

Analytical description of finite size effects for RNA secondary structures

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The ensemble of RNA secondary structures of uniform sequences is studied analytically. We calculate the partition function for very long sequences and discuss how the cross-over length, beyond which asymptotic scaling laws apply, depends on thermodynamic parameters. For realistic choices of parameters this length can be much longer than natural RNA molecules. This has to be taken into account when applying asymptotic theory to interpret experiments or numerical results.

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I. INTRODUCTION

Folding of biopolymers is a fundamental process in molecular biology without which life as we know it would not be possible. In biopolymer folding, well-characterized interactions between individual monomers make a polymer fold into a specific structure believed to minimize the total interaction free energy. The apparent simplicity in the formulation of this biopolymer folding problem is in sharp contrast with the immense challenges faced in actually describing biopolymer folding quantitatively caused by the intricate interplay of monomer-monomer interactions and the constraint that the monomers are connected into a chain of a certain sequence. The biological importance of biopolymer folding paired with this immense intellectual challenge has sparked numerous computational and theoretical studies [1]. These studies do not only attempt quantitative predictions of specific structures but also focus on more fundamental properties of the biopolymer folding problem such as its phase diagram.

While the bulk of the work concentrates on the folding of proteins due to its overwhelming importance in pharmaceutical applications, recently RNA folding has been identified as an ideal model system for biopolymer folding [2, 3]. RNA is a biopolymer of four different bases G, C, A, and U. The most important interaction among these bases is the formation of Watson-Crick(WC) base pairs, i.e., A-U and G-C pairs. This comparatively simple interaction scheme makes the RNA folding problem very amenable to theoretical approaches without losing the overall flavor of the general biopolymer folding problem. Again, a lot of effort has been devoted to understanding fundamental properties of RNA folding such as the different thermodynamic phases an ensemble of RNA molecules can be in as a function of temperature, an external force acting on the molecules, and the sequence design [4–9].

All these theoretical approaches are concerned with the phase behavior of RNA molecules in the *thermodynamic limit*. In order to compare these theoretical predictions with numerical or actual biological experiments it is thus important to know which role *finite size effects* play, i.e., at which size of a molecule the universal predictions of the asymptotic theories are expected to hold. In this publica-

tion we precisely aim to answer this question. We study homogeneous RNA sequences which allows us to *analytically* solve for the universal asymptotic behavior as well as the cross-over length below which the universal theory is not applicable any more. We find that this cross-over length is very strongly dependent on the sequence of the molecule. For realistic energy parameters we find that the cross-over length can be as large as 10,000 bases. This is about the largest size of naturally occurring RNAs as well as the largest length of RNA molecules amenable to quantitative computational approaches. Thus, we conclude that finite size effects have to be seriously taken into account when describing RNA folding by asymptotic theories.

This article is organized as follows: In Sec. II, we briefly review the definition of RNA secondary structure. In Sec. III, we analytically derive the finite size effects of the simplest model of RNA folding namely a homogeneous sequence without loop entropies. While this model is mainly treated for pedagogical purposes, in Sec. IV, we sketch how the result can be generalized to more realistic models of RNA folding. In Sec. V, the behavior of the cross-over length N_0 is discussed. We find that N_0 depends mostly on the degree of branching of the RNA molecules and a simple approximate formula is derived. These results are shown to be consistent with the numerical values obtained using experimentally known energy parameters for specific sequences in Sec. VI. We point out how enormous finite size effects in the RNA secondary structure formation problem can be. The detailed derivations of the partition function and the cross-over length are relegated to various appendices.

II. REVIEW OF RNA SECONDARY STRUCTURES

A. Definitions

RNA usually occurs as a single-stranded polymer with four types of monomers (bases) G, C, A, and U. The strand can bend back onto itself and form helices consisting of stacks of stable Watson-Crick pairs (A with U or G with C).

An RNA secondary structure describes which bases are

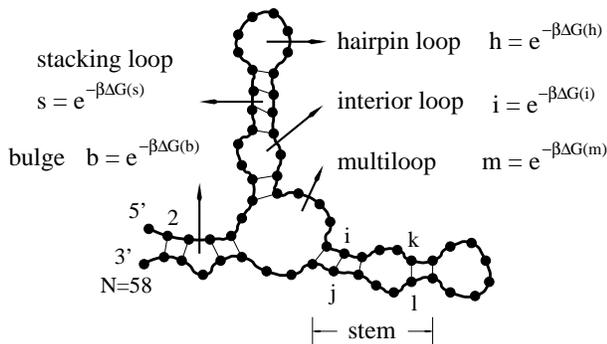


FIG. 1: An RNA secondary structure. The thick line stands for the backbone of the molecule and thin lines stand for base pairings. The solid dots represent monomers. 5' and 3' show the head and tail of this RNA of length 58. Many different loops formed when RNA folds are also defined in the figure.

bound and can be written as a set of binding pairs (i, j) , where i and j denote the i^{th} and j^{th} base of the RNA polymer respectively. For example, the secondary structure \mathbf{S} of Fig. 1 is written as

$$S = \{(2, 57), (3, 56), \dots, (i, j), \dots, (k, l), (37, 44)\}.$$

In this study, we apply the common approximation to exclude the so-called pseudo-knots [2], i.e., for two base pairs (i, j) and (k, l) , the configurations $i < k < j < l$ and $k < i < l < j$ are not allowed. As a result, the analytical studies become more tractable.

