Adhesion transition of flexible sheets

Arthur A. Evans 1,∗ and Eric Lauga 2,†

1Department of Physics, University of California, San Diego, 9500 Gilman Drive, La Jolla, California 92093-0354, USA
2Department of Mechanical and Aerospace Engineering, University of California, San Diego, 9500 Gilman Drive, La Jolla, California 92093-0411, USA

(Received 8 February 2009; published 25 June 2009)

Intermolecular forces are known to precipitate adhesion events between solid bodies. Inspired by a macroscale experiment showing the hysteretic adhesion of a piece of flexible tape over a plastic substrate, we develop here a model of far-field dry adhesion between two flexible sheets interacting via a power-law potential. We show that phase transitions from unadhered to adhered states occur as dictated by a dimensionless bending parameter representing the ratio of interaction strength to bending stiffness. The order of the adhesion transitions, as well as their hysteretic nature, is shown to depend on the form of the interaction potential between the flexible sheets. When three or more sheets interact, additional geometrical considerations determine the hierarchical or sequential nature of the adhesion transitions.

DOI: 10.1103/PhysRevE.79.066116 PACS number: 46.50.+a, 46.32.+x, 46.70.De, 46.70.Hg

I. INTRODUCTION

As fabrication technology and nanoscale engineering increase in complexity, it becomes vital to understand small-scale interactions between material components. Surface-tension-mediated forces play a large role in self-assembly not only at the macroscale [1] but also for microelectromechanical and nano electromechanical systems, and as such, a large amount of work has been done in studying the adhesive forces involved [2–4]. Carbon nanotubes (CNTs) have attracted significant attention since they were discovered to exhibit novel electrical and mechanical properties, and it has been found that CNTs can adhere to each other under the influence of capillary forces [5–7]. At these scales, fluid-regulated forces are not the only factors that must be examined. Dispersion (or van der Waals) forces may become more important than at larger scales, and the microscopic intermolecular forces of extended media start to have a macroscopic effect on structural stability [8,9].

In addition to the progress in nanotechnology, many biological systems also display adhesion phenomena whose origins can be traced to intermolecular forces. Geckos are known to adhere to smooth surfaces, without any liquid interface. The microscopic arrays of hairs or setae on the base of the gecko foot are therefore believed to be the source of such effective dry adhesion [10–14]. In cellular biology, cytoskeletal morphogenesis is regulated by complex biopolymer networks. This networked series of thin elastic filaments adhere and interact with one another to form a scaffolding for eukaryotic cells. Mechanical properties of macromolecules such as actin filaments or DNA can be measured by force or deflection analysis at small scales, and polymers adsorbed onto a surface or "zipped" to another molecule can be peeled apart by applying optical tweezers or other external pulling forces [15–18].

Most of the research into interactions between materials at these scales involve close range, contact, and sometimes capillary forces, and this is the limit considered by many models and experiments to date [15–17]. However, long-range forces due to fixed charge distributions, polar, or even nonpolar interactions can lead to adhesion events if the right conditions are satisfied. Previous work has characterized the van der Waals attraction between thin flexible objects, both theoretically [19,20] and experimentally [21]. In this paper, we aim to develop an understanding of the physical mechanisms by which long-range interaction forces compete with elasticity in the adhesion of thin flexible structures. We first introduce and motivate the prototypical system of interest using a macroscale experiment showing the hysteretic adhesion of a piece of flexible tape over a plastic substrate. We then develop a model of far-field dry adhesion between two elastic slender sheets, interacting via a power-law potential, and study numerically their relative adhesion. We uncover that phase transitions from unadhered to adhered states occur as dictated by a dimensionless bending parameter representing the ratio of interaction strength to bending stiffness, as well as the form of the interaction potential between the flexible sheets. We then generalize our model in order to study the interactions between several sheets and show that additional geometrical considerations determine the hierarchical or sequential nature of the adhesion transitions in that case.

