.

Condensed phases of polymers are ubiquitous in the living world. Examples range from the organization of DNA in viruses, to the native state of globular proteins. For example, DNA condensation is required in order to fit the complete viral genome, which can be tens of microns long, into the viral capsid typically tens of nanometers in diameter. As the persistence length of the DNA (50 nm) is comparable to the linear dimension of the viral capsid (10–100 nm), one would expect it to pack into ordered, crystalline configurations. Indeed, cryo-electron-microscopy studies of the T7 virus have revealed a circular, inverse-spool arrangement of DNA in the capsid [1]. The observed condensed state of DNA is a result of many competing effects such as chain and solvent entropy, electrostatic interactions, bending stiffness of the polymer, and interaction with capsid proteins [2].

These observations motivate the study of the competition between chain entropy and bending energy of a polymer chain under extreme confinement, when only compact configurations are allowed. The resulting transition between an entropy dominated disordered state and an energetically favored crystalline one is captured by a simple lattice model of a compact semiflexible polymer, first introduced by Flory almost 50 years ago in the context of polymer melting [3]. It has also served as the starting point for constructing simple lattice models of protein folding [4].

In this Letter, we provide an analytic solution of the Flory model in two dimensions using field theoretical methods. We resolve the long-standing debate [5, 6, 7, 8, 9, 10] about the nature of the transition, where some numerical results [7] and approximate analytic approaches [8] were indicative of a first-order transition, while other numerical work [9, 10] and theoretical arguments based on similarities with the exactly solvable six-vertex model [5, 9], seemed to imply a continuous transition. We show that the melting transition of a single semiflexible compact polymer is continuous. Furthermore, we characterize the transition precisely by calculating the exact values of the critical exponents that accompany it.

Semiflexible loop model In the Flory model a configuration of a polymer chain is described by a random walk on a square lattice (cubic, in three dimensions) which visits every site of the lattice exactly once. The energy associated with a configuration is $B\epsilon > 0$, where B is the number of bends in the chain. At large temperatures ($k_B T \gg \epsilon$) bends proliferate for entropic reasons, thus reducing the persistence length of the polymer. In the low-temperature limit ($k_B T \ll \epsilon$) bends are very unlikely and the persistence length is large, of the order of the lattice size, as seen in fig. 1. The central question we address here is the nature of the transition between the disordered high-temperature phase and the ordered low-temperature phase.

The interplay between the entropy and the energy is encoded in the partition function of the Flory model

$$Z_F = \sum_{\Gamma_F} w^{B(\Gamma_F)}, \quad (1)$$

where $w = \exp(-\epsilon/k_B T)$ is the Boltzmann weight associated with a bend, and $B(\Gamma_F)$ is the number of 90° bends in the compact chain configuration Γ_F . For $T \rightarrow 0$ ($w \rightarrow 0$) the equilibrium state is one that

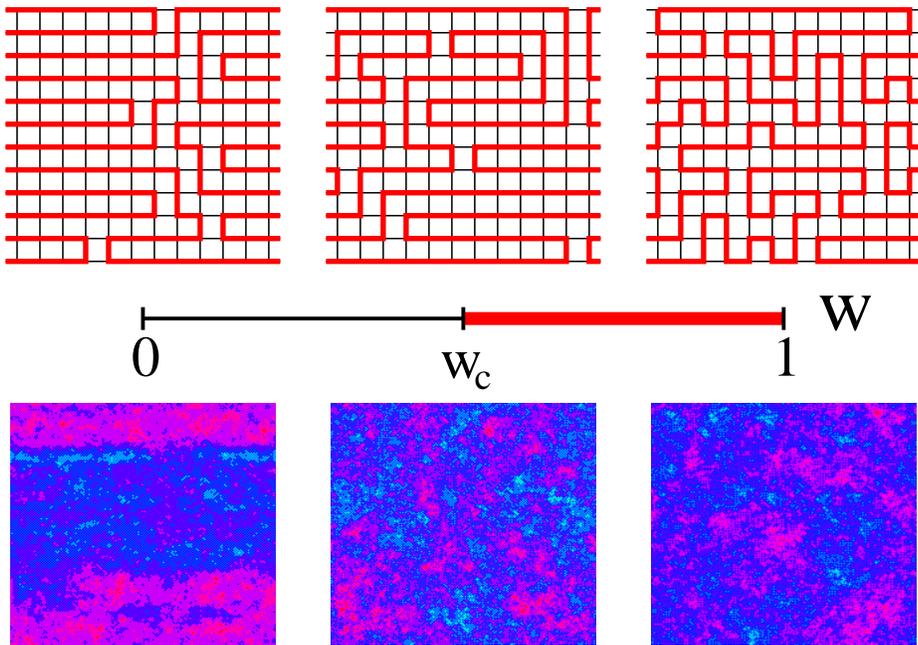


FIG. 1: Phase diagram of the Flory model. The Boltzmann weight of a single bend in the chain is w . For $w = 1$ the model describes a flexible compact polymer with many bends present in a typical polymer configuration. For $w < 1$ there is an energy penalty associated with bending and we are dealing with a semiflexible chain. For $w_c \leq w \leq 1$ the chain is critical. At $w = w_c$ there is a continuous transition to an ordered chain configuration; the order parameter is the difference between the number of vertical and horizontal links. The polymer configurations shown in the top three panels (with periodic boundary conditions imposed in the horizontal direction) are typical for $w = 1/3$, $w = w_c \approx 1/2$, and $w = 1$. The density plots below them are of the second height component in the related loop model, for the same values of w , clearly showing the transition from a rough to a flat interface.

