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Detailed Terms
Polymer-mediated entropic forces between scale-free objects

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The number of configurations of a polymer is reduced in the presence of a barrier or an obstacle. The resulting loss of entropy adds a repulsive component to other forces generated by interaction potentials. When the obstructions are scale invariant shapes (such as cones, wedges, lines, or planes) the only relevant length scales are the polymer size \(R_0\) and characteristic separations, severely constraining the functional form of entropic forces. Specifically, we consider a polymer (single strand or star) attached to the tip of a cone, at a separation \(h\) from a surface (or another cone). At close proximity, such that \(h \ll R_0\), separation is the only remaining relevant scale and the entropic force must take the form \(F = A k_B T / h\). The amplitude \(A\) is universal and can be related to exponents \(\eta\) governing the anomalous scaling of polymer correlations in the presence of obstacles. We use analytical, numerical, and \(\epsilon\)-expansion techniques to compute the exponent \(\eta\) for a polymer attached to the tip of the cone (with or without an additional plate or cone) for ideal and self-avoiding polymers. The entropic force is of the order of 0.1 pN at 0.1 \(\mu\)m for a single polymer and can be increased for a star polymer.

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I. INTRODUCTION

A host of single molecule manipulation techniques [1–5], employing atomic force microscopes (AFMs) [6], microneedles [7], and optical [8,9] and magnetic [10] tweezers, have provided quite detailed studies of shapes and forces in long polymers. The positional accuracy of an AFM tip [5,11] can be as good as a few nm, while forces of order of 1 pN can be measured, with measurements carried out in nearly biological conditions [12,13]. The main thrust of current experimental research is to extract specific information about molecular shapes and reactions from force-displacement curves. The interpretation of such data is complicated by a host of factors such as interactions of the molecule with the probe, and nonequilibrium effects arising from rapid and large imposed deformations. Nonetheless, much recent activity is devoted to obtaining the equilibrium free energy landscape of long molecules by processing results from repeated nonequilibrium measurements (see, e.g., Refs. [14–16]). Here, we consider another contribution to the force arising from the loss of entropy in the presence of constraints imposed by the probes. While most likely weaker than forces due to interaction potentials, the enhanced AFM sensitivities are indeed approaching the range where entropic forces of long polymers in a solvent can be significant (even when the deformation of the polymer is relatively slight). From the theoretical perspective, the attraction of entropic forces on polymers is their independence of microscopic details [17].

At a finite temperature \(T\), a long homopolymer in a good solvent fluctuates between a large number of configurations, and its “macroscopic” properties (such as size and elasticity beyond a characteristic persistence length) are governed by entropic considerations. To illustrate entropic elasticity, consider the hypothetical situation of a polymer with one end fixed to the origin, while the other end explores allowed positions at \(\vec{r}\). In equilibrium, due to rotational symmetry \(\langle \vec{r} \rangle = 0\). The symmetry is broken if a weak force \(\vec{F}\) is applied by a pointlike probe at the end of the polymer. Then, in \(d\)-dimensional space \(\langle \vec{r} \rangle = (R_0^d / dk_B T) \vec{F}\), where \(R_0^d\) is the mean squared end-to-end distance in the absence of external force [17]. This is mostly a realignment, without substantial change of polymer conformations for \(F \ll F_{\text{cr}} \equiv dk_B T / R_0\). For \(R_0 = 0.1 \mu\)m at room temperature \(F_{\text{cr}} \approx 0.1\) pN is the crossover value that separates weak and strong forces; stronger force \((F \gg F_{\text{cr}})\) leads to substantial stretching of the polymer and associated reduction in entropy. Similar considerations come into play when the attachment points are replaced by macroscopic obstacles (probes): Under a strong stretching force polymer configurations are not very much influenced by the shapes of the probes, and convenient models (such as Ref. [18]) have been developed for the resulting force-extension relations. Here, we focus on the opposite limit of strong compression in which the loss of entropy, and the corresponding force, are dominated by confinement within boundaries.

In this work we consider setups where a polymer is attached by one end to an object (the probe), while the other end is free. The probe is then brought to the vicinity of a second solid obstacle, and the loss of polymer entropy leads to a force between the two objects. In particular, we consider confining (probe and obstacle) shapes that are self-similar (at least on scales comparable to \(R_0\)) at separations \(h \ll R_0\). Possible setups include sharp AFM tips approaching flat surfaces covered with polymers, as in experimental studies [19–23] of polymer brushes. Entropic considerations in a dense brush must account for interactions between polymers [24], complicating the calculation of forces exerted on the AFM tip [25]. Here we consider the simpler cases of a single linear or star polymer, avoiding the dense limit. For \(h \ll R_0\) dimensional considerations suggest the particularly simple force-separation relation \(F = A k_B T / h\), where the “universal” amplitude \(A\) only depends on basic geometrical properties and gross features of the polymer.

In a previous paper we outlined some results pertaining to entropic interactions, and critical exponents, for a polymer...
attached to a cone [26]. In this paper we expand on the
details of the calculations, as well as generalize and expand
the results to other setups. Section II demonstrates how the
force amplitude $A$ is related to the exponent $\eta$, characterizing
the anomalous scaling of polymer correlations. The latter exponent depends on the shape of confining boundaries and is the focus of computations that follow. In Sec. III we outline the
general approach to computing $\eta$ for ideal polymers near
scale-invariant surfaces, while in Sec. IV we make explicit
computations for cones and plates. The case of self-avoiding
linear polymers in $d = 3$ is treated by Monte Carlo simulations
in Sec. V. Perturbative $\epsilon = 4 - d$ expansions are employed in
Sec. VI to compute $\eta$ for linear and star self-avoiding polymers
attached to cone tips or to a contact point between a cone and
a plate.

II. POLYMER-MEDIATED FORCES

Let us first consider an idealized setup in which a polymer
is attached to the tip of a solid cone, approaching a solid plate
(or another cone). This exemplifies a geometry of obstacles in which the only (nonmicroscopic) length scale is provided by the tip-plate (or tip-tip) separation $h$. The polymer itself undergoes self-similar fluctuations, spanning length scales intermediate between microscopic (monomer size, or persistence length $a$) and macroscopic. The latter is set by the typical end-to-end distance $R_0$, or by the radius of gyration $R_g$ (which differs from $R_0$ only by a multiplicative factor of order of unity). The typical size of the polymer grows with the number of monomers $N$ through the scaling relation $R_0 \sim R_g \propto N^{\nu}$. The self-similar shape of the obstacles also presumably extends only up to a characteristic scale $H$, say the height beyond which the cone is terminated or changes shape.

Neglecting any structure (and hence energy) associated with
the polymer, variations in free energy as the cone (plus tip-
attached polymer) approaches the plate are entirely entropic
in origin and proportional to $k_B T$. The entropic force has
dimensions of energy divided by length, and at separations $a \ll h \ll R_0 \ll H$ the relevant length scale is $h$, and it is reasonable to posit

$$ F = A \frac{k_B T}{h}. \tag{1} $$

This is because the polymer configurations are constrained only on the scale of confinement $h$; increasing the length of the polymer or the size of the cone (as long as $h \ll R_0 \ll H$) should not influence the force. Furthermore, the entropy change is independent of $a$.

Indeed, the simple force law of Eq. (1) applies to all circumstances where the separation provides the only relevant length scale and follows easily from various polymer scaling forms (see, e.g., the derivation below), such as in Refs. [27–29]. The dimensionless amplitude $A$ will depend on geometric factors characterizing the confining boundaries such as the
opening angle of a cone $\Theta$ (and if tilted, on the corresponding angle). It will also depend on factors that characterize the
scaling of polymeric fluctuations, thus differing in cases of ideal and self-avoiding polymers, and for linear, star, and brush polymers. In the following we demonstrate that in a number

of setups, the amplitude $A$ can be related to variations of the
universal, but shape dependent) exponent $\eta$, characterizing polymeric correlations.

