

# How double-stranded DNA breathing enhances its flexibility and instability on short length scales

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We study the unexpected high flexibility of short dsDNA which recently has been reported by a number of experiments. Via the Langevin dynamics simulation of our Breathing DNA model, first we observe the formation of bubbles within the duplex and also forks at the ends, with the size distributions independent of the contour length. We find that these local denaturations at a physiological temperature, despite their rare and transient presence, can lower the persistence length drastically for a short DNA segment in agreement with experiment.

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The DNA is a double helix of two single-stranded (ss) backbone chains paired by complimentary bases via hydrogen bonding and further stabilized by stacking interaction between adjacent base-pair (bp) planes [1]. Owing to recent advances of single molecule techniques, intensive studies have been done on structural changes and mechanical behaviors of double-stranded (ds) DNA constrained by external forces and twists. The continuum elastic model called worm-like chain (WLC) model has been very useful in analytically describing the micron-scale conformations and elastic response of such DNA [2]. The persistence length, the measure of segmental orientation correlation, is about 50 nm (equivalently, about 150 bases along contour) for dsDNA. On the contrary numerous biological facts suggest DNA loops more readily on much shorter length scales. Indeed, Cloutier and Widom showed that the DNA has much higher cyclization probability than predicted by a WLC of persistence length of 50 nm [3]. Also, Wiggins *et al.* showed the DNA on short length scale has an elastic behavior distinct from that of WLC [4], while Yuan *et al.* very recently reported the persistence lengths are as short as 11 nm for DNA fragments consisting of 10–20 bp [5].

In this paper, we demonstrate that the higher flexibility of dsDNA indeed emerges on shorter scales due to local denaturation. Because of the large initiation energy, the fraction of the open bases is less than 1% at the physiological temperature, and, once formed, bubbles decay shortly on the order of 50  $\mu$ s [6], seemingly little affecting the DNA stability. We show that despite their rare and transient presence the bubbles give a drastic enhancement of the flexibility as the chain gets shorter. For a very short duplex fragment, another type of local denaturation, namely, forks at the free ends are entropically favorable, dominating over the bubbles to enhance the flexibility.

It was suggested and estimated that the baseflips [5] and kinks [7] can enhance the bending flexibility and looping probability. Based on a simple two state model, the looping

probability was evaluated using transfer matrix method [7]. Although all these calculations are suggestive of their significance, the bubbles have not yet been explicitly accounted for with regard to their size distributions and realistic energetics.

We consider a homogeneous dsDNA as the duplex of two interacting single strands described by the effective energy

$$\mathcal{H} = \mathcal{H}_1 + \mathcal{H}_2 + \mathcal{V}_{12}. \quad (1)$$

The  $\mathcal{H}_i$  is the elastic energy of the single strand  $i$  ( $=1$  or  $2$ ), which, in a discrete representation, is given by

$$\mathcal{H}_i = \sum_{n=2}^{N-1} \frac{\kappa}{2} (\mathbf{r}_{n-1}^{(i)} - 2\mathbf{r}_n^{(i)} + \mathbf{r}_{n+1}^{(i)})^2 + \sum_{n=1}^{N-1} \frac{k}{2} (|\mathbf{r}_{n+1}^{(i)} - \mathbf{r}_n^{(i)}| - b)^2, \quad (2)$$

where  $\mathbf{r}_n^{(i)}$  is the three-dimensional position vector of  $n$ th bead in strand  $i$  ( $n=1, 2, \dots, N$ ) and  $b$  ( $\cong 0.34$  nm) is an average distance between neighboring beads within the ss. The first term accounts for bending energy with the bending modulus  $\kappa b^3$  for the ss, which is related to its persistence length ( $L_p$ ) via  $\kappa b^3 = L_p k_B T$  [8]. The second term, the stretching energy of each strand, is introduced to impose chain inextensibility condition. An appropriate value of  $k$  is numerically found by matching simulation of either force-extension curves or the mean end-to-end distances of ssDNA resulting from this model, with corresponding theoretical expression of the inextensible WLC without the stretching term. The  $\mathcal{V}_{12}$  is the pairing energy between complimentary bases. To describe the bp openings due to thermal excitation, namely, thermal bubbles and forks, the interaction is represented by  $\mathcal{V}_{12} = \sum_{n=1}^N \mathcal{V}_n(r_n)$  where