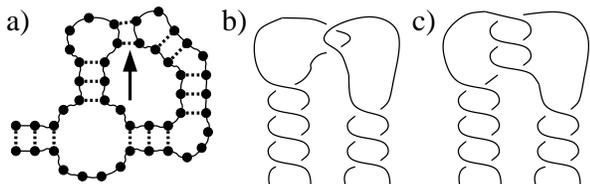


FIG. 2: Pseudo-knots in RNA structures: The base pairings indicated by the arrow in a) create a pseudo-knot. b) the short pseudo-knots (called “kissing hairpins”). c) the long pseudo-knots in 3 dimensions.

This exclusion of pseudo-knots is reasonable. For long pseudo-knots, the double helix structure would require threading one end of the molecule through its loops many times so they are kinetically difficult to form. Thus, these pseudo-knots occur infrequently in natural RNA structures [2, 3]. Short pseudo-knots, on the other hand, do not contribute much to the total free energy because of the relatively low binding energies for short sequences and the strong electrostatic repulsion of the backbone since the polymer backbone is highly charged. By excluding pseudo-knots, we will stay close to commonly used algorithms that compute the exact partition function which can be applied to test our model.

B. Interaction energies

Since the tertiary interactions between structures are in general much weaker than the interactions among the secondary structures [2, 3], we will follow the common approaches and take into account only the energy contribution from the secondary structures.

If we assign a Gibbs free energy $\Delta G(\mathbf{S})$ to each secondary structure \mathbf{S} , the partition function of the ensemble of all structures is given by

$$Z = \sum_{\mathbf{S}} e^{-\Delta G(\mathbf{S})}. \quad (1)$$

The Gibbs free energy is commonly used to describe the secondary structure since it contains entropic contributions from the formation of loops as well as enthalpic terms from the formation of base pairs. The total free energy $G(\mathbf{S})$ is the sum of the energy contributed from each elementary piece such as the stacking of base pairs and the connecting loops. The largest contribution are the stacking energies between adjacent WC pairs, and these energies depend on the type of bases in the pairs. While the typical value of the stacking energy is on the order of $k_B T$ at room temperature, both the enthalpic and entropic terms are on the order of $10k_B T$. Thus, the stacking energy will become repulsive with a moderate increase of temperature to around $80^\circ C$ and the RNA molecule denatures.

III. MOLTEN PHASE

In order to get a qualitative understanding of finite size effects, we first follow previous works [3, 5, 6, 8, 10] and assume that the Gibbs free energy is the sum of the binding free energies ϵ_{ij} of each base pair in the structure,

$$\Delta G(\mathbf{S}) = \sum_{(i,j) \in \mathbf{S}} \epsilon_{ij}, \quad (2)$$

and neglect the entropic energies due to the formation of loops for the rest of this section.

The binding free energies ϵ_{ij} in this model are differences between the gain in chemical binding energy and the loss in the configurational entropy associated with the formation of the base pairs. Since both contributions are large and comparable, the realistic values of the ϵ_{ij} strongly depend on temperature. Since we do not describe spatial degrees of freedom in this model, it does not describe denaturation of the RNA molecule and we restrict ourselves from here on to a parameter regime where the majority of the bases is paired, i.e., where a significant fraction of the ϵ_{ij} is negative.

In order to obtain analytical insights into the finite size effects, we additionally assume that the binding free energy ϵ_{ij} is a constant ϵ_0 , independent of the identities of the bases. Thus, in our simplified model $\Delta G(\mathbf{S}) = \epsilon_0 \times |\mathbf{S}|$ where $|\mathbf{S}|$ stands for the number of pairs in \mathbf{S} .

$$G(N) = G(N-1) + \sum_{k=1}^{N-1} G(k) \cdot G(N-k)$$

FIG. 3: Recursive relation for a simple model of an RNA molecule. The wavy lines stand for undetermined structure and the corresponding partition function. The arch represents a binding pair between bases k and N . The assumption of excluding pseudo-knots separates the last term into two independent parts since two pairs can not go across each other.