A chain starting at the tip of a cone provides a first approximation to a polymer linked to the tip of an AFM probe as depicted in Fig. 1(a). With the cone far away from a plate ($h \gg R_0$), the number of configurations of the polymer grows with the number of monomers as

$$ N_c = b z^N N^{\nu(\eta)-1}, \tag{2} $$

where the effective coordination number $z$, as well as the prefactor $b$, depend on the microscopic details (such as the scale $a$), while the “universal” exponent $\nu$ only depends on the cone angle. When the cone touches the plate as in Fig. 1(c), the number of configurations is reduced to $N_{cP}$ with the same form as Eq. (2), but with a different exponent $\nu_P(\eta)$. We henceforth use an exponent subscript “s” (as in $\nu_s$) to refer to the above cases, with “s = c” for cone and “s = cp” for cone + plate; the absence of a subscript (as in $\nu$) signifies a free polymer. The work done against the entropic force in bringing in the tip from afar to contact the plate can now be computed from Eq. (1) as

$$ W = \int_a^{R_0} dh A \frac{k_B T}{h} = A k_B T \ln \frac{R_0}{a} = A \nu k_B T \ln N. \tag{3} $$

This work can also be computed from the change in free
energies between the final and initial states, due to the change in entropy, as

$$ \Delta F = -T \Delta S = T S_c - T S_{cp} = k_B T (\nu_c - \nu_{cp}) \ln N, \tag{4} $$

FIG. 1. (a) Polymer attached to the tip of a solid cone with apex semiangle $\Theta$ (configuration “c”); positions are described by the spherical coordinates $r, \theta$ and azimuthal angle $\phi$ (not shown). (b) The tip, where the polymer is attached, is at a distance $h \ll R_0$ from the plate. (c) The tip touching the plate (configuration “cp”). (d) Tip is at a finite distance from a plate to which the polymer is attached. (e) Polymer attached to both surfaces.
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FIG. 2. Examples of three-dimensional figures without a length scale; gray surfaces indicate truncation only for graphical representations. (a) Circular cone, (b) two-dimensional sector of a circle (in three-dimensional space), (c) pyramid, (d) wedge, (e) plane, (f) line, (g) point. A polymer is to be attached to the point (g), to the apex of (a), (b), (c), or to any point on the edge of (d) or the entirety of (e) or (f). The plane and line can also be semi-infinite with a polymer attached to their edge or end point.

with the entropy $S = -k_B \ln N$ computed from Eq. (2). By equating $W$ and $\Delta F$ we find

$$A = \frac{\gamma_s \eta_p}{\nu} = \eta_{cp} - \eta_c;$$

the final result obtained from the scaling law [17]

$$\gamma_s = (2 - \eta_s)\nu;$$

where the exponent $\eta_s$ characterizes the anomalous decay of correlations ($\sim 1/r^{d+2+\eta_s}$).

The arguments presented in the previous paragraph rely only on the fact that configurations of the obstacles lack a length scale for both $h \to 0$ and $h \to \infty$. Consequently, similar reasoning can be applied to a variety of objects such as those depicted in Fig. 2, or combinations of such objects. The force prefactor is then related to the exponents in the initial and the final states by

$$A \equiv \frac{\gamma_{initial} - \gamma_{final}}{\nu} \equiv \eta_{final} - \eta_{initial}.$$

Consider, for instance, an ideal polymer in free space (or held by a pointlike object [Fig. 2(g)]). Its number of configurations is $z^N$, that is, $\gamma \equiv \gamma_0 = 1$, and correspondingly $\eta_0 = 0$, since $\nu = \nu_0 = 1/2$. Subscript 0 henceforth indicates exponents describing ideal polymers. If the end of the polymer is brought into contact with a plane, then by the method of images [30] it can be shown that in any space dimension $d$ the number of configurations scales as $zN^{-1/2}$, that is, with $\gamma \equiv \gamma_0 = 1/2$, and corresponding to $\eta_0 = 1$. Thus, for a long polymer held by a pointlike object at a distance $h$ from a plane, the entropic force in Eq. (1) will have a prefactor $A = 1$. This result is valid for any $d$. For self-avoiding polymers in $d = 2$, the exponents are $\gamma = 43/32$, $\gamma_p = 61/64$ [31], and $\nu = 3/4$ [32], leading to $A = 25/48 \approx 0.52$, while in $d = 3$, we get $\gamma \approx 1.158$ [33], $\gamma_p \approx 0.70$, and $\nu \approx 0.59$ [34], leading to $A \approx 0.78$. Similar exact results can be derived for more complex self-avoiding polymers, such as star polymers or branched polymers of arbitrary topology in $d = 2$ [28].

Note that the force amplitude does depend on the surface to which the polymer is attached. For example, consider the configuration in Fig. 1(d) where the polymer is attached to the plate. The initial configuration corresponds to a polymer attached to an infinite surface, thus $A_{initial} = \eta_p$, while the final configuration is the same as in Fig. 1(c), with $A_{final} = \eta_c$. This leads to $A = \eta_{cp} - \eta_p$, as opposed to $A = \eta_{cp} - \eta_c$ for the case that the polymer is attached to the cone. There are a number of detailed studies of small surfaces (such as AFM tips) [35–41] compressing a single polymer (or few polymers) grafted to a surface. However, all these studies concern non-scale-invariant geometries.

When the obstacles are separated by $h$, the loss of polymer entropy [and the corresponding pressure leading to the force in Eq. (1)] is concentrated in the confinement region. The part of the polymer that wanders away from this area is relatively unperturbed and does not contribute to the force. Therefore, for large $N$, Eq. (1) is independent of polymer size. This is not the case when the entire polymer contributes to the force, as for a polymer held between two plates, where the force is proportional to $N$. Thus, the argument fails when dimensionality of the system is changed between the initial and the final states. For example when a three-dimensional polymer connected to a two-dimensional plane approaches a parallel plane, in the final configuration the confined polymer is essentially two-dimensional. Free energies in the initial and final state have different extensive parts, and the polymer-mediated force depends on the number of monomers [42].

It is worth reiterating the idealizations leading to Eqs. (1) and (7). Real obstacles are self-similar over a range of length scales; for example, an AFM tip may be rounded or abruptly change its angle. Equation (1) will then be applicable to a corresponding interval, and there may be nontrivial crossovers between the different regimes [43]. By focusing on entropy, we have assumed that the only interaction between the polymer and the obstacles is due to hard-core exclusion. Attractive interactions between the polymer and surface will introduce temperature dependent corrections and additional size scales [17]. Weak interactions are asymptotically irrelevant, but strong interactions may lead to a phase in which the polymer is absorbed onto the obstacles [27], rendering the entropic considerations presented here inappropriate. At high-enough temperatures we expect entropic interactions to be dominant [17]. However, the absorption of AFM constrained polymers to attractive probes is indeed an interesting topic which falls outside the purview of this paper.

III. IDEAL POLYMERS NEAR SCALE-INARIANT SURFACES

Having reduced the computation of the force to calculation of correlation functions, the remainder of the paper is dedicated to computing exponents characterizing polymer correlations. The behavior of idealized polymers, that is, in the absence of self-avoidance and other interactions, is closely related to the diffusion problem: The spatial configuration of a polymer of $N$-segments can be regarded as the space-time trajectory of a random walker viewed at integer times up to $t = N$. Summing over all trajectories starting at $r$ and ending at $r'$ is then equivalent to considering the probability of diffusion...
from \( r \) to \( r' \) in time \( t \), which satisfies the diffusion equation

\[
\frac{\partial P(r',t)}{\partial t} = D \nabla^2 P(r',t).
\]

The prime sign on the Laplacian indicates the spatial derivatives with respect to components of \( r' \), and the initial condition is \( P(r',t=0) = \delta^d(r - r') \). The diffusion constant \( D \) is chosen such that in free space the mean squared distance coincides with the random walk value of \((r - r')^2 = a^2N = 2dDt\) in \( d \) space dimensions, and thus \( D = a^2/2d \). If a polymer is confined by repulsive walls, its configurations correspond to paths of a random walker that does no cross these boundaries. In the language of diffusion, the exclusion of paths that cross the boundaries is accomplished by assuming that they are absorbing surfaces.