$$\mathcal{V}_n(r_n) = D e^{-(r_n - r_0)/a} [e^{-(r_n - r_0)/a} - 2], \quad (3)$$

is the Morse potential [9,10],  $r_n = |\mathbf{r}_n^{(1)} - \mathbf{r}_n^{(2)}|$  is the distance between  $n$ th bp,  $r_0$  and  $a$  are the bond distance and range which correspond to mean and fluctuation of DNA diameter, respectively, and  $D$  is the potential depth. Whenever bps are unbound, the duplex becomes no more than two single strands with net persistence length as short as twice of  $L_{ss} = 1-4$  nm [11], while for bound bps it takes  $L_{ds} = 50$  nm, the persistence length of long DNA, due to stacking interaction in the double strand. In order to include

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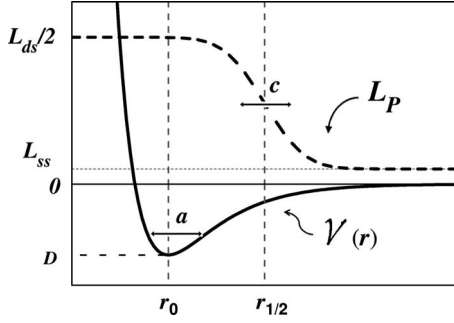


FIG. 1. A schematic figure showing the profile of persistence length  $L_p$  per single strand (dashed curve) along with the Morse potential  $\mathcal{V}(r)$  (bold curve) as function of bp distance  $r$ .

the stacking and destacking that cause such variation in the persistence length in the model, we consider  $\kappa(n) = L_p(n)k_B T / 2b^3$ , and propose the persistence length per single strand  $L_p(n)$  takes the form

$$L_p(n) = L_{ds}/2 - (L_{ds}/2 - L_{ss})\tilde{\theta}(r_{n-1} - r_{1/2}) \times \tilde{\theta}(r_n - r_{1/2})\tilde{\theta}(r_{n+1} - r_{1/2}). \quad (4)$$

Here  $\tilde{\theta}(r) = [1 - \text{erf}(r/c)]/2$  is a steplike form function which smoothly increases from 0 to 1 over the width  $c$  and  $r_{1/2}$  is the distance at which  $L_p$  is  $(L_{ds}/2 + L_{ss})/2$  (see Fig. 1).  $L_p$  depends on three consecutive bp distances naturally as it represents the cooperativity of the stacking interaction [12]. The variation in  $L_p$  depending on the bp distance  $r$  is schematically illustrated in Fig. 1: when all of the three consecutive bps are outside the range  $r_{1/2}$ , namely, they are unbound one other,  $L_p$  takes the single-stranded persistence length  $L_{ss}$  while as any one of the three are bound,  $L_p$  is reduced to  $L_{ds}/2$ . The  $c$  is comparable to the Morse potential width  $a$  since the  $L_p$  varies due to the bp unbinding.

Using this energy model, we simulate the dynamics and equilibrium distribution of the bp distance, via the Langevin equation,

$$\Gamma \frac{d}{dt} \mathbf{r}_n^{(i)}(t) = - \frac{\partial \mathcal{H}}{\partial \mathbf{r}_n^{(i)}} + \boldsymbol{\xi}_n^{(i)}(t), \quad (5)$$

where frictional coefficient per base is  $\Gamma = 6\pi\eta R = 1.88 \times 10^{-11} \text{ kg s}^{-1}$  with  $\eta = 0.001 \text{ kg m}^{-1} \text{ s}^{-1}$  for viscosity of water and  $R = 1 \text{ nm}$  for effective radius of nucleotide.  $\xi_n$  is the Gaussian and white noise satisfying  $\langle \xi_{n\alpha}^{(i)} \rangle = 0$ ,  $\langle \xi_{n\alpha}^{(i)}(t) \xi_{n'\alpha'}^{(j)}(t') \rangle = 2\Gamma k_B T \delta_{ij} \delta_{m'} \delta_{\alpha\alpha'} \delta(t-t')$  for each Cartesian component  $\alpha$  and  $\alpha'$ . The values of potential depth and range,  $D = 0.07 \text{ eV}$ ,  $a = 0.05 \text{ nm}$  are chosen so that the 300 bp DNA fragments have the melting temperature 350 K [8].