minimizes B , while for $T \rightarrow \infty$ ($w \rightarrow 1$) entropy dominates and there is a large, macroscopic number of bends in a typical chain configuration; see fig. 1. In order to investigate the transition between the high and low temperature phase we construct a mapping of the polymer problem to a lattice model of fluctuating loops, following Nienhuis [11].

We define the semiflexible loop model (SFL), whose configurations Γ_L are all possible ways of covering the square lattice with non-intersecting loops. The partition function of the SFL model reads

$$Z_L = \sum_{\Gamma_L} n^{N(\Gamma_L)} w^{B(\Gamma_L)}, \quad (2)$$

where each of the $N(\Gamma_L)$ loops in the configuration Γ_L carries a weight $n > 0$, and bends are weighted by w as in the Flory model. Note that the $n \rightarrow 0$ limit of Z_L/n yields the Flory model partition function for a single *closed* compact polymer. Instead of explicitly computing Z_L the approach we take is to construct a continuum limit of the SFL model in the form of a field theory. Computations of correlation functions in this field theory lead to the phase diagram and exact scaling exponents for any n , including the polymer ($n \rightarrow 0$) limit.

Height model Since loops in Γ_L do not intersect they can be interpreted as contour lines of the height of an interface. In the SFL model, the height is a three component scalar field. To see this, we first note that bonds not covered by loops form so-called *ghost* loops, if periodic boundary conditions are imposed. Then we assign to each real and ghost loop an orientation (clockwise or anticlockwise) independently and randomly. By doing so, each bond of the square lattice is found in one of four states, labeled **A**, **B**, **C**, or **D**, depending on whether it is covered by a real or a ghost loop oriented in one or the other direction.

We parametrize $n = 2 \cos(\pi e)$, with $0 \leq e < 1/2$, and assign a weight $\exp(\pm i\pi e)$ (resp. $\exp(\pm i\pi/3)$) to each orientation of a real (resp. ghost) loop. Tracing over the two loop orientations, we recover the desired weights: n for the real loops and 1 for the ghost loops.

The heights for the SFL model are defined at the centers of the elementary square plaquettes of the lattice. Height differences between neighboring plaquettes are given by *vectors* \mathbf{A} , \mathbf{B} , \mathbf{C} , or \mathbf{D} . Since each vertex is shared by four bonds, all necessarily in different states, a single relation, $\mathbf{A} + \mathbf{B} + \mathbf{C} + \mathbf{D} = 0$, results from requiring that the total height increment when traversing a closed lattice path be zero. Thus, we conclude that out of the four vectors three are linearly independent and that the heights are three dimensional. For the remainder of the paper we adopt the following normalization: $\mathbf{A} = (-1, +1, +1)$, $\mathbf{B} = (+1, +1, -1)$, $\mathbf{C} = (-1, -1, -1)$, and $\mathbf{D} = (+1, -1, +1)$. The partition function of the SFL model can now be rewritten as a sum over height configurations, since a height configuration uniquely specifies an oriented loop configuration.

The continuum limit of the SFL model is obtained by coarse-graining the height field over domains much smaller than the system size and much bigger than a single plaquette. This yields a continuous height function $\mathbf{h}(\mathbf{x}) = (h^1(\mathbf{x}), h^2(\mathbf{x}), h^3(\mathbf{x}))$ defined over the plane \mathbf{x} . For $n \leq 2$ the height field is assumed rough [12] and the partition function can be written as a sum (path integral) over all height functions with weights given by $\exp(-S)$, where

$$S = \frac{1}{2} \int d^2\mathbf{x} K_{ij} (\partial h^i \cdot \partial h^j) + \frac{i}{4\pi} \int d^2\mathbf{x} (\mathbf{e}_0 \cdot \mathbf{h}) \rho(\mathbf{x}) + \gamma \int d^2\mathbf{x} \cos(2\pi h^2) . \quad (3)$$

This particular form of the coarse-grained free energy S is dictated by the symmetries of the loop model which involve translations in the space of heights, and permutations of the height-difference vectors [12]. The value of the background charge

$$\mathbf{e}_0 = -\frac{\pi}{2}(1/3 + e, 0, 1/3 - e) \quad (4)$$

follows from imposing the correct weighting of loops that close at infinity ($\rho(\mathbf{x})$ is the curvature of the \mathbf{x} -surface), while the non-zero elastic constants

$$K_{11} = \frac{\pi}{8}(5/3 - e), \quad K_{13} = \frac{\pi}{8}(e - 1/3), \quad K_{22} = K_{22}(e, w) \quad (5)$$

are partially fixed by the loop ansatz which states that loop weights do not flow under renormalization [13].