II. MACROSCALE EXPERIMENT

An example of an adhesion transition between elastic bodies due to long-range interactions may be demonstrated using everyday materials, namely, a piece of adhesive tape and a plastic substrate. As shown in Fig. 1, this tape can be shown to exhibit complex adhesion properties. The tape is first given a static charge distribution by applying it to a piece of plastic and then removing it swiftly. The tape is initially suspended at the distance shown in Fig. 1(a), sticky side away from an uncharged substrate. As the suspension distance is slightly decreased [Fig. 1(b)], the tape becomes weakly attracted to the surface. At a critical distance [Fig. 1(c)], the attraction suddenly pulls the tape completely to the surface, where it lays flat along the majority of the substrate.
Our analysis will be limited to the regime where \( \varepsilon = h/L \ll 1 \). The thickness of each sheet is denoted \( a \).

\[
E = \frac{1}{2} B_1 \int_0^L y_1''(x_1)^2 dx_1 + \frac{1}{2} B_2 \int_0^L y_2''(x_2)^2 dx_2 + \int_0^L \int_0^L V[x_1, x_2, y_1(x_1), y_2(x_2)] dx_1 dx_2,
\]

where \( B_i \) is the bending modulus of the \( i \)-th sheet and \( x_i \) is the horizontal distance. The function \( V \) describes the interaction potential-energy density between the two sheets, as yet unspecified (note that the integration along the widths of the sheets has already been performed formally in \( V \)). Extremizing this functional yields mechanical equilibrium, as shown by the following system of coupled integro-differential equations with boundary terms:

\[
B_1 y_1'' + \int_0^L dx_2 \frac{\partial V}{\partial y_1} = 0, \tag{2}
\]

\[
B_2 y_2'' + \int_0^L dx_1 \frac{\partial V}{\partial y_2} = 0, \tag{3}
\]

\[
y_1'' [y_1(0) = 0, \quad y_1''(0) = 0], \quad y_2'' [y_2(0) = 0, \quad y_2''(0) = 0]. \tag{4}
\]

The boundary conditions are set by the physical conditions of the sheets. While there are many possible cases that could be examined, we will consider the common physical scenario in which the sheets are fixed and clamped on the left \( y_i(0) = 0, \quad y_i''(0) = 0, \quad i = 1, 2 \) and the right edges of the sheets are force- and moment-free \( y_i''(L) = y_i'''(L) = 0, \quad i = 1, 2 \).

The potential \( V \) can be chosen to describe the physical mechanism responsible for the adhesion between the sheets [3,10,17]. In this paper, we are considering a general long-range potential of the form \( V \sim 1/r^n \), where \( r = \sqrt{(x_1-x_2)^2 + (y_1(x_1)-y_2(x_2))^2} \) and \( n \) is a positive integer. More specifically, we set
\[ V = \sum_{n=1}^{N} \frac{A_n W \sigma^n}{(x_1 - x_2)^2 + y_1(x_1) - y_2(x_2)]^{2n+2}}, \tag{6} \]

where \( \sigma \) is the van der Waals-like radius, \( W \) is the strength of the interaction, and \( N \) is the number of singular modes. The sign of \( A_n \) determines whether the interaction is attractive or repulsive. We will examine the more specific form of this general potential where only two terms remain: an attractive term \( n = p \) with \( A_p = -1 \) and a repulsive term \( n = q \) with \( A_q = +1 \). This is the familiar Lennard-Jones-like potential that is used to model intermolecular interactions \[9\]. We will also work with the case that \( \sigma > a \), so no “true” contact between the sheets will occur. In a related study, Oyharcabal and Frisch \[19\] used a van der Waals-like medium-range potential with values of \( p = 3 \) and \( q = 9 \) to model the attraction between a thin filament and a nonpolar substrate. Other examples include the van der Waals interaction between two filaments \( (p = 6, q = 12) \), polarized attraction between two sheets \( (p = 2, q > p) \), Coulombic attraction \( (p = 1, q > p) \), and many others (see Ref. \[9\] for a review). In fact, a surface with an arbitrary charge distribution can be represented by a standard multipole expansion and, in a suitable far-field regime, a charged polymer or conducting elastic sheet can be modeled by this potential as well. Very generally, by specifying the values of \( p \) and \( q \), any number of potential interactions can be represented, except in the rather exceptional cases in which a power-law potential model is insufficient.