To compute variations in the number of states, we are interested in the total survival probability \( S(r,t) \equiv \int P(r,r',t)d^d r' \), which also satisfies the equation \([44]\)

\[
\frac{\partial S(r,t)}{\partial t} = D \nabla^2 S(r,t),
\]

with absorbing boundary conditions. The initial condition for survival probability is \( S(r,t=0) = 1 \), everywhere inside the space where the particle can diffuse, and \( S(r,t) = 0 \) on the absorbing boundaries. (While the space and time variables are continuous in the above equation, appropriately discretized equations can be applied to diffusion or random walk on a lattice.)

In the absence of confinement, the number of configurations \( N \) of an ideal polymer grows exponentially with its length \( N \). On a regular discrete lattice, \( N \) is simply related to lattice coordination number \( z \) by \( N = z^N \). The presence of repulsive boundaries reduces the number of configurations to

\[
N(r,N) = z^N S(r,kN),
\]

where the prefactor \( k = a^2/(2dD) \) converts the dimensionless number of steps \( N \) to time \( t \) in the diffusion equation. Numerical solutions of Eqs. (8) and (9) can be easily found for many geometries, and analytical solutions are available for a number of simple shapes [45]. The generally complicated solutions simplify to some extent for the self-similar shapes depicted in Fig. 2. In a coordinate system centered on a special point of the object (such as tip of a cone) the dimensionless function \( S \) can only depend on the dimensionless vector \( \mathbf{w} = r/\sqrt{Dt} \). Thus, \( S(r,t) = H[w] \), and Eq. (9) reduces to

\[
\nabla^2_H w + \frac{1}{2} \mathbf{w} \cdot \nabla w H = 0,
\]

where the subscript \( w \) indicates derivatives with respect to components of \( w \). In terms of these dimensionless variables, the function \( H \) vanishes on the absorbing surfaces and increases to unity away from the surface (at separations of order one). For some geometries, the solution to Eq. (11) can be presented in terms of a radial distance \( w \) and a combination of angular variables. For example, generalized cones in \( d \)-dimensional space are described by a single polar angle \( \theta \) and \( d - 2 \) azimuthal angles \( \phi, \psi, \ldots \). (While \( d \)-dimensional wedges can be treated similarly, it is more convenient to note that the solution is independent of the coordinates parallel to the wedge, while the cross-section perpendicular to the wedge resembles a cone.) Furthermore, for \( w \ll 1 \) the distance dependence is expected to be a simple power law \( w^\eta \Psi(\theta, \phi, \ldots) \). In this limit, the second term in Eq. (11) becomes negligible, and the problem reduces to solving the Laplace equation \( \nabla^2 \tilde{w} = 0 \). The boundary conditions are now implemented solely in terms of angular variables, and the problem is reduced to finding a non-negative function \( \Psi \) that satisfies these conditions and the corresponding \( \eta_0 \) for which such solution exists. The dependence of the latter value on the boundary shape is emphasized by denoting it as \( \eta_0 \). Since \( S \sim w^{\eta_0} \equiv (r/\sqrt{Dt})^{\eta_0} \) for a random walk that starts a short distance \( |r| = a \) away from the cone, the survival probability is \( S \sim (a/\sqrt{Dt})^{\eta_0} \), and consequently the number of configurations of an ideal polymer grows as \( z^N N^{-\eta_0/2} \equiv z^N N^{\eta_0} \). We see that \( \gamma_0 = 1 - \eta_0/2 \) as in Eq. (6) for \( \eta_0 = 1/2 \).

IV. IDEAL POLYMERS NEAR CONES AND PLATES

As an example, consider a polymer attached to the apex of a cone, as in Fig. 1(a), or to a point of contact between a cone and a plane, as in Fig. 1(c). When generalized to \( d \) dimensions, this system has cylindrical symmetry, and \( \Psi \) is independent of the \( d - 2 \) azimuthal angles. Therefore, the Laplace equation simplifies to

\[
\frac{1}{(\sin \theta)^{d-2}} \frac{d}{d\theta} \left[ (\sin \theta)^{d-2} \frac{d \Psi}{d\theta} \right] + \eta_0(d - 2 + \eta_0) \Psi(\theta) = 0,
\]

with an appropriate boundary condition on \( \Psi(\theta) \). For an isolated cone, the function \( \Psi \) must be positive and regular outside the cone, with \( d \Psi/d\theta|_{\theta=\pi} = 0 \) to avoid a cusp on the symmetry axis, and \( \Psi(\theta) = 0 \) on the cone surface. For the cone plus plane, the appropriate solution is positive and vanishes at both \( \theta = \Theta \) and \( \theta = \pi/2 \). The first case was considered by Ben-Naim and Krapivsky [46] in connection with diffusion near an absorbing boundary [47], and we follow their derivations. The solution in general \( d \) requires the use of associated Legendre functions, but simplifies in a few cases described below.

(i) For \( d = 2 \), the problem of a cone coincides with that of a wedge. In higher space dimensions the wedge remains equivalent to the two-dimensional case, as the function \( H \) is independent of the \( d - 2 \) coordinates parallel to the wedge. Thus, the following results for \( d = 2 \) are also applicable to wedges in any \( d \). Equation (12) now reduces to \( \Psi^\eta \equiv \eta_0^2 \Psi = 0 \), which is solved by linear combinations of \( \sin(\eta_0 \theta) \) and \( \cos(\eta_0 \theta) \). The requirement that \( \Psi \) is positive and regular and vanishes on the object(s) yields

\[
\eta_0 = \frac{\pi}{2(\pi - \Theta)} \quad \text{and} \quad \eta_{cp0} = \frac{2\pi}{\pi - 2\Theta}.
\]

Both results (depicted in Fig. 3) go to a finite value as \( \Theta \to 0 \), reflecting the strong reduction in configurations due to the remnant (barrier) line, and \( \eta_0 \to \infty \) when the boundaries confine the polymer to a vanishing sector.

This computation can be easily generalized to the case of two cones (wedges) ("s = cc") with apex semicoordinates \( \Theta_1 \) and \( \Theta_2 \) touching at their tips with a common symmetry axis. If one of the wedges is tilted by an angle \( r \) relative to that axis,
then the survival probability $S$ will be determined by the larger sector of free space. The resulting exponent is

$$
\eta_{c0} = \frac{\pi}{\pi - (\Theta_1 + \Theta_2 - \pi)}.
$$

By proper choices of $\Theta$‘s and $\tau$ we can reproduce the results of Eq. (13) as particular cases.

(ii) For $d = 4$, the substitution $\Psi = u/\sin \theta$ simplifies Eq. (12) to $u'' + [\eta_0 + (\eta_0 + 1)^2]u = 0$, solved by a linear combination of $\sin[\eta_0 + 1\theta]$ and $\cos[\eta_0 + 1\theta]$, and we find

$$
\eta_{c0} = \frac{\Theta}{\pi - \Theta} \quad \text{and} \quad \eta_{c0} = \frac{\pi + 2\Theta}{\pi - 2\Theta},
$$

depicted by the bottom and top dashed lines, respectively, in Fig. 3. The cone exponent $\eta_{c0}$ vanishes linearly with $\Theta$; a needle in four dimensions is “invisible” to a random walker.

As in the two-dimensional case, we may generalize to two touching cones aligned along common axis with apex angles $\Theta_1$ and $\Theta_2$, with

$$
\eta_{c0} = \frac{\Theta_1 + \Theta_2}{\pi - (\Theta_1 + \Theta_2)}.
$$

Unfortunately, when one of the cones is tilted, the problem loses its azimuthal symmetry and Eq. (12) with its simple solutions is no longer valid.