In our simulation, we employed 50 nm for the persistence length of dsDNA ( $L_{ds}$ ) and 4 nm for that of ssDNA ( $L_{ss}$ ) with the stretching constant  $k = 41 \text{ pN/nm}$ . For the parameters in  $L_p(n)$ ,  $r_{1/2}$  was chosen to  $(r_0 + r_c)/2 = 2.1 \text{ nm}$  where  $r_c$  is the cutoff distance at which a bp is regarded as unbound in the simulation, and  $c$  is 0.045 nm [8]. We checked that small variation in the parameters values does not affect significantly the main result of our simulation.

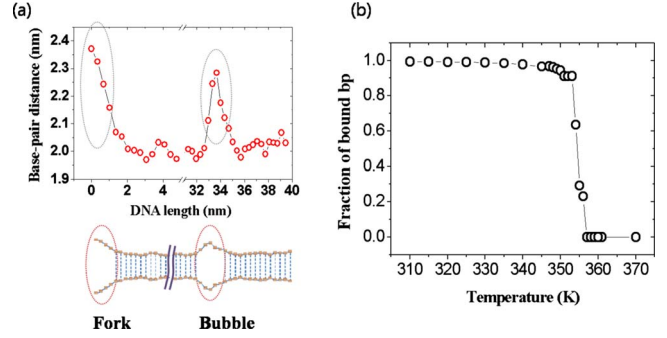


FIG. 2. (Color online) (a) A snapshot of bp distance in a duplex of 300 bp ( $\sim 100 \text{ nm}$ ) long obtained from simulations of the breathing DNA model. The shape of duplex thickness therefrom drawn manifests breathing, formation of bubble in the middle of the duplex and forks at the ends. (b) The simulation data on fraction of bound bp in the 300-bp-long duplex vs temperature. It shows a sharp transition into denaturation state at melting temperature ( $\approx 350 \text{ K}$ ). At physiological temperature ( $\sim 310 \text{ K}$ ), the fraction less than 1% is denatured (unbound) either as bubbles or forks.

Figure 2 shows a snapshot profile of equilibrated bp distance along the contour of 300 bp long or about 100 nm, with free ends, and the fluctuating DNA thickness of about 2 nm therefrom constructed. The figure shows indeed the local denaturation exist in the forms of bubbles within the contour and fork at the ends. From the simulation we find the fraction of bound pairs undergoes a sharp transition from near unity to zero in agreement with experiment [13] at the melting temperature, which is about 350 K. It indicates that at the physiological temperature (310 K), the fraction of the bp that forms the local denaturation is less than 1%.

Shown in the Fig. 3 are the size distributions of the bubbles [ $P_b(n)$ ] and forks [ $P_f(n)$ ] for the various contour lengths of the DNA fragments. The distribution of the bubble size with  $n$  bp open is remarkably independent of its position and the DNA length. For relatively large bubbles it follows the Poland-Sherega form [14]

$$P_b(n) \sim \frac{e^{-n\Delta}}{n^\alpha}, \quad (6)$$

where  $\alpha$  is the statistical factor for a loop formation and  $\Delta$  is the average energy in  $k_B T$  to unbind a bp. These factors are

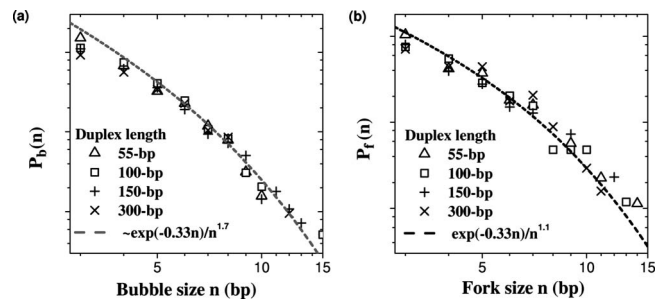


FIG. 3. (a) The bubble size distribution  $P_b(n)$  at 310 K for contour length  $L = 55, 100, 150,$  and  $300 \text{ bp}$ . (b) The fork size distribution  $P_f(n)$  at 310 K for the same lengths.