### B. Dimensionless equations

The system described by Eqs. (2)–(5) is nondimensionalized by scaling the vertical displacements by \( h \) and horizontal distances by \( L \). In what follows, variables are understood to be dimensionless. In that case, Eqs. (2)–(5) become

\[ y_1^{m}(x_1) + \Pi_{p,1} I(x_1, y; p, 2) - \Pi_{q,1} I(x_1, y; q, 2) = 0, \tag{7} \]

\[ y_2^{m}(x_2) - \Pi_{p,2} I(x_2, y; p, 1) + \Pi_{q,2} I(x_2, y; q, 1) = 0, \tag{8} \]

\[ y_1(0) = 1, \quad y_1'(0) = y_1''(1) = y_1'''(1) = 0, \tag{9} \]

\[ y_2(0) = 0, \quad y_2'(0) = y_2''(1) = y_2'''(1) = 0, \tag{10} \]

where \( \Pi_{p,2} = \rho \sigma^{p} L^{2-p} W / B_1 \) is a dimensionless quantity, and where we have defined the integral \( I(x_1, y; \alpha, k) \) as

\[ I(x_1, y; \alpha, k) = \int_0^1 \frac{y_1(x_1)}{[(x_1 - x_2)^2 + (y_1(x_1) - y_2(x_2)]^{2n+2}}, \tag{11} \]

with \( y = y_1(x_1) - y_2(x_2) \).

### C. Asymptotics

We now take advantage of the long-wavelength approximation \( \epsilon \ll 1 \) to simplify the integrals of the form \( I(x_1, y; \alpha, k) \). Introducing the substitution \( x_1 - x_2 = \epsilon^\alpha \xi \), we obtain

\[ I(x_1, y; \alpha, 2) = \frac{1}{\epsilon^{\alpha+1}} \int_{-\infty}^{\infty} \frac{y_1(x_1) - y_2(x_1 - \epsilon \xi)}{(\xi^2 + [y_1(x_1) - y_2(x_1 - \epsilon \xi)]^{2n+2})} d\xi. \tag{12} \]

Expanding to leading order in \( \epsilon \),

\[ I(x_1, y; \alpha, 2) = \frac{1}{\epsilon^{\alpha+1}} \int_{-\infty}^{\infty} \frac{y_1(x_1) - y_2(x_1)}{(\epsilon^2 + [y_1(x_1) - y_2(x_1)]^{2n+2})} d\xi + o\left(\frac{1}{\epsilon^{\alpha+1}}\right), \tag{13} \]

\[ = \frac{1}{\epsilon^{\alpha+1}} \int_{-\infty}^{\infty} \frac{y_1(x_1) - y_2(x_1)}{(1 + \epsilon^2)^{2n+2}} + o\left(\frac{1}{\epsilon^{\alpha+1}}\right), \tag{14} \]

\[ = \epsilon^{\alpha+1} \int_{-\infty}^{\infty} \frac{y_1(x_1) - y_2(x_1)}{(1 + \epsilon^2)^{2n+2}} + o\left(\frac{1}{\epsilon^{\alpha+1}}\right), \tag{15} \]

where we have used \( \epsilon = \xi [y_1(x_1) - y_2(x_1)] \). The other integrals in Eqs. (7) and (8) are evaluated similarly. Physically, Eq. (15) expresses the fact that in the long-wavelength limit \( \epsilon \ll h \), each sheet sees the other one as being locally flat and therefore, at leading order, the integration along the horizontal direction can be performed first.