This simplified energy model serves as the basis of our study for the more realistic energy model.

As it stands, this energy model and the more realistic energy model we will introduce later describe only homogeneous sequences. However, it has been argued that this energy model can be applied to random RNA sequences at high enough temperature when the disorder is sufficiently weak [10]. Under this weak disorder, there exist many structures with nearly degenerate energies and the corresponding scaling laws match the predictions of the simplified energy model. Only as the temperature is lowered, a strong disorder phase arises. This low temperature phase is characterized by a small number of distinct low-energy structures, and is referred to as the “glass phase” in analogy with studies of other disorder system. However, this glass phase is not within the scope of this article.

The partition function of the molten phase model can be obtained through the recursive relation in Fig. 3. This figure shows how the possible ways of binding can be decomposed into two cases where the last base N is either unbound or bound to some base k . If we define $G(N+1)$ as the partition function of an RNA of length N , the relation reads

$$G(N+1) = G(N) + \sum_{k=1}^{N-1} G(k) \cdot q \cdot G(N-k), \quad (3)$$

where $q = e^{-\beta\epsilon_0}$. Together with the boundary condition $G(1) = 1$, this equation allows calculation of the exact value of the partition function $G(N)$ in $O(N^2)$ time. The Vienna package [11] is able to calculate this exact value with more complete sequence dependent energy parameters based on the similar scheme. This recursive equation (3) also leads us to the analytical expression for the partition function. By introducing the z -transform,

$$\widehat{G}(z) = \sum_{N=1}^{\infty} G(N) \cdot z^{-N}, \quad (4)$$

and applying it to Eq. (3), we get a quadratic equation for $\widehat{G}(z)$ as

$$z\widehat{G}(z) - 1 = \widehat{G}(z) + q\widehat{G}^2(z), \quad (5)$$

from which $\widehat{G}(z)$ can be solved. For large sequence lengths, the partition function $G(N)$ is obtained by per-

forming the inverse z -transform on $\widehat{G}(z)$, and can be approximated (see Appendix A) as

$$G(N) = \frac{1}{2\pi i} \oint \widehat{G}(z) z^{N-1} dz. \quad (6)$$

$$\approx A(q) N^{-\theta} z_c^N(q) \left[1 - \frac{N_0(q)}{N} + O\left(\frac{1}{N^2}\right) \right], \quad (7)$$

where $z_c(q) = 1 + 2\sqrt{q}$ is the branch point of $\widehat{G}(z)$, $\theta = 3/2$, $A(q) = [(1 + 2\sqrt{q})/4\pi q^{3/2}]^{1/2}$ and $N_0(q) = 3(1 + 4\sqrt{q})/16\sqrt{q}$. This asymptotic analytical formula is only determined by the behavior of $\widehat{G}(z)$ near the branch point z_c . The exponent $\theta = 3/2$ indicates the characteristic universal behavior of this partition function for long sequences. The non-universal cross-over length $N_0(q)$ characterizes how long a sequence has to be for the universal laws to hold. Here, we find an explicit analytical formula for N_0 as a function of parameter q .

From the formula of $N_0(q)$, we can see that the cross-over length in this simple model is on the order of 1 for all values of q . However, in the following sections we will show that the cross-over length may vary over several order of magnitudes from several bases to 1,000,000 bases. Thus, the loop entropies of the more realistic energy model would greatly modify the behavior of cross-over length.

IV. INCLUDING LOOP ENTROPIES

To get a more quantitative understanding of the cross-over length, we now take into account the loop entropies and introduce Boltzmann factors, s , b , i , h , and m , for the different types of loops (see Fig. 1). The values of these free energy parameters have been carefully measured [15] such that our model can be applied quantitatively to realistic RNA molecules. Typically, the free energy of a stacking loop is large and negative ($s \gg 1$) while the free energies for all the other loops tend to be large and positive, leading to Boltzmann factors much less than one. The binding energy ϵ_0 of the simple model introduced above is now absorbed into these loop free energies. As mentioned in Sec. III, we still restrict ourselves to a temperature regime below denaturation which is determined by the true energy model.

Again, we want to calculate the partition function of the structure ensemble and derive the cross-over length as a function of the loop parameters. This calculation in principle follows along the lines of Sec. III, but is technically much more elaborate because of the more complicated energy model. A reader more interested in the final results than in the technical details is advised to directly skip to the Sec. V.

In order to calculate the partition function, we separate the secondary structure into two categories as shown in Fig. 4. One is the bubble structure which contains only hairpins and multiloops. The other is the stem structure, which connects the bubbles, containing only stack-

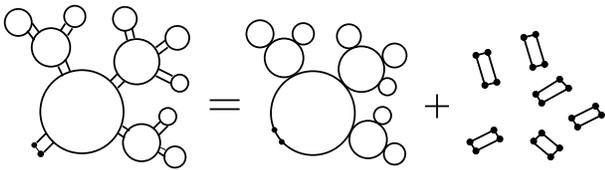


FIG. 4: Separation of stems from the bubble structure.

ing loops, bulges, and interior loops. We will study each of them individually and later combine them together.

