Influence of two bulge loops on the stability of RNA duplexes

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ABSTRACT

Fifty-three RNA duplexes containing two single nucleotide bulge loops were optically melted in 1 M NaCl in order to determine the thermodynamic parameters ΔH° , ΔS° , ΔG_{37}° , and T_M for each duplex. Because of the large number of possible combinations and lack of sequence effects observed previously, we limited our initial investigation to adenosine bulges, the most common naturally occurring bulge. For example, the following duplexes were investigated: 5'GGCAXYAGGC/3'CCG YX CCG, 5'GGCAXY GCC/3'CCG YXACGG, and 5'GGC XYAGCC/3'CCGAYX CCG. The identity of XY (where XY are Watson–Crick base pairs) and the total number of base pairs in the terminal and central stems were varied. As observed for duplexes with a single bulge loop, the effect of the two bulge loops on duplex stability is primarily influenced by non-nearest neighbor interactions. In particular, the stability of the stems influences the destabilization of the duplex by the inserted bulge loops. The model proposed to predict the influence of multiple bulge loops on duplex stability suggests that the destabilization of each bulge is related to the stability of the adjacent stems. A database of RNA secondary structures was examined to determine the naturally occurring abundance of duplexes containing multiple bulge loops. Of the 2000 examples found in the database, over 65% of the two bulge loops occur within 3 base pairs of each other. A database of RNA three-dimensional structures was examined to determine the structure of duplexes containing two single nucleotide bulge loops. The structures of the bulge loops are described.

Keywords: bulge loops; secondary structure; thermodynamics

INTRODUCTION

RNAs are involved in a wide variety of biological activities. RNAs can function as catalysts, either alone or in conjunction with protein (Altman 1990; Cech 1990; Pace and Brown 1995; Scott 2007). Ribosomal RNA has a direct involvement in protein biosynthesis (Noller et al. 1992; Nissen et al. 2000). MicroRNAs are intermediates in the silencing of RNAs, used to target and destroy viral and other RNAs (Plasterk 2002); and siRNAs (small interfering RNA) are involved in the destruction of their RNA targets (Ahlquist 2002). More than a dozen classes of riboswitches, RNA sequences that respond to specific small molecule ligands ranging from amino acids to coenzymes, are known currently (Winkler et al. 2004; Roth and Breaker 2009). A new class of RNA has emerged recently: lncRNA (long noncoding RNA) has been shown to be involved in chromatin remodeling and subsequent epigenetic effects (Guttman et al. 2009; Rinn and Chang 2012). The realization that RNA molecules play an active role in processes of gene expression and are not merely passive structural components has sparked interest in determining the structure of these RNA molecules. In addition, many diseases are mediated through RNA intermediates. Retroviruses, such as the

AIDS virus, are RNA–protein complexes. Antisense therapy relies on the binding of synthetic oligonucleotides to RNA targets (Uhlman and Peyman 1990; Chan et al. 2006). Understanding the structure of RNAs will allow rational design for drug therapy.

The structure and stability of fully base-paired duplex regions of RNA have been extensively investigated (Turner et al. 1988; Xia et al. 2001; Chen et al. 2012). In order to develop the nearest-neighbor model, the thermodynamics of various single-stranded RNA motifs (helices, bulges, internal loops, hairpins, dangling ends, multibranch loops, etc.) are studied in short RNA oligonucleotides. Based partially on these thermodynamic parameters, an algorithm was developed to predict secondary structure from sequence, and it currently predicts 73% of known base pairs (Mathews et al. 1999). The accuracy of the prediction is based on the measured thermodynamics of the various motifs. However, to fully utilize the rapid accumulation of nucleic acid sequence information, further refinements in structure prediction are necessary (Mathews et al. 1999; Andronescu et al. 2007; Lu et al. 2009).

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A bulge occurs when a duplex is interrupted by one or more unpaired nucleotides in one of the strands. Such motifs have been determined to be important in many biological processes including protein binding (Peattie et al. 1981; Wu and Uhlenbeck 1987; Harper and Logsdon 1991; Lilley 1995), intron splicing (Schmelzer and Schweyen 1986; Parker et al. 1987), feedback regulation (Climie and Friesen 1987), and tertiary folding (Woese and Gutell 1989).

Although their biological significance has been studied and they are one of the most common motifs in RNA secondary structure (Michel and Dujon 1983; Woese and Gutell 1989), relatively few studies have been done on the thermodynamics of single nucleotide bulges in RNA (Fink and Crothers 1972; Groebe and Uhlenbeck 1988; Longfellow et al. 1990; Znosko et al. 2002; Blose et al. 2007; McCann et al. 2011; Lim et al. 2012). This has resulted in a lack of thermodynamic characterization of single bulges, and no studies have looked at the influence of two bulge loops in an RNA duplex. This study examines the influence of two bulge loops on the stability of RNA duplex formation and develops a model to predict the free energy contribution for the insertion of two single nucleotide bulge loops into an RNA duplex. These results can be incorporated in secondary structure prediction programs to improve the prediction of RNA secondary structure.