In the previously studied infinite-temperature limit ($w \rightarrow 1$), the elastic constant $K_{22} = \pi(1 - e)/(5 - 3e)$ is also fixed by the loop ansatz, and the cosine term is absent from S [12]. For $w < 1$ the cosine potential accounts for the energy penalty associated with bends. Its scaling dimension in the $w = 1$ theory, $x_B = \pi/K_{22} = (5 - 3e)/(1 - e)$, is greater than two for $0 \leq e < 1/2$ ($0 < n \leq 2$), and is therefore an *irrelevant* perturbation. It has the effect of renormalizing the elastic constant K_{22} which ultimately drives the transition to the crystalline phase at $w = w_c$.

Phase diagram and critical exponents The field theory of the SFL model implies the following phase diagram for any $n \leq 2$, including the polymer limit $n \rightarrow 0$. For $w = 1$ the interface described by the height field is *rough* and the loop model is critical, i.e., it is characterized by a power-law distribution of loop sizes. As w is reduced, the bending stiffness of the loops increases, and the height remains rough until a critical value $w = w_c$. At w_c there is a *roughening transition* [14] to a smooth interface; see fig. 1. In the loop model this corresponds to a phase with a vanishing density of bends. The roughening transition is continuous, in the Kosterlitz-Thouless universality class. At the transition the cosine term in S becomes relevant in the renormalization group sense [14]. In other words, the scaling dimension of the operator $\cos(2\pi h^2)$ at $w = w_c$ is 2. Computing this dimension using the field theory defined by S then leads to the prediction $K_{22}(e, w_c) = \pi/2$. One important test of this result comes from the $n = 1$ ($e = 1/3$) SFL model, which maps to the exactly solvable six-vertex model. In this case $w_c = 1/2$ and $K_{22} = \arcsin(1/2w)$ have been computed previously [15].

In the polymer limit the transition described by the field theory takes us from a critical geometry of a disordered compact polymer, at low bending stiffness, to an ordered state, at high bending stiffness;

w^{-1}	K_{22}	θ_{th}	θ_{num}
1	$\pi/7$	5/56	0.088
1.2	0.584	0.004	0.004
1.4	0.736	-0.094	-0.096
1.6	0.924	-0.214	-0.22
1.8	1.210	-0.404	-0.40
$w_c^{-1} \approx 2$	$\pi/2$	-5/8	-0.7

TABLE I: Numerical transfer matrix results. Exact (for $w = 1, w_c$) and numerically determined values of the elastic constant K_{22} were used to compute the exponent θ_{th} from eq. (6), for various values of the bend weight w ; this compares favorably with the direct numerical determination θ_{num} .

see fig. 1. The critical phase is characterized by scaling exponents ν and θ . Namely, the probability distribution $p(r, l)$, for the end-to-end distance r of a polymer of length l , has a scaling form $p(r, l) = r^\theta f(r/l^\nu)$ [16]. Using the field theory defined by eq. (3) these exponents can be computed exactly [12]. We find

$$\nu = \frac{1}{2}, \quad \text{and} \quad \theta = \frac{3\pi - 16K_{22}}{8\pi}; \quad (6)$$

note that θ varies continuously with w via the elastic constant K_{22} . The calculation of ν gives another important check on the field theory since $\nu = 1/2$ follows directly from the fact that compact polymers completely fill the plane. Finally, exact values for K_{22} in the $w = 1$ and $w = w_c$ case, lead to predictions $\theta(w = 1) = 5/56$ and $\theta(w = w_c) = -5/8$. These values of θ have a simple physical interpretation. Namely, in the absence of bending stiffness ($w = 1$) the ends of the chain repel (positive θ), as is the case for open chain conformations of polymers in a good solvent. Somewhat surprisingly, this entropic repulsion decreases and turns to attraction (negative θ) as the chain stiffness is increased.

Numerical transfer matrix results The main result of this paper, the prediction of continuous melting of compact polymers in two-dimensions, rests on the validity of the proposed field theory of the Flory model, eq. (3). To further test the field theory we have made use of a numerical transfer matrix approach [12] to compute the elastic constant K_{22} and the exponent θ . Results are shown in table I. We see that eq. (6) is satisfied to a very good approximation and that the exact values for the two quantities at $w = 1$ and $w = w_c$ are confirmed by the numerical computations.

Experiments DNA adsorbed on a lipid bilayer provides a laboratory for testing theories of two-dimensional polymers. Previous experiments [17] on this system have measured the theoretically predicted scaling exponents derived from the two-dimensional self-avoiding walk model. These measurements were done at low DNA concentrations. As the amount of DNA on the surface is increased it was observed that single chain conformations become more compact. To study the transition discussed here compaction to areas of linear dimensions approaching the DNA persistence length would be needed. It would then be interesting to monitor the two ends of the DNA using fluorescent labels, as the amount of compaction is increased. This would provide direct information about $p(r, l)$ and the exponent θ computed above.

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