(iii) For $d = 3$, Eq. (12) becomes

$$
(1 - \mu^2) \frac{d^2 \Psi}{d\mu^2} - 2\mu \frac{d\Psi}{d\mu} + \mu_0(\eta_0 + 1)\Psi = 0,
$$

where $\mu = \cos \theta$. The general solution to this equation is given by regular (rather than associated) Legendre functions

$$
\Psi(\theta) = a_1 P_{\eta_0}(\mu) + a_2 Q_{\eta_0}(\mu).
$$

Note that $P_{\eta}(1) = 1$ for any $\alpha$, while $P_{\eta}(-1)$ diverges for noninteger $\alpha$. Similarly, $Q_\eta(\pm 1)$ is divergent. The linear combination in Eq. (18) can be made regular at $-1$ by a proper choice of $a_1/a_2$. For the geometry described in Fig. 1(a) the solution must be regular for $\theta \leq \theta \leq \pi$. Instead of using combinations of $P$ and $Q$, we can simply use $P_{-\eta}(-\cos \theta)$,

which will be regular at $\cos \theta = -1$. The value of $\eta_0$ is then determined by requiring

$$
P_{\eta}(\cos \theta) = 0.
$$

Since $\Psi$ cannot change sign in the physically permitted region, the smallest possible $\eta_0$ must be chosen. The corresponding solution has been described in detail in Ref. [46]. In particular, in the limit $\Theta \to 0$, the exponent $\eta_0$ vanishes through a logarithmic singularity ($-1/|\ln \Theta|$), while for $\Theta \to \pi$ it diverges as $\eta_0 \approx 2.405/(\pi - \Theta)$.

For the geometry of a polymer attached to the apex of a cone touching an infinite plane [Fig. 1(c)], we need to solve Eq. (17) with $\Psi(\theta)$ vanishing at both $\Theta$ and $\pi/2$. The latter condition corresponds to $\mu = 0$ and can be assured by setting in Eq. (18) the ratio

$$
\frac{a_2}{a_1} = -\frac{P_{\eta}(0)}{Q_{\eta}(0)} = \frac{2}{\pi \tan(\pi \eta_0/2)}.
$$

With the above choice, the angular function can be conveniently written as

$$
\Psi(\theta) = \pi \sin(\pi \eta_0/2)P_{\eta}(\cos \theta) + 2 \cos(\pi \eta_0/2)Q_{\eta}(\cos \theta).
$$

For noninteger $\alpha$, $P_{\eta}(\mu)$ and $Q_{\eta}(\mu)$ are linearly independent and both solve Eq. (17) [48]. Thus, for noninteger $\eta_0$, Eq. (21) can be replaced by

$$
\Psi(\theta) = P_{\eta}(\cos \theta) - P_{\eta}(\cos \theta).
$$

Again, we must choose the smallest $\eta_0$ for which this function vanishes for $\theta = \pi$. The resulting exponents (which cannot be cast as simple functions) are plotted as solid lines in Fig. 3. From bounds on the roots of Legendre polynomials [49–51], we find that in the limit $\Theta \to \pi/2$, the exponent diverges as $\eta_0 \approx 2\pi/(\pi - 2\Theta)$, while as $\Theta \to 0$, $\eta_0 \to 1$. Indeed, the solution in Eq. (21) has no roots for $\eta_0 < 1$. Asymptotic expressions for $P_{\eta}(\mu)$ and $Q_{\eta}(\mu)$ near $\mu = 1$ are known and can be used to determine positions of roots in such limit, indicating that $\eta_0 \approx 1 + 1/|\ln(\Theta)|$.

It is worth noting that in all cases above ($d = 2, 3, 4$) the cone + plane exponent has the identical divergence $\eta_0 \approx 2\pi/(\pi - 2\Theta)$ as $\Theta \to \pi/2$. This may be justified by arguing that a diffuser confined between an almost flat cone and the plate encounters absorbing boundaries not dissimilar to that of a wedge.

(iv) In general $d$, with the substitution $\Psi(\theta) = \sin^{-d} \theta u(\mu)$, where $\mu = \cos \theta$ and $\delta = (d - 3)/2$, Eq. (12) becomes

$$
(1 - \mu^2) \frac{d^2 u}{d\mu^2} - 2\mu \frac{d u}{d\mu} + \left[\eta_0 + (\eta_0 + 1)^2\right] u = 0,
$$

which is solved by associated Legendre functions [48] that can be chosen in several forms such as $P_{\eta_0+d}(\pm \mu)$, $Q_{\eta_0+d}(\pm \mu)$, or their combinations. The particular choice of one or a pair of linearly independent functions depends on the boundary conditions and the specific values of $\eta_0$ and $\delta$. For the case of cone in different dimensions the reader is referred to Ref. [46]. To find a proper form for $u$ in (integer or fractional) $d$ dimensions, we can closely follow the case of $d = 3$:
For a single cone we may choose \( u = P_{n0}^{-\delta}(-\cos \theta) \), since \( \sin^{-\delta} \theta P_{n0}^{-\delta}(-\cos \theta) \) has no cusp at \( \theta = \pi \). The value of \( \eta_0 \) will be set by the requirement that the function vanishes for \( \theta = \Theta \). For cone + plate configurations we may choose
\[
u = a_1 P_{n0}^{-\delta}(-\cos \theta) - a_2 P_{n0}^{-\delta}(\cos \theta),
\]
which for \( a_1 = a_2 \) vanishes on the plate (\( \theta = \pi \)), and will be a suitable solution as long as \( \eta_0 \) is not integer and is chosen such that the function vanishes for \( \theta = \Theta \).

For all \( d \), the cone becomes a plate for \( \Theta = \pi/2 \). Correlations with one point approaching a surface are easily obtained by the method of images [30,52] leading to \( \eta_p = \eta_0 = 1 \), which is clearly seen from the intersection of the curves in Fig. 3.

Note that in \( d = 3 \) both cone and cone + plate exponents approach their limit \( \Theta \rightarrow 0 \) via a logarithmic singularity, while for \( d = 4 \) they approach that limit linearly. For intermediate dimensions \( 3 < d < 4 \), the limiting behavior as \( \Theta \rightarrow 0 \) is given by
\[
\eta_p = \frac{\Gamma(1 - \epsilon/2)}{\sqrt{\pi} \Gamma(1/2 - \epsilon/2)} \Theta^{1-\epsilon} \quad \text{and} \quad \eta_{cp0} = 1 + \frac{4\Gamma(2 - \epsilon/2)}{\sqrt{\pi} \Gamma(1 - 2 - \epsilon/2)} \Theta^{1-\epsilon},
\]
where \( d = 4 - \epsilon \). The singular behavior in the above equations is a power law \( \Theta^{p0} \) with \( p_0 = d - 3 \), which has a simple geometric interpretation. The ideal polymer can be regarded as a self-similar object with fractal dimension 2, while the remnant of the cone as \( \Theta \rightarrow 0 \) is a self-similar object with fractal dimension 2, while the intersection of the two entities (random walk and remnant line) spans a space of dimension \( d - (2 + 1) = p_0 \). Indeed, for \( d < 3 \) the limiting value is different from the case without any cone indicating the finite probability of intersection of the polymer with the semi-infinite barrier line.

In both \( d = 2 \) and \( d = 4 \), we found simple expressions for the ideal polymer attached to the contact point between two coaxial cones of opening angles \( \Theta_1 \) and \( \Theta_2 \). However, in general \( d \) the solution has the form in Eq. (24). The vanishing of \( u \) on the first cone requires \( a_1/a_2 = \cos(\Theta_1)/\cos(-\Theta_1) \). With this choice, the only free parameter left is the exponent \( \eta_0 \). To satisfy the condition \( u(\theta = \Theta_2) = 0 \), we must have
\[
P_{n0}^{-\delta}(-\cos \Theta_1) P_{n0}^{-\delta}(-\cos \Theta_2) - P_{n0}^{-\delta}(\cos \Theta_1) P_{n0}^{-\delta}(\cos \Theta_2) = 0.
\]

Unfortunately, values of \( \eta_0 \) satisfying the above equation cannot be expressed as simple functions. However, for small \( \Theta_1 \) and \( \Theta_2 \), we can prove that
\[
\eta_{col}(\Theta_1, \Theta_2) \approx \eta_{col}(\Theta_1) + \eta_{col}(\Theta_2),
\]
that is, the exponent for two sharp cones is approximately the sum of exponents for individual cones of opening angles \( \Theta_1 \) and \( \Theta_2 \).