RESULTS

In the database of RNA secondary structures, 20,292 single nucleotide bulge loops were found. Of these, >10% (2250) are located within a duplex containing a second bulge loop. The definition of a duplex used here is two bulges separated by a stretch of canonical base pairs (including wobble pairs) and terminated by a least 1 base pair (bp) on each end of the helix beyond the bulges. These bulge loops were characterized by both the relative orientation (BB, two bulge loops on the same side of the duplex; Bb, bulge loops on opposite strands with the second bulge 3' of the first bulge; bB, bulge loops on opposite strands with the second bulge 5' of the first bulge) (see Table 2 for example secondary structures) and position of the two bulge loops relative to each other. The results of this search are summarized in Table 1. Several represent highly conserved motifs, including position 926 (*Escherichia coli* numbering; helix 28) of the small ribosomal subunit RNA: bB separated by 3 bp. Many of the bulges were separated by only a few base pairs. In fact, there are very few single nucleotide bulges separated by more than 9 bp. This is partially a result of the fact that the average duplex in rRNA is only 7 nucleotides (nt) long.

To determine the influence of two bulge loops on duplex formation, we measured the thermodynamic parameters for duplex formation for the oligonucleotides listed in Table 2. We examined only Group I adenosine bulges because adenosine is the most prevalent bulged nucleotide (Gutell et al. 2000) and Group I bulges eliminate any positional ambiguity (Blose et al. 2007). Fifty-three oligonucleo-

TABLE 1. Number of base pairs separating two single-nucleotide bulge loops in RNA secondary structures

Number of base pairs between the two bulge loops	BB	Bb	bB
1	28	19	167
2	113	22	39
3	111	51	779
4	56	36	32
5	278	12	28
6	97	11	50
7	145	5	81
8	53	12	11
9	0	2	3
10	0	0	1
11	1	0	1
12	0	3	3
13	0	0	0

The RNA Strand database of RNA secondary structures (Andronescu et al. 2008) containing 4666 structures of which 2898 contained bulge loops was analyzed. The database was searched for single nucleotide bulge loops. A total of 20,292 bulge loops was identified of which 2250 were located in duplexes with two single nucleotide bulge loops.

tides were examined (18, BB; 15, Bb; and 20, bB). The oligonucleotides are listed based upon the orientation of the bulge loops relative to each other and then based upon the number of base pairs in the stem regions. Residues in bold are the proposed bulged nucleotides. Structural studies of the RNAs are beyond the scope of this study; however, in choosing sequences to be investigated, we did screen for alternate structural possibilities. For the non-self-complementary sequences, the individual strands were melted and showed no discernable transition.

The parameters are the average values derived from fits of the melt curves and from T_M^{-1} versus $\log(C_t/4)$ (non-self-complementary) or (C_t) (self-complementary) plots. The parameters from the two methods agree within 15% for most duplexes in Table 2, suggesting that the two-state model is a reasonable approximation for these transitions (Freier et al. 1986; Allawi and SantaLucia 1997).

The two exceptions are (CGGCACG)₂ and (CGAGCCG)₂, which are relatively low melting duplexes with difficult to fit lower baselines. Although the enthalpy values derived from the melt data for these two oligomers differ by >15%, the free energy values are similar (−6.1 and −5.8; −6.2 and −6.0) for (CGGCACG)₂ and (CGAGCCG)₂, respectively. These free energy values give reasonable values for the two oligomers so we have chosen to include them in our analysis. The average deviations in thermodynamic parameter values are 6.5%, 7.4%, and 1.8% for ΔH° , ΔS° , and ΔG_{37}° , respectively.

The free energy contribution of each bulged nucleotide was calculated from the experimental data according to Equation 3 and presented in Table 3. As observed with a single bulge loop, insertion of multiple bulge loops into a duplex

TABLE 2. Thermodynamic parameters for duplex formation of multiple single nucleotide bulges^a