A. Stem structure

In principle, the loop free energy depends on the length of a loop. Thus, unbound bases also contribute to the total free energy. This contribution has been experimentally measured for small loops and behaves logarithmically with length when the loop is large. However, in the following we show that the free base energy of unbound bases provides only a negligible effect on the behavior of a stem and thus on the cross-over length if the stacking energy is relatively large.

To explore all possible ways of pair bindings, again a graphical recursion relation is helpful. Such a recursion relation is shown in Fig. 5. Starting from a closed pair on the left, the following loop can be either a stacking loop, a bulge or an interior loop which correspond to the terms on the right hand side. To study the influence of a free energy for unbound bases, we assign the Boltzmann factors, b and \tilde{i} , to each unbound base in a bulge and an interior loop respectively. If we define the partition function of stem structures with N bases and the first and the last bases of which is paired as $S(N-1)$, which corresponds to the left hand term, this relation in Fig. 5 is formulated as

$$S(N-1) = sS(N-3) + 2 \sum_{k=3}^{N-2} b\tilde{b}^{N-k-1} S(k-2) + \sum_{a=3}^{N-3} \sum_{b=a+1}^{N-2} \tilde{i}^{N-(b-a)-3} S(b-a). \quad (8)$$

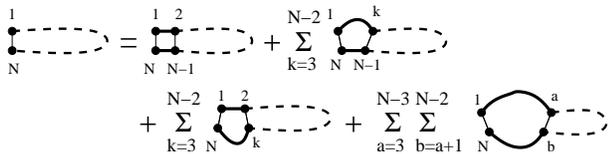


FIG. 5: Recursion relation for stem structure. The dashed lines stand for the undetermined structures. Thick lines represent the backbone and thin lines stand for pair bindings.

To perform the z-transform, we have to consider the initial conditions, $S(1) = 1$, $S(2) = 0$, $S(3) = s$, and $S(4) = 2b\tilde{b}$. These initial conditions arise because certain

structures are not allowed, e.g., since a base can not be shared in two base pairings, a stem with 3 bases does not exist and this leads to $S(2) = 0$. Also when the length of a stem is small, certain loops which require many bases are not allowed, e.g., the only available structure for a stem with 4 bases is a stacking loop. Including these conditions, we apply the z-transform on Eq. (8) with the definition

$$\widehat{S}(z) = \sum_{N=1}^{\infty} S(N)z^{-N}, \quad (9)$$

to resolve the convolution. Solving for $\widehat{S}(z)$ gives us

$$\widehat{S}(z) = \frac{1}{z} \left[1 - \frac{s}{z^2} - \frac{2b \cdot \tilde{b}}{z^2(z - \tilde{b})} - \frac{i \cdot \tilde{i}^2}{z^2(z - \tilde{i})^2} \right]^{-1}. \quad (10)$$

Again the partition function $S(N)$ can be obtained by applying the inverse z-transform on $\widehat{S}(z)$.

To illustrate the effect from the unbound bases, we show how their Boltzmann factors affect the average behavior of the long stem structure. The average quantity of a certain type of loop or unbound base can be calculated as $\tau \cdot \partial_{\tau} \ln S(N)$ where τ is the corresponding Boltzmann factor. Here, we specifically calculate the average number of unbound bases per interior loop, which is defined by $\tilde{i} \cdot \partial_{\tilde{i}} \ln S(N) / i \cdot \partial_i \ln S(N)$ in the large N limit, as a function of \tilde{i} (see Appendix B for detailed calculations).

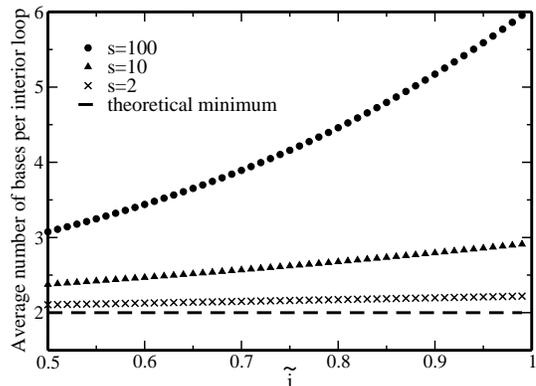


FIG. 6: Average number of unbound bases per interior loop v.s. \tilde{i} for different stacking energy. Following the measured free energies [12], we chose the typical values $b = 0.01$, $\tilde{b} = 0.85$, and $i = 0.05$ for the other Boltzmann factors.