V. Self-Avoiding Polymers: Simulations

Self-avoiding walks (SAWs) provide a convenient model for exploring universal aspects of swollen (coil) polymers with short-range interactions. The number of SAWs in three-dimensional space (without obstacles) is governed by the critical exponent \( \gamma \approx 1.158 \) [33] (corresponding to \( \eta \approx 0.03 \), while in two dimensions, \( \gamma = 43/32 \) [53] \( (\eta = 5/24) \). (For ideal polymers these exponents do not depend on \( d: \gamma_0 = 1 \) and \( \eta_0 = 0 \).) Several important results regarding \( \gamma \) (with obstacles) for polymers confined by wedges or planes are known [31,34,54–57]. For example, the exponent \( \gamma_{\text{wedge}} \) of a SAW confined to a wedge (in \( d = 2 \) or \( d = 3 \)), and anchored at its sharp end, depends on opening angle and diverges to \( -\infty \) as the confining angle vanishes. Polymers attached to the tip of a two-dimensional sector in \( d = 3 \), and to the apex of a cone also were studied [58]. Numerous analytical [59,60] and numerical [34,57] studies of SAWs anchored to flat surfaces in \( d = 3 \) find \( \gamma_0 = \gamma_f \) in the range of 0.70 [57] to 0.68 [61] \( (\eta_0 = 0.81 \) or 0.84, respectively). (Again, for ideal polymers these exponents do not depend on \( d: \gamma_p = 1/2 \) and \( \eta_p = 1 \).) We are particularly interested in geometries depicted in Fig. 1, and provide below our numerical and analytical estimates of the relevant exponents.

We employed a dimerization method [62,63] to numerically generate SAWs on a cubic lattice. In this (recursive) method two \( N/2 \)-step SAWs are joined in an attempt to create an \( N \)-step walk. If the resulting walk does not self-intersect, the process is successful and the \( N \)-step walk is used to generate even larger SAWs. If the combined walk has intersections, it is rejected and both \( N/2 \)-step components are discarded. The process is then repeated. As the rejection rate increases slowly with \( N \), this is an extremely efficient method for creating an unbiased collection of SAWs. We generated \( 10^8 \) SAWs of lengths \( N = 16,32, \ldots, 1024 \). Each SAW was attached to the origin, and we checked whether it intersects the imposed obstacles.

The probability that a SAW does not intersect the confining boundaries is the ratio of the number of SAWs satisfying the geometrical constraints to the total number of such walks, that is, \( p_N = N^{\gamma} / N^{N_{\text{wall}}-1} \), and therefore \( p_N / p_{2N} = 2^{\Delta \eta} \). This result becomes accurate for \( N \rightarrow \infty \), and we extract the limiting value of \( \gamma \) by plotting its finite-\( N \) estimates as a function of \( 1/\sqrt{N} \). The estimated errors in the extrapolated values of the exponents are caused both by the limited sample size and by values of \( N \) that are not long enough. The resulting exponents \( \Delta \eta \equiv \eta_p - \eta \equiv (\nu - \gamma)/\nu \) (solid symbols in Fig. 4) are somewhat lower than the corresponding values for ideal polymers (indicated by dotted lines). The difference primarily originates in the change in \( \eta_p \) as a result of self-avoiding interactions. (This trend is slightly moderated by a small shift of \( \eta \) in free space.) Figure 3 of Ref. [26] presents \( \Delta \gamma \) as a function of \( \Theta \), in which form results for SAWs are much closer to those of ideal polymers due to multiplication by the self-avoiding and ideal values of \( \nu \), respectively.

The dashed line in Fig. 4 represents the result of an \( \epsilon = 4 - d \) expansion [58] that treats both the self-repulsion of the polymer as well as its repulsion by the two-dimensional surface of the cone, as weak perturbations. A renormalization group computation is then carried out to the lowest order in \( \epsilon \), resulting in \( \Delta \eta = (3\epsilon/4) \sin \Theta \). For small \( \Theta \) this expression resembles the expected behavior, but for \( \Theta > \pi/2 \) the function (incorrectly) decreases. This is because, faced with the weakly repulsive potential at the surface of the cone, the polymer simply jumps from the decreasing exterior into the larger internal space of the cone. This approach also produces
an incorrect Θ → 0 behavior. Deficiencies of the results of Ref. [58] can be remedied by exclusion of the entire interior of the cone, as was done by Cardy [31,55] for the wedge geometry. The analogous computations for a cone, described in the following section, are more complicated and build upon

\[ G_0(x,x') = \sum_{n,m} \frac{(-1)^l}{2 \sin(\rho_n \pi \Gamma(\rho_n - l))} \int_{\rho_n-1/2}^{\rho_n+1/2} \frac{\rho_{n,m}^{l-1/2}}{\rho_{n,m}^{l-1/2} (1 - \cos \theta)} r_{n,m}^{l-1/2} \delta_{n,m}^{l-1/2}(\psi,\phi) \]

where the summation is made over a triplet of integers \( n > 0, l \geq 0, \) and \(-l \leq m \leq +l\). The exponent \( \rho_n \) labeled by the integer \( n \) is the root of the equation

\[ P_{\rho_n-1/2}^{-1/2} (1 - \cos \theta) = 0. \]

In the above equation, \( r^< \) and \( r^> \) refer to the smaller and larger radial coordinate for \( x \) and \( x' \). Such division of the Green’s function is appropriate since one endpoint of the polymer is close to the tip of the cone and the other end is far away.

For an interacting Green’s function, one should subtract polymer configurations which self-intersect. In the perturbative analysis, the strength of self-repulsion is indicated by a parameter \( u_r \), and to lowest order one must subtract contributions forming a single intermediate loop, such that

\[ G_1 = G_0 - u \int d^4x' G_0(x,x') G_0''(x'',x'') G_0(x'',x'). \]

Similar expansions arise in the context of quantum field theories, in which language the first term in the last equation is the “free” propagator (in the presence of external boundary conditions), while the next term is the “one-loop” correction. The latter should be regularized by replacing the middle Green’s function (computed at identical points) with the difference between the Green’s functions in the presence and absence of external boundaries, that is, \( G_0'' = G_0 - G_0'' \). In a renormalization group treatment, the parameter \( u \) changes with scale of observation, ultimately arriving at a fixed point value of \( u^* = 2\pi^2 \epsilon \). This universal value is a characteristic of the polymer and independent of obstacles or boundary conditions. To find the scaling behavior for \( r \gg a \), it is sufficient to include only the term \( n = 1, l = m = 0 \) from the sum in Eq. (29) in the nonloop propagators, arriving at

\[ G_0_{l,m=0}(x,x') = \frac{\rho_{n,m}^{-l-1}}{4\pi^2} \left( \frac{\sin \rho_1 (\pi - \theta)}{\sin \theta'} \frac{\sin \rho_1 (\pi - \theta')}{\sin \theta} \right). \]

(32)

(Corrections from higher-order terms are in higher powers of \( a/r \), similar to the analysis by Cardy [31,55].) However, the
loop (Green’s function at identical points) can be of small size; in fact, we have regularized this Green’s function to cancel out the divergent contribution in the limit of a shrinking loop. Therefore, for the loop propagator one should consider the entire sum.

The integral corresponding to the loop variable is over the whole space: The azimuthal and spherical angles \( \phi’’ \) and \( \psi’’ \) are trivial due to rotational symmetry. Once summed over spherical harmonics indexed by the same \( l \), we get a factor of \( l(l+1) \). The integral over the radial coordinate, \( r’’ \), should produce a logarithm, later to be exponentiated to a power law. Along the line of Cardy’s analysis, this logarithm is due to the contribution from the region \( r < r’’ < r’ \). Finally, an integration over \( \theta’’ \), the polar angle, completes the integral; however, the latter is complicated by the fact that the roots of the nonalgebraic Eq. (30) should be computed.