Oligomers ^c	T_M^{-1} vs $\log C_T$ plots				Average of curve fits			
	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)
BB								
GAGUUCAG C CAAG C	59.1	174.5	4.9	29.1	58.7	173.3	4.9	29.0
GAGCCGAG C CGGC C	68.8	191.1	9.5	51.0	66.7	184.4	9.5	51.4
CACCGGCAC G GGCCG G	54.5	139.6	11.2	66.0	57.8	149.0	11.6	67.0
GCAUGAUG CG AC AC	41.6	117.7	5.1	26.8	36.7	101.3	5.2	26.4
GUAGGAGC CA CC CG	36.3	98.7	5.7	30.1	31.2	81.2	6.0	33.8
GCAGUUCACG CG CAAG GC	47.9	134.2	6.3	35.5	51.7	146.6	6.3	35.4
GCAGCCACG CG CGG GC	41.3	110.7	7.0	40.6	40.8	108.9	7.0	40.9
GCAGCCGACG CG CGGC GC	61.2	167.0	9.4	52.2	66.9	184.7	9.6	51.9
GCAUGCGUACG CG ACGCA GC	32.4	78.1	8.0	50.8	29.6	69.3	8.1	54.6
GGCAGAGCG CCG C CGC	41.2	108.0	7.7	45.9	35.1	88.4	7.7	47.7
GGCACAGCC CCG G CGG	45.3	120.0	8.1	47.9	48.5	130.4	8.1	47.1
GGCAUUAGGC CCG AA CCG	53.6	151.6	6.6	37.1	61.1	177.0	6.3	36.0
GGCAGCAGGC CCG CG CCG	50.6	136.5	8.3	48.0	45.9	121.5	8.3	49.2
GGCAUUAUAGGC CCG AUA CCG	54.2	152.4	7.0	39.4	53.7	150.6	7.0	39.7
GGCAUUUAGGC CCG AAA CCG	41.6	111.1	7.1	41.4	41.1	109.0	7.3	42.7
GGCAGGCAGGC CCG CCG CCG	66.7	179.0	11.2	60.1	61.0	161.6	10.9	60.9
GGCAUUUUAGGC CCG AAAA CCG	35.5	92.8	6.7	38.5	37.4	98.3	6.9	40.1
GGCAUAAUAGGC CCG AUUA CCG	61.5	174.5	7.4	41.4	62.6	178.1	7.3	40.9
Bb								
GAGUAC C C CAUGAG	35.6	96.6	5.6	36.4	39.9	111.6	5.4	34.6
GAGGAACC G C CCUUGGAC	60.4	161.5	10.3	57.7	60.6	162.1	10.3	57.8
CAGUCCAC G G CAGGUGAC	70.7	189.1	12.0	63.2	65.1	172.2	11.7	63.8
CAGUGCAC G ^d G CACGUGAC	64.8	174.0	10.8	63.6	65.6	176.2	10.9	64.0

Continued

TABLE 2. Continued

Oligomers ^c	T_M^{-1} vs log C_T plots				Average of curve fits			
	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)
CAGGUACC G ^d G CCAUGGAC	67.0	179.8	11.3	65.2	67.8	181.8	11.4	65.4
CGAGC CG GC CGAGC	17.8	37.8	6.1	44.4	28.1	71.9	5.8	38.9
CGAUGA GC GC ACUACG	53.3	152.7	5.9	33.5	47.3	132.8	6.0	33.8
CGAGUAC CG GC CAUGAGC	23.5	61.0	6.1	43.6	26.7	61.7	6.0	41.0
GCAGGCC GC CG CCGGACG	59.3	158.3	10.2	62.5	50.4	131.0	9.8	64.6
GCAUGCGCA GC CG ACGCGUACG	51.9	135.9	9.8	63.4	53.6	141.1	9.8	63.1
GUGAGC C AC CAC CGAGUG	42.3	116.8	6.1	40.2	48.4	136.6	6.1	39.5
GGCAG GCC CCG GCACGG	40.4	108.6	6.7	45.3	37.2	98.5	6.7	45.8
GUCAUCA CUG CAG AGUAGAC	49.3	136.9	6.8	38.7	56.2	159.6	6.7	37.7
GGCAGCGC GCC CCG CGCGACGG	76.8	203.6	13.7	73.2	70.6	185.3	13.2	73.8
CGCCAGCAC AGCC GCGG CGUGAUCGG	88.9	236.1	14.7	68.7	82.3	219.4	14.2	68.9
bb								
G CAUGAC CAGUAC G	36.9	106.5	3.9	22.4	35.5	101.5	4.1	23.9
G GUACAC CACAUG G	38.1	106.0	5.2	33.3	38.1	105.8	5.2	33.4
C GUCGACAG GACAGCUG C	72.5	199.7	10.6	59.6	63.0	170.3	10.1	60.6
C GGUACCAG GACCAUGG C	62.7	169.7	10.1	60.4	58.1	155.8	9.8	60.9
C GGUACCAG GACCAUGG C	52.2	137.6	9.5	61.4	51.8	135.8	9.7	63.2
C GUGCACAG GACACGUG C	63.7	169.7	11.1	65.7	69.7	187.4	11.6	65.7
CG GCACG GCACG GC	23.0	54.3	6.2	43.9	36.5	98.5	6.0	39.3
GC ACUACG CGAUGA GC	52.3	152.3	5.0	28.4	50.4	145.6	5.2	29.1
CG GUACACG GCACAUG GC	26.3	66.3	5.8	38.2	27.9	70.9	5.9	39.3
GC GGCCAGC CGACCGG CG	54.7	142.2	10.6	67.4	53.3	139.3	10.2	65.3
GC AGGUGAGC CGAUCCAC CG	56.2	152.1	9.0	51.3	51.9	138.6	8.9	52.0
GC GUGCACAGC CGACACGUG CG	64.9	174.7	10.7	63.1	65.7	176.7	10.9	63.8