From Fig. 6, we can see that this average number is barely affected by \tilde{i} when the Boltzmann factor s for a stacking loops is large. This can be easily understood as follows: For an interior loop, the unbound bases always introduce a strong energy penalty since less bases are available for stacking. This penalty is much larger than

Notice that the leading singularity in \widehat{G} is again from the branch cut induced by the square root. Thus, as expected, the inverse z -transform leads to the same universal behavior (7) with an exponent $\theta = 3/2$ as the simple model we studied first. However, non-universal quantities such as the cross-over length N_0 will depend on the parameters of the extended model.

D. Minimum hairpin length constraint

For natural RNA molecules, a hairpin loop needs to have at least 3 unbound bases due to the width of the double helix (which implies $j - i > 3$ in the secondary structure). This minimum hairpin length constraint is easily taken into account in our calculations. Under the constraint, a bubble structure which contains at least one hairpin must have at least 5 bases. Thus, we adopt the initial condition, $B(1) = B(2) = B(3) = 0$, when we perform the z -transform on Eq. (11). The summation range in Eq. (11) is then changed and it simply leads to a substitution of h by h/z^3 in all subsequent equations.

This substitution is reasonable since z represents the Boltzmann factor for the free energy of one single base. The minimum hairpin length constraint reduces the number of available bases for binding by 3, so it introduces an energy penalty $3 \cdot \ln z$ which causes the Boltzmann factor h to be divided by z^3 . In this way, we can easily introduce any kind of minimum length constraint via a similar substitution. For example, if we require a bulge loop to have at least two unbound bases instead of one, the replacement of b by b/z will include this constraint.

Note, this principle also helps us to understand equation (10) for the stem structure. The terms with different powers of z , namely s/z^2 , b/z^3 , and i/z^4 , arise because a stacking loop reduces the number of available bases for pairing by 2, a bulge loop has at least one unbound base, and an interior loop has a minimum of 2 unbound bases, respectively.

V. BEHAVIOR OF THE CROSS-OVER LENGTH

In this section, we will use the general results of Sec. IV in order to calculate the cross-over length in our model for sequences with loop entropies.

A. Large stacking energy approximation

To derive the cross-over length, we need to solve for the branch point z_c that is defined by the vanishing of the term under the square root in Eq. (17). In principle, we can always obtain the numerical value of the branch point and expand $\widehat{G}(z)$ around this point to obtain a numerical value for the cross-over length (see Appendix A). We will refer to this numerical value as the homogeneous cross-

over length since it is obtained under the homogeneous molten phase model.

However, since this calculation involves finding the root of a fourth order polynomial, no meaningful analytical expression can be found in general. Thus, we resort to a large s approximation in order to obtain an analytical expression. This approximation is justified since the Boltzmann factor of a stacking loop, s , is usually much larger than 1 while the loop Boltzmann factors b , i , h , and m are less than one. In this approximation, from Eq. (17) we find the branch point $z_c \approx \sqrt{s}$, i.e., the free energy per base is $f = -kT \cdot \ln(z_c) \approx \frac{1}{2} \Delta G(s)$. This can be easily interpreted since we expect most bases to form pairs due to the favorable stacking loops such that the free energy per base is half of the free energy of a stacking loop.

B. Cross-over length

Including the minimum hairpin length constraint introduced in Sec. IV.D, we expand the branch point z_c near \sqrt{s} . Then, the approximated analytical formula for the cross-over length becomes

$$N_0 = \frac{3s^{3/4}}{8\sqrt{hm}} \left[(\sqrt{s} - 1)^2 + h + \frac{9b}{2\sqrt{s}} + \frac{11i - 6b}{4s} + O(s^{-3/2}) \right]. \quad (18)$$

It has a straight forward interpretation: The simplest possible structure is a long stem with one hairpin at the end. Every additional branching of the structure requires formation of one hairpin and one multiloop. Since upon formation of the hairpin and multiloop, at least 3 bases become unbound, the Boltzmann factor for a branching is $hm/s^{3/2}$. The prefactor in Eq. (18) is up to the numerical factor of $3/8$ the inverse square root of this expression. Thus, we conclude that the cross-over length which becomes larger as the Boltzmann factor for a single branching becomes smaller can be interpreted as the minimum length that allows a certain degree of branchings.

Since the Boltzmann factors b and i appear only in the higher order terms, they barely modify the cross-over length and can be neglected altogether. This is consistent with the fact that b and i only play roles in the stem structure, but not in the bubble structure. The leading term of the approximated analytical formula (18) will be referred to as the large s cross-over length.