Considering the above complications, we focus instead on a cone with small opening angle, that is, \( \theta \to 0 \). In this limit, one can see that the leading singularity is due to the \( l = 0 \) term in the Green’s function corresponding to the loop. Note that a sharp cone is invisible to higher spherical partial waves. The loop Green’s function is then obtained from the sum

\[
G_{l=0}^{\phi} \left( x’, x \right) = \frac{1}{4\pi^2 \rho c^2} \sum_{n=1}^{\infty} \int \frac{n^2 \pi^{n-1} \sin^2 \theta n^{n-1}}{\sin^2 \theta n^{n}}.
\]

Equations (32) and (33) are used to perform the integral over \( \theta’’ \) and summation over all \( n \). The Green’s function dependence on the radius is then obtained as

\[
G_1 \propto \frac{r_{\phi}^{n_0}}{r_{\phi}^{n_0+1/2}} \left[ 1 + \epsilon \ln \frac{r_{\phi}}{r_{\phi}^{n_0}} \left( \frac{4\pi}{\rho} \ln \Theta + 0.16 \Theta \right) \right],
\]

where the second term in the bracket is the correction due to the loop integral, and thus it is proportional to \( \epsilon \). Another dependence on \( \epsilon \) is introduced through the exponent \( n_{c0} \) whose value in \( d = 4 - \epsilon \) dimensions is given by Eq. (25). The radial logarithm can be exponentiated yielding the renormalized exponent,

\[
\eta_c = \eta_{c0} + \left( \frac{\ln \Theta}{4\pi} + 0.16 \Theta \right) \epsilon.
\]

As the first correction to \( \eta \) in empty space appears at the order of \( \epsilon^2 \), the above (first-order) result for the cone vanishes logarithmically as \( \Theta \to 0 \). The logarithm in the angular variable is suggestive of another exponentiation to obtain a power-law in the limit of \( \Theta \to 0 \). First note, however, that expanding \( \eta_{c0} \) to the first order in \( \epsilon \) yields

\[
\eta_{c0} \approx \frac{\Theta}{\pi} + \left( \frac{\ln \Theta}{\pi} - 0.22 \Theta \right) \epsilon,
\]

producing another contribution to \( \ln \Theta \), originating from the expansion in \( 4 - \epsilon \) dimensions of the phantom polymer (as opposed to the perturbative terms from the one-loop computation). Putting all these pieces together, we obtain

\[
\eta_c \approx \frac{1}{\pi} \left( 1 - \frac{3}{4} \epsilon \ln \Theta - 0.06 \epsilon \right),
\]

which can be recast as a power law

\[
\eta_c \approx \frac{1}{\pi} \left( 1 - 0.06 \epsilon \right) \Theta^{1 - \frac{1}{2} \epsilon}.
\]

As expressed in Eq. (25), the exponent \( \eta_{c0} \) vanishes with cone angle as \( \Theta_0^{\rho_0} \) with \( \rho_0 = d - 3 = 1 - \epsilon \) for the phantom polymer (Fig. 3). The above equation indicates that the vanishing of \( \eta_c \) for a self-avoiding polymer is governed by the modified exponent \( p = 1 - 3\epsilon/4 \). This dependence is shown for \( \epsilon = 1 \) as the lower solid line in Fig. 4. We may interpret this result as follows: The self-avoiding condition swells the polymer at all scales compared to a random walk. As a result, the fractal dimension of the polymer is reduced from 2 to \( \nu^{-1} = 2 - \epsilon/4 \) at the linear order in \( \epsilon \). Whether or not a fractal coil and the one-dimensional needle intersect depends on the dimensionality \( d \) of space; the domain of their intersection is given by the codimension of the polymer + needle, that is, \( d - 1 - \nu^{-1} = 1 - 3\epsilon/4 \approx p \). It would be interesting to see if this connection holds in higher orders in perturbation theory.

The current numerical results are not accurate enough to test this conjecture.

Finally, we note that for two touching cones of small angles \( \Theta_1 \) and \( \Theta_2 \), we find the analog of Eq. (27) to first order in the \( \epsilon \) expansion, namely,

\[
\eta_{cc}(\Theta_1, \Theta_2) \approx \eta_c(\Theta_1) + \eta_c(\Theta_2).
\]

B. Cone-plate

When constrained by a cone touching a plate, the Green’s function is given by

\[
G_0(x, x’) = \sum_{n=1}^{l} \left( \frac{(-1)^l \pi}{2} \frac{\Gamma(n_0 + l + 1)}{n_0 \rho_0 l} \right) \frac{P_{n_0-1/2}^{-l-1/2} \cos \Theta}{\sin \Theta} \frac{\partial_{\rho_0} \left[ P_{n_0-1/2}^{-l-1/2} \cos \Theta - P_{n_0-1/2}^{-l-1/2} \cos \Theta \right]}{\sin \theta} \frac{Y_m(\psi, \phi)}{\sqrt{\sin \theta}} Y_m(\psi’, \phi’),
\]

where the sum is again over the triplet of integers \( n > 0, l \geq 0 \), and \(-l \leq m \leq +l\). The exponent \( \rho_0 \) is now the \( n \)th root of the transcendental equation

\[
P_{n_0-1/2}^{-l-1/2} \cos \Theta - P_{n_0-1/2}^{-l-1/2} \cos \Theta = 0.
\]
Again for the nonloop propagators, it suffices to keep only the first term in the sum
\[
G_{0}^{[n=1]}(x,x') = \frac{r_0^{n-1}}{4\pi^2 r_{\Theta}^{n+1}} \sin \theta \sin \theta' \sin \theta 
\]
(42)
However, the intermediate loop should be summed entirely. Here again, we focus on a sharp cone. It is more convenient to express the regularized Green’s function for the intermediate loop as
\[
G'_{0} \equiv G_{0} - \tilde{G}_{0} = (G_{0} - G_{0}') + (G_{0}' - \tilde{G}_{0}),
\]
(43)
where \(G_{0}'\) is the Green’s function in the presence of the plate alone and is independent of the cone angle \(\Theta\). Also note that \(\tilde{G}_{0}\) is the Green’s function in empty space. Each bracket on the right-hand side of Eq. (43) can be treated separately. The first bracket, to the leading order for a sharp cone, is given by
\[
\left( G_{0} - G_{0}' \right)_{l=m=0}(x',x'') = \frac{1}{4\pi^2 r_{\Theta}^2} \sum_{n=1}^{\infty} \frac{1}{n} \frac{\sin^2 \theta}{\sin^2 \theta''} - \frac{2}{\sin^2 \theta''}.
\]
(44)
Also, the second bracket in Eq. (43) is merely the Green’s function in the presence of a plate. The latter can be cast as a series expansion by using the method of images:
\[
(G_{0} - \tilde{G}_{0})(x,x') = -\sum_{n_{l} m_{l} > 0}^{\infty} \Gamma(n_{l} + 1) \frac{P_{n_{l} - 1/2}(\cos \theta)}{2\Gamma(n_{l} - 1)} \times P_{n_{l} - 1/2}(\cos \theta') Y_{l_{m}}(\psi,\phi) Y_{l_{m}}^{*}(\psi',\phi').
\]
(45)
The Green’s functions appropriate to the external legs and the loop should be inserted in Eq. (31) to obtain, in the limit of \(\Theta \to 0\),
\[
G_{1} \propto \frac{r_{\Theta}^{n_{\Theta}}}{r_{\Theta}} \left[ 1 + \frac{1}{\pi} \ln \frac{r_{\Theta}}{r_{\Theta}^{e}} + \frac{1}{8} + \frac{1}{\pi} \ln \Theta + 0.66 \Theta \right].
\]
(46)
where the exponent \(n_{\Theta}^{p_{0}}\) is given by Eq. (25). Upon exponentiation in \(r\), we find
\[
\eta_{cp} = \eta_{cp0} + \left( -\frac{1}{8} + \frac{1}{\pi} \ln \Theta + 0.66 \Theta \right) \epsilon.
\]
(47)
Note that the loop correction does not vanish for \(\Theta \to 0\), instead going to the limiting value of \(-\epsilon/8\) due to the presence of the plate. Expanding \(\eta_{cp0}\) to the first order in \(\epsilon\),
\[
\eta_{cp0}(\Theta,\epsilon) \approx 1 + \frac{4\Theta}{\pi} + \left( -\frac{4}{\pi} \ln \Theta - 1.52 \Theta \right) \epsilon,
\]
(48)
we find a logarithmic dependence in \(\Theta\). Summing both contributions, the exponent \(\eta\) to the lowest order in \(\epsilon\) and \(\Theta\) becomes
\[
\eta_{cp} = 1 - \frac{\epsilon}{8} + \frac{4\Theta}{\pi} \left( 1 - \frac{3}{4} \epsilon \ln \Theta - 0.86 \epsilon \right).
\]
(49)
As in the previous section, we exponentiate the logarithm in \(\Theta\) to obtain the power-law dependence
\[
\eta_{cp} \approx 1 - \frac{\epsilon}{8} + \left( \frac{4}{\pi} - 0.86 \epsilon \right) \Theta^{-1/4}.
\]
Revealingly, the exponent \(p = 1 - 3\epsilon/4\) is the same as in the case of a single cone, whereas the amplitude has changed. Furthermore, it approaches a constant value in the limit of vanishing angle. This result is shown for \(\epsilon = 1\) as the upper solid line in Fig. 4.