### D. Identical sheets

Having derived above the general system of equations for two interacting sheets, we now consider the simplified case where the sheets are identical. Defining \( z(x) \) as the distance between the sheets \( z(x) = y_1(x) - y_2(x) \), Eqs. (7)–(10) become

\[ z''(x) + \Omega \left(\frac{1}{\epsilon^2} - \frac{\beta}{\zeta^2}\right) = 0, \tag{16} \]

\[ z(0) = 1, \quad z'(0) = 0, \tag{17} \]

\[ z''(1) = 0, \quad z''''(1) = 0, \tag{18} \]

where \( \Omega = (2WJ_p L^{2-p})/h \), \( \beta = (\sigma/h)^{p-n} J_q / J_p \), and \( J_a = \Gamma[(1 + \alpha)/2] / \Gamma(1 + \alpha/2) \). Note that the divergent behavior \( z = 0 \) is prohibited thanks to the repulsive part of the potential in Eq. (16). The dimensionless quantity \( \Omega \), which we refer to as the bending parameter, is a measure of the relative importance of the interaction forces to the elastic forces, while \( \beta \) is a dimensionless van der Waals-like radius with a numerical prefactor. Hence, the symmetric system is completely described by the four parameters \( \{\Omega, \beta, p, q\} \).

### E. Numerics

The symmetric nonlinear system described by Eqs. (16)–(18) is solved numerically on an adaptive grid to an
IV. ADHESION TRANSITION

A. Main result

The main result of this paper is illustrated in Fig. 3. For the particular values $p = 3$, $q = 9$, and $\beta = 0.05 J_0 / J_3$, we display the nondimensional bending energy of the sheets $E_B$—i.e., the sum of the first two terms in Eq. (1)—as a function of the bending parameter $\Omega$. Although here we have chosen specific values for $\{p, q, \beta\}$, the results are similar for other values, with some possible qualitative differences highlighted in the sections below.

The sudden shape changes quantified by the bending energy are reminiscent of the behavior observed experimentally in Sec. II. For $\Omega \ll 10^{-2}$, the two sheets are essentially free standing, as indicated in Fig. 3(a). As $\Omega$ is increased, the sheets are attracted weakly to one another, resulting in a small shape change [Fig. 3(b)]. At a critical value of $\Omega$, the bending energy jumps discontinuously and the sheets abruptly snap together [Fig. 3(c)]. As $\Omega$ increases further, the sheets become more tightly bound, with the unclamped portion becoming smaller. If $\Omega$ is then decreased, the system exhibits hysteretic behavior, with the sheets remaining adhered as shown in Fig. 3(d) [dashed lines indicate the previous shape from Fig. 3(c)]. As $\Omega$ is decreased further, still there is another discontinuity in the energy, and the sheets once again take on a qualitatively different shape (arclike), as displayed in Fig. 3(e). Remarkably, there exists a second (smaller) hysteresis loop on this branch of the energy profile. The jump in the energy at this second hysteresis corresponds to a large change in the contact between the sheet end points and the slope of the sheets. Finally, for decreasing $\Omega$, the sheets return to the positions shown by Fig. 3(a) via another sharp transition.

We also plot in Fig. 4 the dimensionless force $y_1'(0)$ and moment $y_2'(0)$, necessary to apply to the left edge of the first sheet to maintain it clamped. For the values of $\{p, q, \beta\}$ considered (6, 12, and $0.15 J_1 / J_6$, respectively), the system models two filaments interacting via van der Waals forces. Much like the shapes themselves, the forces and moments undergo sharp transitions and exhibit hysteresis. Note that if the sheets were free to interact, they would adhere along their entire length, and a force would need to be applied to one end in order to peel them apart. In essence, the same
Note the lack of a second hysteresis loop for these values of the adhesion parameter. In analogy to the external field of the magnetic system, there is an energy penalty associated with deforming the sheets and at the end of the unadhered length. As the bending parameter is increased further, the spatial location of the energy minima shifts, as displayed by Fig. 5(c). The length of the adhered region (or domain wall) $\delta$ increases with $\Omega$ as $\delta \sim \Omega^{-1/4}$, as expected from the boundary layer scaling arising from Eq. (16). This scaling is confirmed by our numerical simulations (not reproduced here). Decreasing $\Omega$ and coming down on the hysteresis loop, at a lower critical value of $\Omega$ the adhered sheets become arclike, and the second local minimum in $\Psi(x)$ disappears [see Fig. 5(d)].