C. Reliability of the large s approximation

For the analytical large s cross-over length to be useful, we have to know how good they agree with the numerical value of the homogeneous cross-over length. Here, we compare these two values for many different choices of energy parameters covering the whole range of realistic

values. Fig. 9 shows how the large s cross-over length approaches the homogeneous one as s gets large. Typical values for the Boltzmann factor of a stacking loop s involving GC pairs are $s \geq 30$ [15], so the approximation is very good in this region. For stacking loops involving AU pairs, s is around 5 so a deviation from the approximated formula in the large s limit can be seen. However, since N_0 only sets the order of magnitude of the length beyond which the asymptotic theory is applicable, the large s cross-over length with a deviation by a factor of 2 at $s = 5$ is still a good estimation.

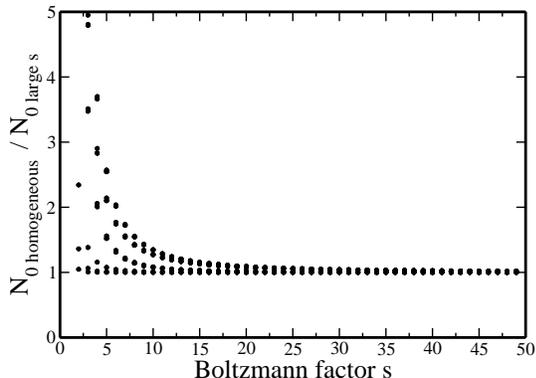


FIG. 9: Ratio between the large s cross-over length and the numerical value of the homogeneous one for many combinations of the parameters: $b, i = \{0, 1\}$ and $h, m = \{0.1, 0.01, 0.001\}$. These choices cover the region of realistic values.

VI. NUMERICAL VERIFICATION

While for a generic RNA sequence, the molten phase model is believed to only apply at sufficiently high temperature, it can be applied to repeated sequences at all temperatures below denaturation since each repeated unit can be viewed as the equivalent of a base in the molten phase (Fig. 10). To illustrate the correctness of the calculations shown in Secs. IV and V, and to get a feeling for typical cross-over lengths, we now compare our large s cross-over length of repeated sequences with the full numerical results. The full numerical result is obtained by using the Vienna package [11] which can calculate the exact value of the partition function and other observables for arbitrary sequences using a realistic energy model.

As an observable, we choose the average size l of a structure. This quantity is defined as

$$l = \sum_{k=1}^{N/2} \sum_{k'=N/2+1}^N P_{k,k'}, \quad (19)$$

where $P_{k,k'}$ is the probability that bases k and k' are paired. The latter probability is also calculated exactly

by the Vienna package. This size measures the average number of base pairs to be crossed when connecting the $N/2^{\text{th}}$ base to base 1 (Fig. 10), which captures the size of the secondary structure. We expect l to obey

$$l \propto N^{1/2} \cdot \left(1 - \left(\frac{N_0}{N} \right)^{1/2} + O\left(\frac{1}{N} \right) \right), \quad (20)$$

where the leading term is the asymptotic behavior [10] and the next term reflects the first expected correction which is a constant independent of N . We determine l for sequences of different lengths and extract the full numerical cross-over length N_0 by fitting data obtained via the Vienna package to Eq. (20). This is then compared to our large s cross-over length.

A. $(GCA)_n$ sequence

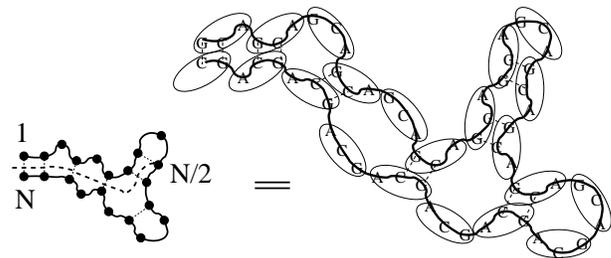


FIG. 10: Equivalence between the $(GCA)_n$ sequence and the molten phase: Three consequent bases GCA are mapped to one single base so the GC/CG stack is equivalent to a binding pair in the molten phase. Then, the smallest interior loop on the left is viewed as a stacking loop and the following interior loop becomes a bulge in the molten phase. The hairpin loop on the right with 2 units of GCA is considered to have two unbound bases in the molten phase.

We apply this scheme to a repeated $(GCA)_n$ sequence. Such sequences naturally occur in the gene for Huntington's disease and their secondary structures are believed to play a role in this disease. Since the GC/CG stack in the secondary structure of the $(GCA)_n$ sequence is much more favorable than any other combination, we can exclude the possibility to have binding pairs other than GC/CG. Thus, GCA is viewed as one unit base in the molten phase. With this equivalence we can use the experimentally determined parameters [12] and calculate the equivalent energy parameters s , b , i , h , and m for the molten phase model. For example, the stacking energy of the molten phase is the sum of GC/CG stacking energy and the free energy for the interior loop of length 2 in the $(GCA)_n$ sequence.