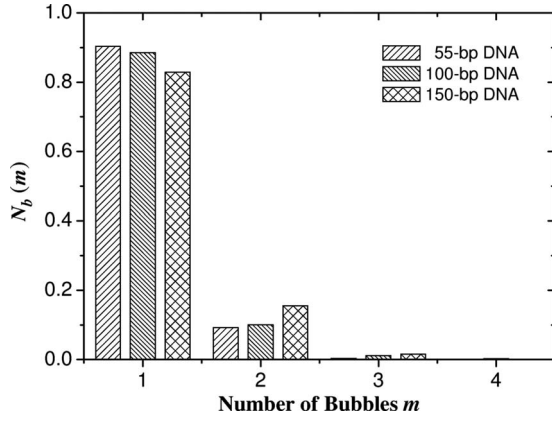


FIG. 4. The relative probability of finding  $m$  bubbles within three DNA lengths at 310 K.

independent of the length and given by those for a long length,  $\alpha=1.7$ ,  $\Delta=0.33$ , at 310 K [8]. The average size of the bubbles for various contour lengths is 1.7 bp at 310 K. This is in agreement with the Peyrard-Bishop-Dauxois model simulation [15]. The length independence of their distribution and small average size implies that the thermal bubbles are transiently excited due to short-range elastic interaction in unconstrained DNA. This is to be contrasted with the dsDNA under mechanical constraints, where the average bubble size is found to be much larger depending on the (negative) twist as well as the contour length [16]. The distribution of  $n$  bp open at a free end is given by

$$P_f(n) \sim \frac{e^{-n\Delta}}{n^\beta}, \quad (7)$$

where  $\beta=1.1$  and  $\Delta=0.33$  are also independent of the contour length within the errors. The average size of the forks is 1.9 bp. Since  $L_{ss}$  is 4 nm in these simulations, they are semi-flexible forks with  $\beta$  much larger than that of the flexible forks [17].

Last, we obtain the  $N_b(m)$ , the relative probability of finding  $m$  bubbles *simultaneously* in a DNA of given length. The result obtained from simulation is presented in Fig. 4, showing that the single bubble occurrence becomes predominant as the DNA length gets shorter. This is mainly due to the large energy cost of bubble initiation; namely, once a bubble is formed, increasing its size is energetically more favorable than opening another bubble elsewhere. Within the short DNAs of our interest, the number of bubbles, if they exist at all, can be regarded to be unity in a good approximation.

To focus on the effects of bubble (excluding the end fork effect) on the overall duplex persistence length, we consider an ensemble of fragments with given contour length  $L$  randomly taken *from inside* a 300 bp long ( $\sim 100$  nm) dsDNA at a physiological temperature. The square of duplex end-to-end distance,  $\mathbf{R}_{ee}^2$ , is defined as arithmetical average of the end-to-end distances of the two single strands. The thermal average of its square  $\langle \mathbf{R}_{ee}^2 \rangle$  is taken over the ensemble of the duplex with a given  $L$ . Suppose that the duplex to be a WLC with a uniform persistence length  $L_p^{eff}$ , we use the well-known relation [18]

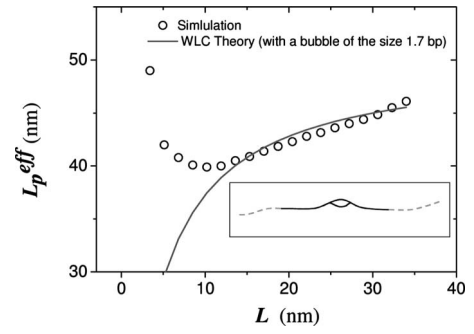


FIG. 5. Effective persistence length of the DNA fragments of the length  $L$  taken from inside a 300-bp-long DNA (empty circles) This is compared with theoretical curve (line) obtained assuming a single bubble of the size given by the mean, 1.7 bp.

$$\langle \mathbf{R}_{ee}^2 \rangle = 2L_p^{eff}L - 2(L_p^{eff})^2[1 - e^{-L/L_p^{eff}}]. \quad (8)$$

Comparing this with simulation data, we evaluate  $L_p^{eff}$  of our duplex for various values of the contour length  $L$ . The result shown in Fig. 5 indicates that  $L_p^{eff}$  decreases from the value 50 to 40 nm as the  $L$  decreases to 10 nm. It reflects the enhanced flexibility of the duplex, which we attribute to the presence of thermal bubbles. Along the DNA length shorter than 50 nm, if any, a single bubble is most likely to exist, as shown in  $N_b(m)$ , Fig. 4. If the single bubble survives as the contour length decreases, its mean size does not change as discussed earlier, consequently yielding larger flexibility.