C. General behavior

Although our investigation to this point has considered particular values for the parameters $\{\beta, p, q\}$, qualitative changes in the hysteresis and transition behavior can be obtained for different values of these parameters. Not only can the hysteresis region be made to shrink, but it can also disappear entirely. In addition, while the transitions seen so far have been first-order, by tuning the model parameters this transition can be made to become second-order.

These different behaviors are illustrated in Fig. 6, where for convenience we have introduced the parameter $b = \beta J_p/\Omega$. As we saw above, there are three characteristic shapes for the sheets that we will denote as weakly bent (W), adhered/clamped (C), and arc shaped (A). The areas that exhibit each of these shapes are depicted in Fig. 6. In Figs. 6(a) and 6(b), we display the total energy (bending plus attraction) for a system with $p = 6$ and $q = 12$. These values model the attraction between two thin nonpolar filaments, and the characteristic shapes seen are similar to those in Fig. 3. If $b$ is increased, representing an increase in the minimum adhesion distance between the fibers, the arc shapes and the associated phase transition vanish. Similarly for the case where $p = 3$ and $q = 9$ [19], increasing the value of $b$ causes the hysteresis region to shrink and the arc shape vanishes [Figs. 6(c) and 6(d)]. Furthermore, the hysteresis region can be made to disappear completely for $p = 2$ and $q = 4$ [Figs. 6(e) and 6(f)].

The transformation of a first-order transition into a second-order transition indicates that there may be a cusp catastrophe in the parameter space we are exploring [25]. For
FIG. 5. (Color online) Bending energy density \([y''(x)]^2\) and interaction energy density \(\Psi(x)\) for various sheet shapes (top) along the hysteresis loop for \(p=3, q=9\), and \(\beta=0.05J_0/J_5\). (a) Weak bending \((\Omega=0.5)\), with ends of the sheets approximately straight (zero bending energy); (b) tightly clamped configuration \((\Omega=0.89)\) with the local minimum in the interaction energy density denoting the end of the unclamped region. The local minimum in the interaction energy is in the same vicinity, but not at the same point; (c) hysteretic-clamped shape \((\Omega=0.5)\), with vertical dashed line indicating the position of the local maximum of the bending energy density, and the vertical dotted line indicates where this local maximum was in (b). The red (outer horizontal dashed) line in the upper inset denotes the weakly bent shape from (a), while the blue (inner horizontal dashed) line denotes the shape from (b). (d) Arc shape \((\Omega=0.11)\), with a notable small local maximum in \(\Psi(x)\) near the end point indicating localized adhesion.

The case \(p=3\) and \(q=9\), we plot in Fig. 7 the bending energy landscape as both \(\Omega\) and \(b=\beta J_3/J_0\) are varied. For small values of \(\beta\), the first-order nature of the transition is apparent. For a fixed value of \(\Omega\), increasing \(\beta\) decreases the area of hysteresis until a critical value is reached where the hysteretic behavior vanishes completely.