Fig. 11 shows the full numerical results for the average size of the secondary structure for a $(GCA)_n$ sequence as a function of the sequence length N . By fitting the

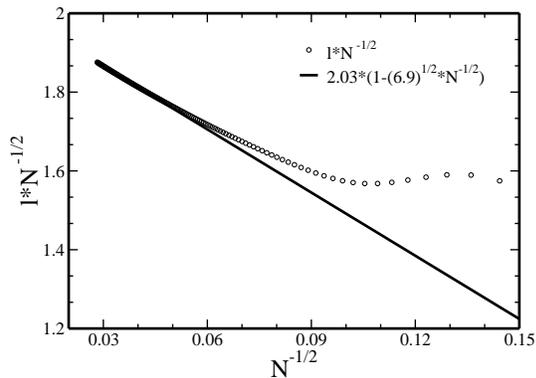


FIG. 11: Average size of the secondary structure of a $(GCA)_n$ sequence at $37^\circ C$. The data is fitted by the expected law (20).

result to the Eq. (22), we get a full numerical value of the cross-over length of 6.9 bases.

To compare this full numerical value with our large s cross-over length, we plug the equivalent energy parameters of the molten phase model into the approximated formula $3s^{1/4}(\sqrt{s}-1)^2/8\sqrt{hm}$. This formula is different from equation (18) because the minimum hairpin length of the corresponding molten phase model only 1 instead of 3. The resulting large s cross-over length is about 2.3 repeat units which corresponds to 6.9 bases in the $(GCA)_n$ sequence. This agrees very well with the full numerical value.

B. $(AU)_n$ sequences

Repeated AU sequences have already been suggested as models for the molten phase by de Gennes in 1968 [9]. For such sequences, we exclude the possibility of AA or UU binding pairs since they are not favorable at all. In a similar fashion as for the $(GCA)_n$ sequence in Fig.10, the smallest bulge has two free bases. However, since the minimum hairpin length is 4 instead of 3, in order to match AU or UA closing pairs, the large s cross-over length is $3s(\sqrt{s}-1)^2/8\sqrt{hm}$.

Plugging in the correct values for the parameters at body temperature results in a large s cross-over length of $N_0 \approx 7700$. A verification of this value is beyond the reach of the numerical procedure using the Vienna package. Since the cross-over length is expected to decrease as the denaturation transition is reached, the full numerical verification could be performed at a higher temperature. Thus, we first study the the temperature dependence of the cross-over length N_0 . To this end we choose to study the homogeneous cross-over length instead of large s cross-over length because it is not clear whether the large s approximation is appropriate when approaching the denaturation temperature. When studying the temperature dependence, it is important to note that the free energies themselves have a very strong temperature dependence which changes Boltzmann factors much more

than the explicit β in the exponent. By looking up the table of enthalpies corresponding to different loops, we determine this temperature dependence of the Boltzmann factors s , b , i , h , and m . The numerical values of the homogeneous cross-over lengths thus obtained are shown in Fig. 12. From this figure, we choose to numerically verify the estimate of the cross-over length at $57^\circ C$.

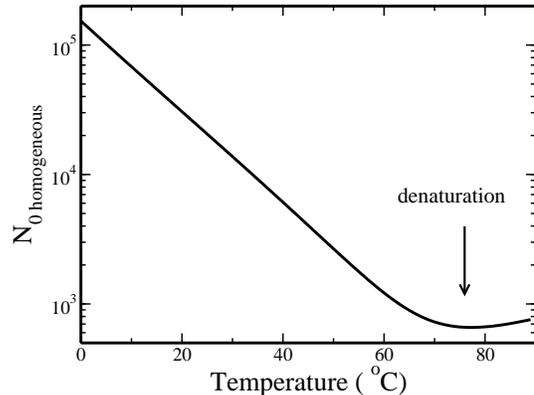


FIG. 12: Numerical values of homogeneous cross-over length for $(AU)_n$ sequence with respect to temperatures. This quantity is obtained by numerically calculating the branch point and then expanding $\hat{G}(z)$ around this point(see Appendix A).

The full numerical verification is done in a similar way as Sec. VI.A. Fitting numerical data at $57^\circ C$ gives a full cross-over length $N_0 \approx 900$. Our homogeneous cross-over length predicts a value $N_0 \approx 1500$ which is of the same order of magnitude as the full numerical result. From the above numerical studies, we conclude that our theory for the large s cross-over length is a good estimate. Thus, the cross-over length of $(AU)_n$ at body temperature is truly around 7,700 bases, which is almost the largest size of naturally occurred RNA sequences. For $(GC)_n$ sequences at body temperature, the situation is even more dramatic with a predicted cross-over length about 105,000 bases. This is beyond the limit of most natural RNAs and suggests that the structure ensemble of $(AU)_n$ and $(GC)_n$ molecules can never be described by the asymptotic theory for any naturally available molecules.