To support this argument quantitatively, we note that the mean end-to-end distance is given by  $\langle \mathbf{R}_{ee}^2 \rangle = \int_0^L \int_0^L ds ds' \langle \mathbf{t}(s) \cdot \mathbf{t}(s') \rangle$  [18], where  $\mathbf{t}(s)$  is the unit tangent vector at the arclength  $s$  from an end of the duplex. The tangent correlation function is given by  $\langle \mathbf{t}(s) \cdot \mathbf{t}(s') \rangle = \exp\{-|s' - s|/L_{ds}\}$  if they are the positions within  $ds$  region and  $\langle \mathbf{t}(s) \cdot \mathbf{t}(s') \rangle = \exp\{-|s' - s|/2L_{ss}\}$  if  $s$  and  $s'$  are the points within the bubble region. Integrating the correlation function along the contour with a bubble of the mean size, 1.7 bp, we evaluate the  $\langle \mathbf{R}_{ee}^2 \rangle$ . Since a bubble occurs with equal probability along the contour we further average the mean square end-to-end distance over all possible position of the bubble, and relate it with the effective persistence length, which is shown by the curve in Fig. 5.

For the length larger than 10 nm, this analytical theory agrees with simulation result remarkably, suggesting that the persistence length reduction is indeed due to a single bubble. As the length decreases below that, this curve departs much from the simulation result for the length shorter than that. It is because even a single bubble is unlikely to occur within such a short length so that the persistence length sharply rises to the double strand value. For a long DNA fragment, on the other hand, only a bubble is found in most cases with its average size fixed independently of DNA length, which results in no significant perturbations on the persistence length.

For the short duplex with free ends, we also have investigated the mean end-to-end distances of DNA fragments length shorter than 50 nm (147 bp) at 310 K. The persistence length calculated in a similar way from these data is shown

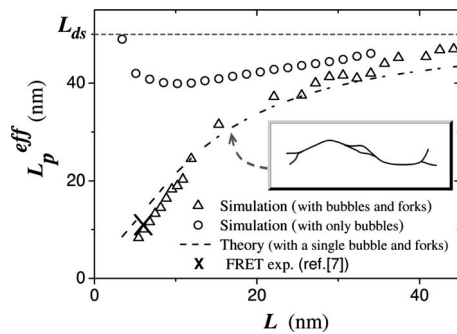


FIG. 6. Effective persistence length of the fragment of DNA with contour lengths  $L$ , where bubbles as well as forks exist (triangle). It is in a good agreement with the WLC theoretical curve with a bubble of the size given as its average, 1.7 bp and two forks of the size given as its average, 1.9 bp, and with the results of Ref. [5] obtained by a FRET experiment.

by triangles in Fig. 6. It clearly shows that  $L_p^{eff}$ , in this range of the contour length, are shorter than  $L_{ds}=50$  nm but converges to the value as the contour length increases. The persistence length in this case is shorter than that of the duplex of the same contour length without the free ends discussed before (indicated by circles). This can be ascribed to the additional form of local denaturation, i.e., the forks at the free ends. To support this quantitatively, we analytically calculated the effective persistence length of the duplex with a single bubble of the size 1.7 bp and two forks of the size 1.9

bp at the ends by integrating the correlation function and following the procedure as before. The close agreement of the theoretical curve with the simulation evidences additional influence of the forks; the two forks combine with and increasingly dominates over a bubble, to enhance flexibility, as the duplex gets shorter.

In Ref. [5], the effective persistence length of dsDNAs of contour length 15~21 bp (equivalently 5~7 nm) was determined to be  $11 \pm 2$  nm by a fluorescence resonance energy transfer (FRET) experiment, which is marked by a cross in Fig. 6. They considered the buffer conditions where the persistence lengths of noninteracting single-stranded DNA have the range of the values 2.7~3 nm [11]. In a duplex, however, the two unpaired single strands are subject to steric and electrostatic repulsions, which may enhance this ss persistence length effectively to about 4 nm. Indeed, our values of  $L_p^{eff}$  estimated from simulation with  $L_{ss}(=4$  nm) are in a good agreement with this experimental value.

Simulating the Breathing DNA model that incorporates locally fluctuating persistence lengths depending upon the bp distances along the contour, we find that the distributions of bubbles and forks are nearly independent of the contour length, reducing the effective persistence length of the duplex for the short lengths. The forks dominate the bubbles in enhancing duplex flexibility.

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