D. Nonidentical sheets

We have considered so far the case where the two sheets are identical. If we allow instead their bending rigidity to be different, we now have two parameters \(\Omega_i=(2WJ_3L^2\rho^\alpha)/(B_i\rho^\beta)\) for \(i=1,2\). In this case, the difference in rigidity causes a change in both the equilibrium point of adhesion and the critical values of \(\Omega_i\). The different shapes can be characterized by a phase diagram that maps the transition points for different values of the two bending parameters, as displayed in Fig. 8. Three distinct regions exist: multistability of both arc shapes and weak bending (MS), arc shapes exclusively (A), and adhered or clamped states (C). As shown in Fig. 8(a), for small values of \(\Omega_2\) and \(\Omega_1\), weak attraction occurs, unless the hysteretic regime has been entered, in which case there will be adhesion (not shown). As \(\Omega_1\) is increased past the fixed value of \(\Omega_2\), the sheets adhere in an asymmetric arc shape [Fig. 8(b)]. This transition is of first order, and the sheets adhere smoothly into a clamped phase [Fig. 8(c)]. In this final clamped state, the position of the adhered portion of the sheets depends on the relative values of the bending parameters [Fig. 8(d)].

E. Adhesion of three sheets

We now consider the adhesion transition for an array of multiple sheets and illustrate the complexity and richness of the system considered on a few examples. Assuming the nearest-sheet interaction for simplicity, with identical potentials, we can easily extend the modeling approach offered above to the case of \(N\) interacting sheets [26]. The equation
of shape for the $i$th sheet $1 \leq i \leq N$ is then given by

$$y_i^{(m)} + \Omega \sum_j \text{sgn}(i - j) \left[ \frac{1}{|y_i - y_j|^p} - \frac{\beta}{|y_i - y_j|^{p+q}} \right] = 0,$$  \quad (19)

where the sum on $j$ runs over nearest neighbors, and $\Omega$, $p$, $q$, and $\beta$ are defined as in the $N=2$ case.

For multiple sheets, any asymmetry in the system now plays a role in determining the order in which sheets adhere to one another. For three sheets, unless there is perfect symmetry between the top and bottom sheets, the adhesion events always occur in a sequential fashion (see Figs. 9 and 10). Specifically, two sheets first come together and then adhere to the third sheet for a further increase in the relevant bending parameter. By locating the middle beam slightly closer to one of its neighbors, this sequence can be made to occur preferentially between two previously chosen sheets (by changing the clamping distance, the competition between bending and interaction energy changes, and in effect one of the bending parameters gets a boost from the geometric asymmetry). For three identical sheets, as the bending pa-
and a region of multiple stability (MS) where arc shapes and weak bending are both possible, (a) arc shapes only and (c) tightly clamped sheets. (a) In this region, the sheets are either bending weakly or, in the hysteretic case, they exhibit arc shapes; (b) as $\Omega_1$ is increased, the sheets snap together into an arc shape (a first-order transition); (c) as $\Omega_1$ increases further, there is a smooth variation from arc shapes into clamped [i.e., a second-order transition (dashed line)]; and (d) if $\Omega_1$ remains fixed and $\Omega_2$ is decreased, the bottom sheet will become relatively more rigid, producing a net shift in the equilibrium adhesion position [the dashed lines indicate the adhesion shape in (c)].

FIG. 8. (Color online) Phase diagram of sheets with asymmetric bending parameter for $p=3$, $q=9$, and $\beta=0.15J_3/J_2$. There is a region of multiple stability (MS) where arc shapes and weak bending are both possible, (a) arc shapes only and (c) tightly clamped sheets. (a) In this region, the sheets are either bending weakly or, in the hysteretic case, they exhibit arc shapes; (b) as $\Omega_1$ is increased, the sheets snap together into an arc shape (a first-order transition); (c) as $\Omega_1$ increases further, there is a smooth variation from arc shapes into clamped [i.e., a second-order transition (dashed line)]; and (d) if $\Omega_1$ remains fixed and $\Omega_2$ is decreased, the bottom sheet will become relatively more rigid, producing a net shift in the equilibrium adhesion position [the dashed lines indicate the adhesion shape in (c)].

FIG. 9. (Color online) Sequential adhesion transition for three sheets $(1,1,1)\rightarrow(1,2)\rightarrow(3)$. Two consecutive first-order transitions occurring for the total bending energy of the sheets as a function of the (identical) bending parameters in the case $p=3$, $q=9$, and $\beta=0.35J_3/J_2$. An inherent asymmetry has been introduced in that the middle sheet is pinned slightly closer to the bottom sheet than to the top one (5% difference in height). (a) Weak attraction; (b) as the bending parameters are increased, a first-order adhesion transition takes place where two of the three sheets adhere; and (c) a second first-order transition occurs when the three sheets adhere. Both transitions display hysteresis.

FIG. 10. (Color online) Same as Fig. 9, but for $\beta=0.45J_3/J_2$. In contrast with the case depicted in Fig. 9, the first adhesion transition is now of second order.

When a fourth sheet is added, the adhesion transitions can be made to occur in either a hierarchical or a sequential fashion. This is illustrated in Figs. 11 and 12.

We show in Fig. 11 an example of sequential adhesion for four identical sheets, similar to the one discussed in the three-sheet case. The values of $p$, $q$, and $\beta$, as well as the geometric asymmetry, have been chosen so that there is a mix of first-order and second-order transitions (see figure captions). The relative distances between the four sheets are as follows. The top sheet is pinned at $y=3h$, the second highest at $y=1.8h$, the third highest at $y=0.7h$, and the final sheet at $y=0$. As all four bending parameters are increased at the same rate, the sheets start by a state of weak attraction, with more bending exhibited by the sheets that are closer to one another. Past a critical value of the bending parameters, there is a second-order transition for these values of the model parameters, and the two bottom sheets adhere [Fig. 11(b)]. As the bending parameters are increased further, another second-order transition takes place and the lower three sheets adhere [Fig. 11(c)]. Finally, a final first-order transition occurs when the four sheets adhere [Fig. 11(d)]. Note the non-monotonic variation of the bending energy in the second-highest sheet (inset of Fig. 11).

An example of hierarchical adhesion transition is displayed in Fig. 12, where we plot the total bending energy profile for a symmetric four-sheet system (i.e., there is no asymmetry in the relative distances between the sheets). As the bending parameters are increased, a second-order phase...
transition leads to an adhesion between two pairs of sheets [Fig. 12(a)]. First-order transitions are also possible for other values of the model parameters (not shown here). As the bending parameters are increased further, a first-order transition occurs and the four sheets all adhere to one another [Fig. 12(b)]. Remarkably, in this case, the first-order transition does not display any hysteresis.

V. CONCLUSION

In this paper, we have studied the prototypical dry adhesion problem between flexible sheets or filaments and focused on their morphological transitions. Motivated by a simple macroscale experiment showing hysteretic adhesion, we have introduced a model of dry adhesion between two elastic slender sheets, interacting via a power-law potential, and studied numerically the transitions in their conformations. Given a particular form of the interaction potential, the system is completely described by a single dimensionless parameter quantifying the relative effect of the long-range attraction and bending rigidity and governing the nature of the adhesion transitions (first or second order). We have also generalized the model to multiple sheets, showing—in particular—that additional geometric considerations dictate the order in which structures adhere to each other. The physical systems modeled here include the interactions between charged sheets or between nonpolar filaments. Future work will focus on the presence of thermal fluctuations allowing the adhered states to “jump” from one state to another. We will also consider the case where the filaments are actuated and will include the effect of hydrodynamic interactions. Finally, using an approach similar to ours, the adhesion of three-dimensional structures such as coiled filaments or planar arrays could be investigated.

ACKNOWLEDGMENTS

We would like to thank Saverio Spagnolie for many useful conversations and help with some numerical aspects of this work. This work was funded in part by the National Science Foundation (Grants No. CTS-0624830 and No. CBET-0746285) (E.L.).

[26] Even for $p=1$ (long ranged or Coulombic attraction), adding contributions from all neighbors does not qualitatively change our results.