C. Distribution of cross-over length for short repeated sequences

Lastly, we want to explore the sequence dependence of the cross-over length for short repeated sequences. We study all possible partially self-complementary repeated sequences of unit length two and three and evaluate their cross-over lengths using the formula for the large s cross-over length.

For non self-complementary repeated sequences, e.g., the $(GCC)_n$ sequence, it is not a priori clear that we can

Together with the following approximation valid in the large N limit,

$$\int_{\mu_0}^{\mu_c} (\mu_c - \mu)^\alpha e^{\mu N} d\mu \approx \Gamma(1 + \alpha) N^{-(1+\alpha)} e^{\mu_c N},$$

the inverse z-transform of $\widehat{G}(z)$ can be expressed as

$$\begin{aligned} G(N) &= \frac{1}{\pi} \int_{\mu_0}^{\mu_c} \sqrt{f(\mu)} e^{\mu N} d\mu \\ &= \frac{\sqrt{-f'(\mu_c)}}{\pi} \Gamma\left(\frac{3}{2}\right) N^{-3/2} e^{\mu_c N} \\ &\quad \cdot \left[1 - \frac{f''(\mu_c)}{4f'(\mu_c)} \cdot \frac{\Gamma(5/2)}{\Gamma(3/2)} \cdot \frac{1}{N} + O\left(\frac{1}{N^2}\right) \right] \\ &= A_0 N^{-3/2} z_c^N \left[1 - \frac{N_0}{N} + O\left(\frac{1}{N^2}\right) \right]. \end{aligned}$$

Here, the partition function of the molten phase model (7) is obtained by setting $f(\mu) = 4q - (e^\mu - 1)^2$. For the general case, we can put the result in terms of z with the substitutions

$$\begin{aligned} \left. \frac{df(\mu)}{d\mu} \right|_{\mu_c} &= z_c \cdot \left. \frac{df(z)}{dz} \right|_{z_c} \\ \left. \frac{d^2 f(\mu)}{d\mu^2} \right|_{\mu_c} &= z_c \cdot \left. \frac{df(z)}{dz} \right|_{z_c} + z_c^2 \cdot \left. \frac{d^2 f(z)}{dz^2} \right|_{z_c}. \end{aligned}$$

After little algebra, the cross-over length is obtained as

$$N_0 = \frac{3}{8} \cdot \left[1 + \frac{z_c \cdot \left. \frac{d^2 f(z)}{dz^2} \right|_{z_c}}{\left. \frac{df(z)}{dz} \right|_{z_c}} \right]$$

for arbitrary $f(z)$.

B. average length of interior loops

The calculation of partition function for the stem structure $S(N)$ is different from that for the molten phase

model because there is no branch cut, so we do not expect the universal behavior $N^{-3/2}$. However, the fact that only the pole with the greatest real part is important in the large N limit is preserved. From Eq. (8), the z-transform of the partition function for the stem structure $\widehat{S}(z)$ has the form $f(z)/g(z)$ where

$$\begin{aligned} g(z) &= z^2(z - \tilde{b})(z - \tilde{i})^2 - s(z - \tilde{b})(z - \tilde{i})^2 \\ &\quad - 2\tilde{b}\tilde{i}(z - \tilde{i})^2 - \tilde{i}^2(z - \tilde{b}). \end{aligned}$$

In the large N limit, the inverse z-transform can be approximated by

$$\begin{aligned} S(N) &= \frac{1}{2\pi i} \oint \widehat{S}(z) z^{N-1} dz \\ &\approx \frac{1}{2\pi i} \oint \frac{f(z)}{g'(z_c)(z - z_c)} z^{N-1} dz \\ &\approx \frac{f(z_c)}{z_c g'(z_c)} z_c^N, \end{aligned}$$

where $z_c = z_c(s, b, \tilde{b}, i, \tilde{i})$ is the pole with greatest real part.

Since z_c^N is the dominating term for large N, the average quantity can be approximated as

$$\begin{aligned} \langle \tau \rangle &= \tau \partial_\tau \ln(S(N)) \\ &\approx N \tau \partial_\tau \ln(z_c) = N z_c^{-1} \tau \partial_\tau z_c. \end{aligned}$$

So the average number of unbound bases per interior loop can be obtained by $(\tilde{i} \partial_{\tilde{i}} z_c) / (i \partial_i z_c)$. Here, the pole z_c can only be solved numerically.

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