Finally we can compute the amplitude of the force according to Eq. (5), using Eqs. (38) and (50), as
\[
A = 1 - \frac{\epsilon}{8} + \left( \frac{3}{2\pi} - 0.80 \epsilon \right) \Theta^{1/4}.
\]
(51)

C. Star polymers

In a typical setup, the amplitude \(A = \eta_{cp} - \eta_{cp0}\) with the exponents being computed in the previous subsections is of order unity. Therefore, according to Eq. (1), the force is roughly 0.1 pN at room temperature in a separation of 0.1 \(\mu\)m. Such a force is at the margin of measurement by current precision apparatus. We can further increase the force by attaching more polymers to the cone tip. The total force is additive for ideal (phantom) polymers; that is, it is proportional to \(f\), the number of arms. For self-avoiding polymers, however, interactions come into play and the result is no longer additive.

In general, there are two types of interactions: a single arm can self-intersect (intra-arm interaction) or two different arms intersect (interarm interaction). The former effect leads to corrections similar to those computed in previous sections and would by itself simply lead to multiplication of the force on a single polymer by \(f\). The latter interaction, originating from intersections between two different arms, is proportional to \(f(f - 1)/2\), the number of interacting pairs. We can then write the overall exponent as \(\eta_{cp} = f \eta_{cp0} + f_{i} \eta_{i} + f_{o} \eta_{o} \theta^{1/4}\), where \(\eta_{o}\) characterizes a single phantom polymer, and \(\eta_{i}\) and \(\eta_{o}\) correspond to intra- and interarm interactions, respectively. A general situation of many-arm polymers in the absence of external boundaries is considered in detail in Ref. [66].

To zeroth order (i.e., in the absence of interactions) the correlation function describing a pair of polymers starting at the same point \(x\), but with different end points at \(x'_{1}\) and \(x'_{2}\), is simply the product of two free Green’s functions, \(G_{0} \otimes G_{0}\). Subtracting configurations in which the two polymers avoid each other yields first to order the interacting Green’s function
\[
G_{1}^{2/5} = G_{0} G_{0} - u \int d^{4}x'_{3} G_{0}^{2}(x,x'_{3})G_{0}(x',x_{1})G_{0}(x',x'_{2}).
\]
(52)
where we have integrated over the intersection point \(x'_{3}\). The prime on the Green’s function indicates that we have only considered the interarm interactions. Also note that regularization is needed since the integral in Eq. (52) is finite. For a long polymer, it is sufficient to include only the first term in the Green’s function, that is, Eq. (32) for a single cone and Eq. (42) for a cone-plate configuration. The leading
logarithmic contribution for a cone is then
\[ G_1^{(2)} = G_0 G_0 \left[ 1 + \epsilon \ln \frac{r}{r_{\text{max}}} \right] \left( \frac{1}{4} - \frac{\Theta}{2\pi} \right) \].

(53)

where \( r_{\text{max}} \) depends only on \( r_1' \) and \( r_2' \). Hence, by exponentiating the radial function, \( \eta'' \) for a cone is obtained as
\[ \eta'_c = \epsilon \left( \frac{1}{4} - \frac{\Theta}{2\pi} \right) \].

(54)

The corresponding \( \epsilon \) expansion for cone + plate gives
\[ \eta'_{cp} = \epsilon \left( \frac{1}{4} - \frac{7\Theta}{3\pi} \right) \].

(55)

These equations then lead to the force amplitude
\[ \frac{A(f)}{f} = 1 - \frac{\epsilon}{8} + \left[ \frac{3}{\pi} - \left( 0.80 + \frac{11}{12\pi} (f - 1) \right) \right] \Theta^{1-3\epsilon/4}. \]

(56)

Interestingly, interarm interactions reduce the force amplitude per polymer. This equation is similar to Eq. (51) with the addition of interarm interactions. Note that the same exponent \( p = 1 - 3\epsilon/4 \) dictates the limiting behavior in Eq. (56) as \( \Theta \to 0 \).

Yet another possible experimental setup is a system consisting of a polymer attached to a cone which is approaching another cone. The entropic force is relatively smaller in this case. We have verified that the exponent in the latter case also vanishes with the opening angle as \( \Theta^p \), providing further support for the conjecture that this exponent reflects the intersection of a fractal polymer and a needle.

VII. DISCUSSION

In summary, we have demonstrated that polymers exert an entropic force \( \mathcal{A} k_B T / h \) on a cone tip, with a “universal” amplitude \( \mathcal{A} \) dependent on geometry, interactions, and internal topology of the polymer. We conjecture that the singular behavior of the amplitude on vanishing cone angle is described by a new exponent, simply related to the fractal dimension of the polymer. There are many other self-similar shapes where a similar force law is expected on the basis of scaling at length scales shorter than an appropriate correlation length.

In this work we concentrated on situations where the polymer is attached to a single surface such as described by Fig. 1(b) or 1(d). However, our results can be easily applied to a variety of other situations. First, we note that our approach is easily generalized to a slightly more complicated situation with a variety of other situations. First, we note that our approach is easily generalized to a slightly more complicated situation with a variety of other situations. First, we note that our approach is easily generalized to a slightly more complicated situation with a variety of other situations.

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APPENDIX: GREEN’S FUNCTIONS

We first derive the Green’s function for the Helmholtz equation in four dimensions; the corresponding function for the Laplace equation is then obtained as a limit. The former satisfies the Helmholtz equation with a \( \delta \) function source:
\[ - (\nabla_i^2 + k^2) G_0(x,x') = \delta(x - x'); \]

(A1)

the latter is obtained by setting the wave vector \( k \) to zero. Consider spherical coordinates \( x = (r,\theta,\psi,\phi) \), where \( r \) is the radius, \( \theta \) is the polar angle, and \( \phi \) is the remaining angular coordinates; a constant \( \theta \) specifies the surface of a cone in four dimensions. The cone is spherically symmetric with respect to \( \psi \) and \( \phi \) (similar to the azimuthal symmetry of the three-dimensional cone). We expand the Green’s function as
\[ G_0(x,x') = \sum_{lm} \frac{1}{r r'} \frac{1}{\sin\theta' \sin\theta} g_{lm}(r,\theta,r',\theta') \times Y_{lm}(\psi',\phi'). \]

(A2)

The prefactors are introduced for later convenience. With this ansatz, Eq. (A1) takes the form
\[ - \left[ \nabla_{(2)}^2 + \left( l + \frac{1}{2} \right)^2 + k^2 \right] g_{lm} = \frac{\delta(r - r')}{r^2} \delta(\cos \theta - \cos \theta'). \]

(A3)

Note that the Laplacian in the above equation acts in two dimensions, that is, \( \nabla_{(2)}^2 = \frac{1}{r^2} \partial_r r^2 \partial_r + \frac{1}{\sin\theta} \partial_\theta \sin\theta \partial_\theta \).