Continued

TABLE 2. Continued

Oligomers ^c	T_M^{-1} vs $\log C_T$ plots				Average of curve fits			
	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)	$-\Delta H^\circ$ (kcal/mol)	$-\Delta S^\circ$ (eu)	$-\Delta G_{37}^\circ$ (kcal/mol)	T_M^b (°C)
CAG GACUG GUCAC GAC	26.1	61.8	7.0	42.1	22.0	48.2	7.0	44.3
GGC CGAGCC CCGAGC CGG	60.7	165.4	9.4	57.2	54.5	146.4	9.1	57.8
GUG GCACAC CACACG GUG	52.8	140.0	9.4	60.3	52.1	137.7	9.4	60.9
GUC ACUACUG CAGAUGA GAC	39.0	104.2	6.6	37.8	41.0	111.1	6.6	37.5
GGC GCGCAGCC CCGACGCG CGG	79.3	208.2	14.7	76.8	77.1	202.1	14.5	76.8
CGC CAGCACAGCC GCGAGUCGUG CGG	86.2	229.9	15.0	70.6	85.1	226.4	14.8	70.6
GCC GUGCUGAGCG CGGACACGAC CGC	85.6	227.6	15.1	71.3	85.0	225.5	15.1	71.6
CGC CACCACAGCC GCGAGUGGUG CGG	74.0	191.4	14.6	75.2	74.1	191.6	14.7	75.2

^aSolutions are 1.0 M NaCl, 10–20 mM sodium cacodylate, 0.5 mM EDTA pH 7.^bCalculated at 10^{-4} M oligomer concentration.^cNucleotides in bold are bulge residues. Top sequence is written 5' → 3' and bottom strand is written 3' → 5'.^dLim et al. (2012).

leads to a destabilization of the duplex. The extent of the destabilization varies widely from 0.8 to 11.4 kcal/mol. In general we observe a smaller influence for the bulges in smaller duplexes. This is somewhat expected as we have previously shown that the influence of single nucleotide bulge loops is dependent upon non-nearest neighbor interactions (Kent et al. 2014). The relationship to predict the influence of single bulge loops on duplex formation is given in the equation below:

$$\Delta G_{37(\text{bulge})}^\circ = -0.51\Delta G_{37(\text{stem})}^\circ + 0.85, \quad (1)$$

where $\Delta G_{37(\text{stem})}^\circ$ is the less stable stem for group I bulge loops and the second least stable stem for group II and group III bulge loops (Kent et al. 2014). Therefore, smaller, less stable stems minimize the influence of bulge insertion on the thermodynamics of duplex formation.

The current model used by RNAstructure and Mfold to predict the free energy contribution of two single nucleotide bulge loops in a duplex is just the sum of the contributions of the individual bulge contributions. To determine how the presence of a second bulge influences the stability of a duplex, several potential models were examined. The model that gave the best fit of the data is described below.

Since the current model to predict the influence of a single nucleotide bulge (Kent et al. 2014) is dependent upon the stability of the adjacent stems (Equation 1), we determined the stability of each of the three stems gener-

ated by the insertion of the two bulge loops into the RNA duplexes listed in Table 2. We then used Equation 1 to predict the influence of two independent bulge insertions. In testing the model, we determined that using the stability of the stem with the second least stability generates a model that predicts the stability of the insertion of two bulges into a duplex with reasonable accuracy (see Discussion).

DISCUSSION

Phylogenetic analysis of two single nucleotide bulge loops

A search of the database of naturally occurring RNA structures found that in cases where duplexes contained two bulge loops, the bulges were often in close proximity to each other. In fact, over half of the structures identified in Table 1 are three or fewer base pairs apart. Since bulge loops were found to be located near one another, we were interested in determining how one bulge loop would influence the thermodynamics for insertion of a second bulge loop near the first.

Possible alternate secondary structures

The oligomers listed in Table 2 are shown as having two single nucleotide bulge loops. However, alternate secondary

structures for some of these duplexes are possible. The two most problematic types of sequences are as follows: (i) structures with a single base pair at the end of the duplex; (ii) two structures with a single base pair separating the two bulge loops. The first structures could potentially form a duplex with terminal mismatches and a single nucleotide overhang. The influence of terminal mismatches and single nucleotide overhangs has been measured and can be used to predict the stability of these duplexes as though they have formed these secondary structures (Serra and Turner 1995). The average difference between these predicted free energy values and the measured values for the 13 oligomers with this type of sequence is 0.3 kcal/mol (average deviation 0.7 kcal/mol). The average difference, 0.2 kcal/mol (average deviation 0.9) between the measured and the predicted values for these duplexes where the influence of two bulge loops on duplex stability is predicted by Equation 2 is smaller but not statistically different. In addition, the inclusion of these oligomers does not alter the overall model for the influence of two bulge loops on duplex stability (see below). Therefore, we have chosen to include these duplexes in our analysis. The second set of problematic structures has the two bulges separated by a single base pair. If the base pair did not form, the structure would represent a 3×1 internal loop. Three duplexes in Table 2 have this potential secondary structure. Assuming that these duplexes form a 3×1 internal loop, the stability of the duplexes can be predicted (Schroeder and Turner 2000). The difference between the measured and these predictions are on average 1.6 kcal/mol; however, these duplexes are also not predicted well as duplexes with two bulge loops, the average difference being almost 2.5 kcal/mol. Therefore, we excluded them from further analysis (Equations 1, 2 and Tables 3, 4).

Thermodynamic contribution of two single nucleotide bulge loops to duplex formation

As previously observed for the insertion of a single nucleotide bulge loop into a duplex, the insertion of two bulge loops destabilized all of the duplexes in this study. The extent of the destabilization is generally larger than observed for the insertion of a single bulge loop; and, perhaps not surprisingly, the range of destabilization is also much wider (Blose et al. 2007). Since the identity of the bulges was constant (two adenosine residues), the magnitude and range of destabilization caused by the bulges must be due to non-nearest neighbor influences, as was observed for the insertion of a single bulge loop into a duplex (Blose et al. 2007). The best predictor of the influence of the two bulge loops on duplex stability was deter-

TABLE 3. Thermodynamic parameters for insertion of two single nucleotide group I bulge loops in RNA duplexes

Base pairs outside bulge	Base pairs between bulge	BB ΔG_{37}° (kcal/mol)	Bb ΔG_{37}° (kcal/mol)	bB ΔG_{37}° (kcal/mol)
1	4	2.2, 2.0	2.2	2.6, 3.7
1	5	3.5		
1	6		3.3, 1.5, 1.2, 0.8	1.1, 2.4, 3.0, 1.2
2	2	2.8, 4.5	4.0	3.9
2	3	7.0, 6.5	5.5	5.5
2	4	7.1	6.4, 6.8	6.4, 6.7
2	5	8.2		8.8
2	6		10.1	8.5
3	1			
3	2	7.8, 9.1	8.0, 11.0	8.3, 4.7
3	3	9.5, 8.2, 8.9	6.7	6.9
3	4	9.6, 9.4	9.1	8.1
3	5			
3	6			11.0, 7.9, 11.4
4	4		11.2	

Measurements were made in 1.0 M NaCl, 10–20 mM sodium cacodylate, and 0.5 mM Na₂EDTA, pH 7. Values calculated as described in text.

mined to be related to the stability of the adjacent stem. Since the two bulges separate the duplex into three stem regions, we examined a number of possible models to provide a predictive model for the stability of duplexes containing two single nucleotide bulge loops. The model that worked the best uses Equation 1 with the $\Delta G_{37}^{\circ}(\text{stem})$ being the second least stable stem of the three stems in the duplex.

For example, for the sequence 5' GCAGGUCACG/3' CG CCAG GC, the three stems are GC/CG, GGUC/CCAG, and CG/GC with predicted stabilities of -3.42 , -5.52 , and -2.36 kcal/mol, respectively. Using the second least stable stem (-3.42 kcal/mol) and Equation 1, the predicted influence of a single bulge loop would be 2.6 kcal/mol. Since there are two bulge loops in the duplex, the influence of both would be predicted to be 5.2 kcal/mol. The measured value for the insertion of the two bulge loops is 6.53 kcal/mol.

Using this model, the influence of two bulge loops on duplex formation is ~ 1 kcal/mol (0.7 kcal/mol) more destabilizing than the predicted influence of the two individual bulge loops (Table 4). Figure 1 displays a graph of the measured influence of the two bulge loops versus the predicted value, assuming there is no interaction between the two bulge loops. The slope of the line is ~ 1 (1.04) ($R^2 = 0.81$), suggesting that there is a strong correlation between the measured values and the predicted values for the insertion of two single nucleotide bulge loops within the same duplex. The fact that the measured values for the duplexes with two bulge loops are on average 0.72 kcal/mol less stable than predicted (value of Y intercept) suggests that there is an interaction between the two bulge loops, which leads to a further destabilization of the duplex relative to the predicted values for the insertion of two bulge loops. Interestingly, we examined the influence of the distance between the bulges (as measured by the predicted influence of the two bulges) versus the difference between the measured and predicted value for the insertion of the bulges and found no correlation between these values (data not shown). This indicates that the observed interaction

TABLE 4. Comparison of models to predict the influence of bulge loops on a duplex^a

Oligomer	ΔG_{37}° (bulge)		(kcal/mol)		
	Measured	Predicted ^b Eq. 1	Difference	Predicted ^c Eq. 2	Difference
BB					
GAGUUCAG C CAAG C	2.15	1.7	0.45	2.4	−0.25
GAGCCGAG C CGGC C	1.97	1.7	0.27	2.4	−0.43
CACCCGGCAC G GGCCG G	3.53	1.7	1.83	2.4	1.13
GCAUGAUG CG AC AC	2.77	3.4	−0.63	4.1	−1.33
GUAGGAGC CA CC CG	4.50	5.0	−0.50	5.7	−1.20
GCAGCCACG CG CGG GC	6.99	5.2	1.79	5.9	1.09
GCAGUUCACG CG CAAG GC	6.53	5.2	1.33	5.9	0.63
GCAGCCGACG CG CGGC GC	7.11	5.2	1.91	5.9	1.21
GCAUGCGUACG CG ACGCA GC	8.25	5.2	3.05	5.9	2.35
GGCAUUAGGC CCG AA CCG	7.79	8.5	−0.71	9.2	−1.41
GGCAGCAGGC CCG CG CCG	9.11	8.5	0.61	9.2	−0.09
GGCAUAUAAGGC CCG AUA CCG	8.89	8.5	0.39	9.2	−0.31
GGCAUUUAGGC CCG AAA CCG	8.22	8.5	−0.28	9.2	−0.98
GGCAGGCAGGC CCG CCG CCG	9.47	8.5	0.97	9.2	0.27
GGCAUUUUAGGC CCG AAAA CCG	9.55	8.5	1.05	9.2	0.35
GGCAUAAUAGGC CCG AUUA CCG	9.42	8.5	0.92	9.2	0.22
Bb					
GAGUAC C C CAUGAG	2.21	1.7	0.51	2.4	−0.19
GAGGAACC G C CCUCCAC	3.27	1.7	1.57	2.4	0.87
CAGUCCAC G G CAGGUGAC	0.83	1.7	−0.87	2.4	−1.57
CAGUGCAC G G CACGUGAC	1.52	1.7	−0.18	2.4	−0.88
CAGGUACC G G CCAUGGAC	1.23	1.7	−0.47	2.4	−1.17
CGAGC CG GC CGAGC	4.04	4.1	−0.06	4.8	−0.76

Continued

TABLE 4. *Continued*

Oligomer	ΔG°_{37} (bulge)		(kcal/mol)		
	Measured	Predicted ^b Eq. 1	Difference	Predicted ^c Eq. 2	Difference
CGAUGA GC GC ACUACG	5.47	5.2	0.27	5.9	−0.43
CGAGUAC CG GC CAUGAGC	6.43	4.1	2.33	4.8	1.63
GCAGGCC GC CG CCGGACG	6.78	5.2	1.58	5.9	0.88
GUGAGC CAC CAC CGAGUG	8.02	6.1	1.92	6.8	1.22
GGCAGC GCC CCG GCACGG	11.02	8.5	2.52	9.2	1.82
GUCAUCA CUG CAG AGUAGAC	6.67	6.0	0.67	6.7	−0.03
GGCAGCGC GCC CCG CGCGACGG	9.06	8.5	0.56	9.2	−0.14
CGCCAGCAC AGCC GCGG CGUGAUCGG	11.25	10.2	1.05	10.9	0.35
bB					
G GUACAC CAC AUG G	2.61	1.7	0.91	2.4	0.21
G CAUGAC CAGUAC G	3.74	1.7	2.04	2.4	1.34
C GUCGACAG GACAGCUG C	1.14	1.7	−0.56	2.4	1.26
C GGUACCAG GACCAUGG C	2.43	1.7	0.73	2.4	0.03
C GGUACCAG GACCAUGG C	3.03	1.7	1.33	2.4	0.63
C GUGCACAG GACACGUG C	1.22	1.7	−0.48	2.4	−1.18
CG GCACG GCACG GC	3.94	4.1	−0.16	4.8	−0.86
GC ACUACG CGAUGA GC	5.47	5.2	0.27	5.9	−0.43
GC GGCCAGC CGACCGG CG	6.78	5.2	1.58	5.9	0.88
CG GUACACG GCACAUG GC	6.73	5.2	1.53	5.9	0.83
GC GGCCAGC CGACCGG CG	6.38	5.2	1.18	5.9	0.48
GC AGGUGAGC CGAUCCAC CG	8.81	5.2	3.61	5.9	2.91
GC GUGCACAGC CGACACGUG CG	8.46	5.2	3.26	5.9	2.56
GGC CGAGCC CCGAGC CGG	8.32	8.5	−0.18	9.2	−0.88
GUG GCACAC CACACG GUG	4.72	6.1	−1.38	6.8	−2.08

Continued

TABLE 4. Continued

Oligomer	ΔG_{37}° (bulge)		(kcal/mol)		
	Measured	Predicted ^b Eq. 1	Difference	Predicted ^c Eq. 2	Difference
GUC ACUACUG CAGAUGA GAC	6.87	6.0	0.87	6.7	0.17
GGC GCGCAGCC CCGACGCG CGG	8.06	8.5	-0.44	9.2	-1.14
CGC CAGCACAGCC GCGAGUCGUG CGG	10.95	8.5	2.45	9.2	1.75
GCC GUGCUGAGCG CGGACACGAC CGC	7.85	8.5	-0.66	9.2	-1.36
CGC CACCACA GCC GCGAGUGGUG CGG	11.35	8.5	2.85	9.2	2.15
Average difference			0.65		0.15
Average deviation			1.3		1.0

^aMeasured values using Equation 3.^bPredicted values using Equation 1 (no interaction between the two bulge loops).^cPredicted values using Equation 2 (interaction between the two bulge loops).

of the two bulges is no greater when the bulges are inserted near each other (1 bp away) or further (up to 6 bp away), nor is the orientation of the bulge relative to each other (BB, Bb, or bB) important. Thus we can use the current model for the prediction of the influence of a single bulge loop to predict the influence of two bulge loops into a single duplex. The model is given below:

$$\Delta G_{37}^{\circ} \text{ (for insertion of two bulge loops)} \\ = 2 \times (-0.51\Delta G_{37(\text{stem})}^{\circ} + 0.85) + 0.72, \quad (2)$$

where $\Delta G_{37(\text{stem})}^{\circ}$ is the second least stable stem of the three stems. This model should improve the prediction

FIGURE 1. Plot of measured free energy change for the insertion of two single Group I nucleotide bulge loops into a duplex, $\Delta G_{37(\text{bulges})}^{\circ}$ versus predicted free energy value for the insertion of noninteracting two single nucleotide bulge loops (Equation 1) into a duplex as described in text. Line is the least-squares fit of the data points.

of RNA secondary structure and RNA stability from sequence.

Table 4 lists the measured free energy values for the insertion of two bulge loops into a duplex and the predicted values using Equation 1 (no interaction between the two bulge loops) and Equation 2 (includes the additional 0.7 kcal/mol interaction between the two bulge loops). The addition of the 0.7 kcal/mol interaction improves the prediction for almost two-thirds (32 of 50 duplexes) of the duplexes in Table 4. The average difference between the measured and predicted free energy value using Equation 2 is 0.15 kcal/mol. The average deviation between the measured and predicted free energy value for the insertion of two bulge loops into an RNA duplex using Equation 2 is 1.0 kcal/mol which is twice the value we measured for the insertion of a single bulge into an RNA duplex (Znosko et al. 2002). Twenty-eight of the 50 duplexes (56%) are predicted within 1 kcal/mol or 0.5 kcal/mol per inserted bulge and another 17 are predicted within 2 kcal/mol of the measured value. The remaining five oligomers (10%) have a predicted value that is more than 2 kcal/mol greater than the measured value. While this seems a large difference, five of the 37 oligomers (14%) with a single nucleotide bulge loop had a measured and predicted value that differed by more than 0.75 kcal/mol (Znosko et al. 2002). An examination of the oligomers in Table 4, which were poorly predicted, does not reveal any obvious structural feature that would explain the differences between the measured and predicted free energy values. Non-nearest neighbor interactions cannot be easily incorporated into current free energy prediction models (Mathews and Turner 2002) and may require an iterative process to arrive at a free energy value of an RNA containing a bulge loop.

Structural analysis of naturally occurring double nucleotide bulge loops

RNA FRABASE was used to search for three-dimensional structures of duplexes containing two bulge loops separated by up to 10 bp (Popenda et al. 2010). A total of 27 unique examples were identified (Ye et al. 1995; Carter et al. 2000; Ben-Shem et al. 2001, 2011; Berglund et al. 2001; Merianos et al. 2004; Berk et al. 2006; Walden et al. 2006; Weixlbaumer et al. 2008; Blaha et al. 2009; Agirrezabala et al. 2010; Armache et al. 2010; Bulkley et al. 2010; Jenner et al. 2010; Jin et al. 2010; Hainzl et al. 2011; Ng et al. 2011; Rabl et al. 2011; Demeshkina et al. 2012; Anger et al. 2013; Hashem et al. 2013; Amunts et al. 2014; Gogala et al. 2014). The secondary and tertiary structures of the duplexes are presented in the Supplemental Material. All of the bulge loop structures display a structure previously identified (Znosko et al. 2002). Of the 54 bulged nucleotides, 24 are intercalated, 27 are extrahelical, and three form base triples. In most of the examples, the bulged nucleotides have no discernible tertiary structural interactions. In four of the 27 structures, the bulge loop nucleotides are involved in tertiary interactions; these are displayed in Figure 2. The first example (panel A) is the only structure where the two bulged nucleotides appear to interact. In this case, the two bulged adenosines (1260 and 1275) on opposite strands are stacked. This leads to continuous stacking of C1262, A1261, A1275, and G1276. There are three examples where the bulged nucleotide is interacting with other residues. In panel B, the bulged U10 is stacked on adenosine residue 28 and forms a base triple with A13 and U24. In panel C, the bulged A5 is paired with G15 of a second molecule in the unit cell. In panel D, the bulged G (926) is stacked on a G residue at position 1505.

FIGURE 2. Secondary structure diagrams (left panel) and three-dimensional presentations (right panel) are shown. The bulged nucleotide is shown in CPK color and the intervening nearest neighbor pairs in red (top strand) and blue (bottom strand). (A) Position 1259 of the *T. thermophilus* small subunit rRNA. (B) Position 9 Bovine TAT-TAR complex. (C) Position 4 model U2-snRNA branchpoint duplex. (D) Position 922 of the *T. thermophilus* small subunit rRNA. Extracted from pdb files: 3UZ7, 1BIV, 1I9X, and 3IZ9.

These examples emphasize the structural flexibility imparted into the RNA when bulges are introduced. Further investigation will be needed to understand the forces responsible for the selection of the correct bulged nucleotide and its orientation in a complex structural motif.

MATERIALS AND METHODS

RNA synthesis and purification

Oligomers were synthesized on CPG solid supports (Applied Biosystems 392 DNA/RNA Synthesizer) utilizing phosphoramidites with the 2' hydroxyl protected as the *tert*-butyl dimethylsilyl ether from Glen Research. Oligomers underwent ammonia and fluoride deprotection, and a crude sample was purified using preparative *tlc* (*n*-propanol:ammonium hydroxide:water, 55:35:10) and Sep-Pak C18 (Waters) chromatography as previously described (Serra et al. 1994). Sample purity was determined through analytical *tlc* or HPLC (C-8) and was >95%.

Melting curve and data analysis

Optical melting experiments were performed as previously described in 1 M NaCl melt buffer (1 M NaCl, 0.02 M cacodylic acid, 0.001 M EDTA, pH 7.0) (Serra et al. 1994). Absorbance versus temperature profiles were fit to a two-state model with sloping base lines using a nonlinear least squares program, Meltwin (McDowell and Turner 1996). Thermodynamic parameters for duplex formation were obtained as previously described (McCann et al. 2011).

Determination of the contribution of bulge loops to duplex thermodynamics

The free energy of duplex formation can be approximated by the nearest-neighbor model (Xia et al. 2001). The free energy contribution of the bulged nucleotides was calculated from the experimental data and the nearest neighbor model according to Equation 3, where $\Delta G_{37}^{\circ}(\text{measured})$ is the experimentally determined value from the melts and $\Delta G_{37}^{\circ}(\text{duplex})$ is calculated from the nearest-neighbor model for the duplex as if it did not contain the bulges.

$$\Delta G_{37}^{\circ}(\text{bulges}) = \Delta G_{37}^{\circ}(\text{measured}) - \Delta G_{37}^{\circ}(\text{duplex}). \quad (3)$$

Phylogenetic analysis

A database of phylogenetically determined RNA secondary structures of 305 small subunit rRNAs, 169 large subunit rRNAs, 16 Group I RNAs, and seven Group II RNAs, was searched for duplexes containing two single nucleotide bulge loops (Andronescu et al. 2008). The loops were characterized by relative position within the duplex.

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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Influence of two bulge loops on the stability of RNA duplexes

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