Given that the boundary is of conical shape, it is convenient to break up the Green’s function in the coordinate \( \theta \). We should then find a completeness relation in the function space of the other variable \( r \). This is obtained by the Kontrovich-Lebedev transform [65]
\[ \frac{1}{\pi} \int_0^\infty d\kappa \sinh \kappa k_{i\kappa-1/2}(r) k_{i\kappa+1/2}(r') = \delta(r - r'), \]

where \( k_\nu \) is the spherical Bessel function of order \( \nu \). Using this relation, one can show that
\[ g_{lm} = \frac{2}{\Gamma(-l + 1)} \left( \frac{1}{r} \phi \right)^{1/2} \left( \frac{1}{r'} \phi' \right)^{-1/2} \times P_{i\kappa-1/2}^{1/2}(\cos \theta) P_{i\kappa+1/2}^{-1/2}(\cos \theta). \]

(A4)
solves Eq. (A3). Here, $\kappa$ is the imaginary frequency ($k = i\kappa$), $\theta_\omega = \min(\theta, \theta')$, and $\theta_\infty = \max(\theta, \theta')$. Note that this equation satisfies the Helmholtz equation in empty space. We discuss the boundary condition below. The completeness relation, together with an identity regarding the Wronskian of Legendre functions, can be exploited to see that Eq. (A4) indeed solves Eq. (A3). By a construction similar to that in Ref. [64], we can analytically continue the complex order of the Bessel and Legendre functions to the real axis to obtain

$$g_{lm} = \kappa \sum_{n=l+1}^{\infty} \frac{i \Gamma(n + l + 1)}{\Gamma(n - l)} \frac{\Gamma(n + l + 1)}{\Gamma(n - l)} \times P_{n-1/2}(-\cos \theta)P_{n-1/2}(-\cos \theta'). \quad (A5)$$

Note that the asymmetry has shifted to the radial variable while the angular coordinates are treated symmetrically. We have used the symmetry $P_{n+1/2}(-x) = (-1)^{n+1} P_{n-1/2}(x)$ for integers $l < n$ to restore the latter symmetry. Next we take the limit $\kappa \to 0$ to find

$$G_0(x, x') = \sum_{n=1}^{\infty} \sum_{l=0}^{\infty} \frac{\Gamma(n + l + 1)}{2\Gamma(n - l)} \frac{\Gamma(n + l + 1)}{\Gamma(n - l)} \times P_{n-1/2}(-\cos \theta')Y_{lm}(\psi, \phi)Y^*(\psi', \phi'). \quad (A6)$$

This equation provides the Green’s function for the Laplace equation in empty space. For the Green’s function in the presence of the conical obstacle, we modify Eq. (A4) to

$$G_0(x, x') = \sum_{n=1}^{\infty} \sum_{l=0}^{\infty} \frac{\Gamma(n + l + 1)}{2\Gamma(n - l)} \frac{\Gamma(n + l + 1)}{\Gamma(n - l)} \times P_{n-1/2}(-\cos \theta')Y_{lm}(\psi, \phi)Y^*(\psi', \phi'). \quad (A8)$$

The integer $n$ represents the $n$th root of the transcendental equation

$$P_{n-1/2}(-\cos \theta) = 0. \quad (A9)$$

The solution for $\rho_n$ also depends on $\Theta$ and $l$. In the limit of $\Theta \to 0$; that is, for a sharp cone, it can be shown that [67]

$$\rho_n \approx l + n + \frac{\Gamma(2l + n + 1)}{\Gamma(l + 3/2)\Gamma(l + 1/2)\Gamma(n)}(\tan \Theta)^{2l+1}, \quad (A10)$$

and the asymptotic behavior in this limit is dominated by the root for $l = 0$. To proceed further, we take advantage of the identities

$$P_{-1/2}(\cos \theta) = \frac{1}{\rho} \sqrt{\frac{2}{\pi \sin \theta}} \sin(\rho \theta), \quad (A11)$$

$$P_{-1/2}(-\cos \theta) = \frac{1}{\rho} \sqrt{\frac{2}{\pi \sin \theta}} \sin[\rho(\pi - \theta)],$$

the second of which dictates

$$\rho_n = \frac{n\pi}{\pi - \Theta} \quad (A12)$$

Hence, using Eqs. (A8)–(A12), we find, for a single cone,

$$G_0|_{l=m=0}(x, x') = \frac{1}{4\pi^2} \sum_{n} \frac{\rho_n^{-1}}{\rho_n^{0}} \frac{\sin \rho_n(\pi - \theta) \sin \rho_n(\pi - \theta')}{\sin \theta \sin \theta'}. \quad (A13)$$

We can similarly find the Green’s function in the space between two cones aligned along a common axis which touch at their tips. Designating the opening angles by $\Theta_1$ and $\pi - \Theta_2$, the available space is characterized by the polar angle $\Theta_1 < \theta < \Theta_2$. The Green’s function (for the Helmholtz equation) is similar to Eq. (A4) with the substitution of $P_{n+1/2}(-\cos \theta_2)P_{n-1/2}(-\cos \theta_2)$ by

$$P_{n+1/2}(\cos \theta_2)P_{n+1/2}(-\cos \theta_2) \quad (A11)$$

$$P_{n-1/2}(\cos \theta_2)P_{n-1/2}(-\cos \theta_2) \quad (A12)$$

and

$$P_{-1/2}(\cos \theta_2)P_{-1/2}(-\cos \theta_2) \quad (A13)$$

Hence, using Eqs. (A8)–(A12), we find, for a single cone,
We then rotate to the real axis where we should find the roots of the transcendental equation
\[ P_{\rho_n-1/2}(-\cos \Theta)P_{\rho_n-1/2}(\cos \Theta_2) - P_{\rho_n-1/2}(\cos \Theta_1)P_{\rho_n-1/2}(-\cos \Theta_2) = 0. \]  \hspace{1cm} (A15)

As we are mainly interested in a cone attached to a plate, we choose \( \Theta_1 \equiv \Theta \) and \( \Theta_2 = \pi/2 \). By rotating to the real axis and taking the limit \( \kappa \to 0 \), the full Green’s function is then obtained as
\[
G_0(x',x) = \sum_{n=0}^{\infty} \sum_{l=-\infty}^{\infty} \frac{\pi}{2} (-1)^l \frac{P_{\rho_n-1}^l(r_\rho)}{r_\rho^{n+1}} \frac{\Gamma(\rho_n + l + 1)}{\sin(\rho_n \pi \Theta_1)} \frac{P_{\rho_n-1/2}^l(\cos \Theta)}{\sqrt{\sin \theta}} \frac{P_{\rho_n-1/2}^l(\cos \Theta_2)}{\sqrt{\sin \theta_2}} Y_{lm}(\psi, \phi)Y_{lm}^*(\psi', \phi').
\]  \hspace{1cm} (A16)

where \( \rho_n \) is the \( n \)th root of the transcendental equation
\[ P_{\rho_n-1/2}(\cos \Theta) = 0. \]  \hspace{1cm} (A17)

For a sharp cone, we can focus on \( l = 0 \), in which case, using Eq. (A11), we find
\[
\rho_n = \frac{n \pi}{\pi/2 - \Theta}.
\]  \hspace{1cm} (A18)

The Green’s function for the cone + plate configuration then reads
\[
G_0|_{x=x_0}(x',x) = \frac{1}{4\pi^2} \sum_{n} \frac{r_\rho^{n+1}}{n r_\rho^{n+1}} \frac{\sin \rho_n(\pi/2 - \theta)}{\sin \theta} \frac{\sin \rho_n(\pi/2 - \theta')}{\sin \theta'}.
\]  \hspace{1cm} (A19)

References: