

# How does RNA fold dynamically?

David Z. Bushhouse<sup>1,2†</sup>, Edric K. Choi<sup>1,2†</sup>, Laura M. Hertz<sup>1,2†</sup> and Julius B. Lucks<sup>1,2,3,4,5\*</sup>

**1 - Interdisciplinary Biological Sciences Graduate Program, Northwestern University, Evanston, IL 60208, USA**

**2 - Center for Synthetic Biology, Northwestern University, Evanston, IL 60208, USA**

**3 - Department of Chemical and Biological Engineering, Northwestern University, Evanston, Illinois 60208, USA**

**4 - Center for Water Research, Northwestern University, Evanston, IL 60208, USA**

**5 - Center for Engineering Sustainability and Resilience, Northwestern University, Evanston, IL 60208, USA**

**Correspondence to Julius B. Lucks:** Department of Chemical and Biological Engineering, Northwestern University, Evanston, Illinois 60208, USA [jb-lucks@northwestern.edu](mailto:jb-lucks@northwestern.edu) (J.B. Lucks) [@DavidBushhouse](https://twitter.com/DavidBushhouse) (D.Z. Bushhouse), [@Laura\\_M\\_Hertz](https://twitter.com/Laura_M_Hertz) (L.M. Hertz), [@LucksLab](https://twitter.com/LucksLab) (J.B. Lucks)

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## Abstract

Recent advances in interrogating RNA folding dynamics have shown the classical model of RNA folding to be incomplete. Here, we pose three prominent questions for the field that are at the forefront of our understanding of the importance of RNA folding dynamics for RNA function. The first centers on the most appropriate biophysical framework to describe changes to the RNA folding energy landscape that a growing RNA chain encounters during transcriptional elongation. The second focuses on the potential ubiquity of strand displacement – a process by which RNA can rapidly change conformations – and how this process may be generally present in broad classes of seemingly different RNAs. The third raises questions about the potential importance and roles of cellular protein factors in RNA conformational switching. Answers to these questions will greatly improve our fundamental knowledge of RNA folding and function, drive biotechnological advances that utilize engineered RNAs, and potentially point to new areas of biology yet to be discovered.

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## Introduction

RNA function has long been known to be deeply intertwined with RNA structure.<sup>1</sup> Recently, progress in high resolution static structural determination from x-ray crystallography,<sup>2</sup> NMR,<sup>3</sup> and cryo-EM<sup>4,5</sup> have given us deep insights into the intricacies of RNA structure beyond simple base pairing. These studies have revealed complex and diverse RNA folding interactions that give rise to the atomic architectures of ligand-binding pockets,<sup>6,7</sup> protein docking sites,<sup>8,9</sup> and catalytic active sites<sup>10,11</sup> that are at the center of RNA functions ranging from

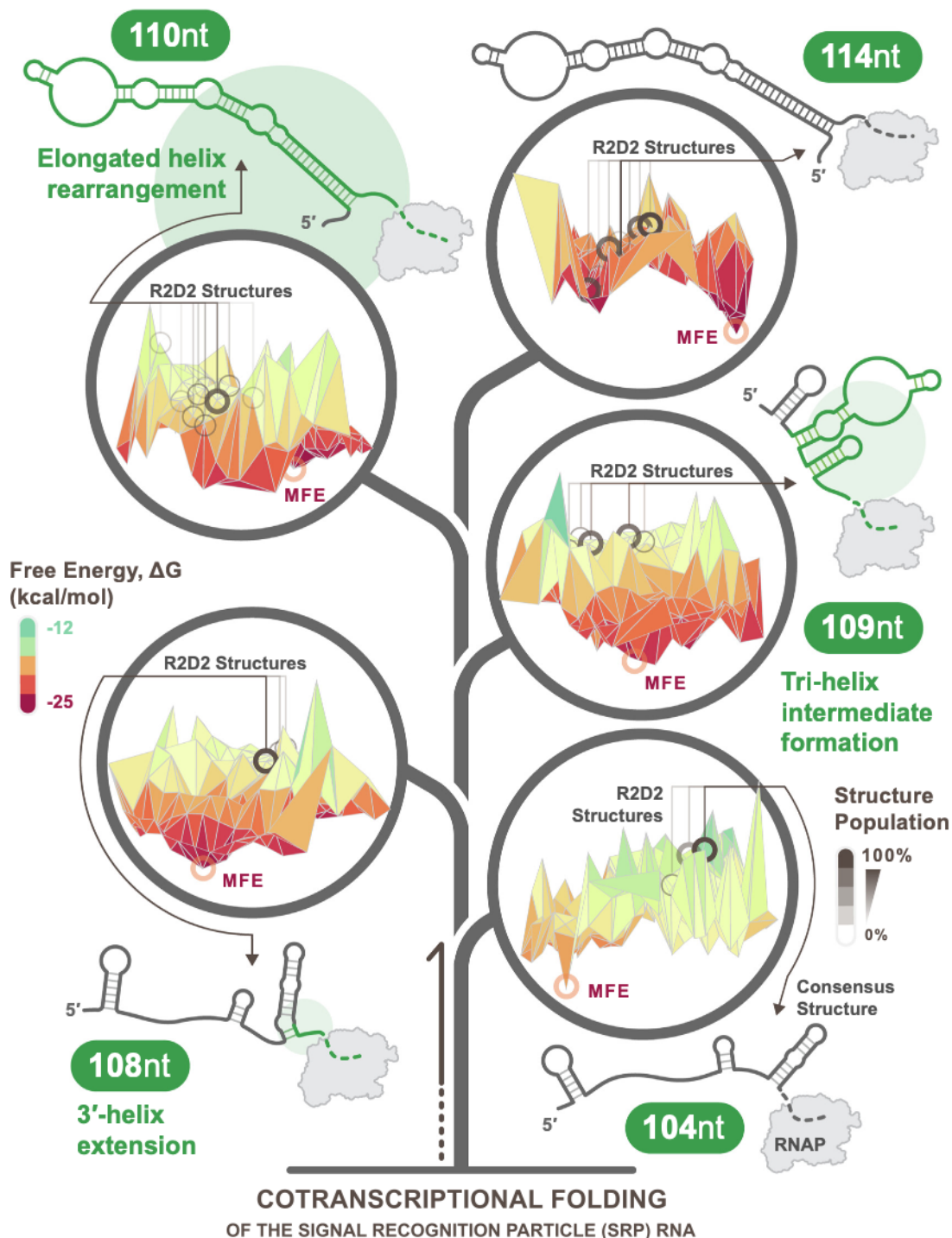
gene expression control,<sup>12,13</sup> RNA processing<sup>14–16</sup> and protein translation.<sup>17,18</sup>

However, a new appreciation for the dynamics of RNA folding suggests that a static viewpoint of RNA structures is only the tip of the iceberg.<sup>19</sup> Whether from RNA folding transitions during transcription,<sup>20</sup> or RNA switching conformations once synthesized,<sup>21</sup> new studies of RNA folding dynamics are proving critical for understanding known RNA function<sup>22,23</sup> and for drug discovery.<sup>24</sup> Well-known examples of these important features are riboswitch RNAs that regulate various aspects of gene expression by changing structural conformations in

response to ligand binding,<sup>25</sup> attenuator RNAs such as the TRAP system that regulate transcription by conformationally switching in response to protein binding,<sup>26,27</sup> and many other RNA regulatory mechanisms. Beyond explicit regulation there are also new appreciations for the role of RNA conformational switching in the activation of the spliceosome,<sup>28</sup> the assembly of the ribosome,<sup>29,30</sup> and as we later speculate for heterogeneous nuclear RNA proteins, in guiding dynamic RNA folding pathways in the cell. Even relatively simple RNA hairpin

structures, such as the one found in the signal recognition particle (SRP) RNA, have been shown to undergo conformational switching during transcription to establish their functional folds<sup>31,32</sup> (Figure 1). These are but a few of a growing list of examples revealing that dynamic RNA folding is important for broad arrays of RNA functions.

While dynamic RNA structure and folding has long been appreciated, advances in biophysical techniques are fueling new discoveries through probing RNA structural dynamics<sup>33–35</sup> and detect-





ing conformational changes tied to function. These studies raise new questions about how to conceptualize RNA folding and how to identify new mechanisms for which RNA dynamics are a core component.

Here we pose three prominent questions for the field that we believe are at the forefront of our knowledge of the importance of RNA folding dynamics and the role of conformational switching in RNA function. The first centers on the most appropriate biophysical framework to describe the complexities that are introduced when considering non-equilibrium conformational switching, especially in the context of a growing RNA chain during transcriptional elongation. The second is about the potential ubiquity of strand displacement – a process by which RNA can rapidly change conformations – and how this process may be present in broad classes of seemingly different RNAs. The third raises questions about the potential importance and roles of cellular protein factors in RNA conformational switching. Answers to these questions will greatly improve our fundamental knowledge of RNA folding and function, drive biotechnological advances that utilize engineered RNAs, and potentially point to new areas of biology yet to be discovered.

## What is the Appropriate Biophysical Framework to Describe Non-equilibrium Conformational Switching?

A biophysical framework for understanding “How RNA Folds” according to the thermodynamics of base pairing has long been established.<sup>1</sup> In thermodynamic renaturation conditions, RNA is understood to fold hierarchically, with secondary structures stabilizing first, creating an architecture to then establish tertiary interactions. Such a folding

regime is well described by a free energy landscape, a multi-dimensional surface that represents the free energies of all possible folds of a given RNA sequence<sup>36</sup> (Figure 1). While powerful, this framework does not capture essential details of cellular RNA folding such as transient structures that can form during cellular RNA synthesis.<sup>35</sup>

RNA folding begins during transcription, where due to the kinetics of RNA base pairing, nascent chains fold before the molecule is completely synthesized.<sup>37,38</sup> While these folds may undergo processing and other cellular interactions, recent studies suggest that for a large fraction of RNAs, these initial folds persist throughout the RNA molecule’s lifetime in the cell.<sup>39</sup> Moreover, nascent RNA structures are important for a wide array of cellular processes, such as the regulation of transcription, translation, and mRNA degradation in prokaryotes,<sup>40–42</sup> and splicing, polyadenylation, and 3’ end processing in eukaryotes.<sup>43–46</sup> Thus, understanding how RNAs fold during transcription is essential for understanding fundamental RNA biology.

Cellular RNA folding is different than the prevailing biophysical paradigm in two important aspects: i) RNAs do not fold all at once from a denatured full-length state, making the renaturation folding model incomplete, and ii) the sequence of the RNA changes during transcription, making the concept of a fixed free energy landscape of an RNA sequence incomplete. Overall, this raises a deep conceptual question: What is the appropriate biophysical framework to describe non-equilibrium nascent RNA folding?

Progress on this question has come primarily from computational studies of kinetic RNA folding processes. Computational modeling has chiefly considered kinetic RNA folding to be a stochastic process, where the RNA transitions to different folding states with different probabilities. These

**Figure 1. Transcription elongation leads to shifting free energy landscapes traversed by dynamically changing RNA structures.** Free energy landscapes were estimated from experimentally informed structural reconstructions of the *E. coli* SRP cotranscriptional folding pathway using Reconstructing RNA Dynamics from Data (R2D2).<sup>32</sup> R2D2 samples possible nascent RNA structures at each intermediate length and selects structures that are most consistent with experimental cotranscriptional RNA structure chemical probing data from each length. To model the possible free energy landscape of folding at each intermediate length, multidimensional scaling (MDS) was performed on the sampled structures at each length and plotted against predicted free energies. For visual clarity, the resulting plots were smoothed by calculating the mean free energy of all structures within a 10x10 coordinate area of the MDS plot. Each unique R2D2-selected structure is marked with a gray circle on the landscapes, where the shading corresponds to how frequently a structure is selected. These structures are notably far from the minimum free-energy structure (red circles) suggesting that the cotranscriptional folding process traverses out-of-equilibrium folding pathways on shifting free energy landscapes as nucleotides are added to the growing RNA chain. The illustrated secondary structures are consensus structures over 100 folding pathway predictions at each length by R2D2, with green highlighting and circles demonstrating the degree of change between previous structures. The addition of each nucleotide during transcription elongation presents an opportunity for significant conformational switching as the entire energy landscape reshapes.

probabilities are governed by transition rates that arise from local energy barriers between nearby possible RNA structures.<sup>47,48</sup> For example, Kinfold simulates the folding of a growing RNA chain by computationally searching for nearby possible structures, and stochastically making or breaking base pairs based on estimated rates to these nearby structures.<sup>49</sup> The Kinfold algorithm uses a similar framework, but chooses between the formation, breaking, or shifting of individual base pairs.<sup>50</sup> However, while these algorithms have great practical utility to simulate an RNA folding trajectory quickly and easily, they do not reveal a deeper understanding of general cotranscriptional RNA folding landscapes. Specifically, the stochastic approach samples one possible folding trajectory at a time out of a myriad of possibilities, making it intractable to gain a global view of possible folding trajectories of a given RNA molecule (Figure 1).

Recently, experimental techniques are also beginning to provide insights into non-equilibrium nascent RNA folding. Single molecule techniques, such as single molecule optical force spectroscopy, are able to measure structural transitions of a growing RNA chain at millisecond time resolution, revealing the switching mechanisms of riboswitches<sup>51</sup> and structural rearrangements of the SRP RNA during transcription.<sup>31</sup> New methods utilizing Förster resonance energy transfer are being used to label nascent RNAs to understand folding properties during key moments in the folding pathway.<sup>20,52</sup> And whole new approaches are using engineered superhelicases to simulate transcription elongation and characterize kinetically controlled decision landscapes.<sup>34,53</sup> Atomic structure determination techniques have also been applied to study transient structures and structural transitions.<sup>5,54</sup> For example, the coupling of NMR with photo-cage labeling of RNA strands has allowed researchers to trigger the conformational switching of an adenine riboswitch by uncaging part of the expression platform and using NMR to watch it invade the aptamer.<sup>54</sup> Recently, single particle cryo-EM has been used to reveal local conformational dynamics in an exterior helix of the *Tetrahymena* ribozyme, and rotational flexibility of the R-loop in the target-bound CRISPR-Cas9 complex.<sup>5,55</sup> And high throughput chemical probing approaches have been developed that can probe nascent RNA structures in transcription elongation complexes,<sup>35</sup> that when combined with new modeling algorithms can reconstruct possible cotranscriptional folding pathways from this experimental data.<sup>32</sup> This approach is revealing details of riboswitch mechanisms, such as strand invasion pathways that allow expression platform folding in the absence of ligand but are blocked by aptamer-ligand interactions,<sup>35,56,57</sup> and rearrangement pathways of the SRP RNA molecule that were corroborated by single molecule techniques.<sup>32</sup> Collectively, these methods have pushed the field forward in

experimentally understanding the physical parameters of cotranscriptional folding pathways and are permitting researchers to better hypothesize how RNA undergoes dynamic conformational switches during transcription.

As an example of how these techniques can reveal new insights into co-transcriptional RNA folding, we combined experimental cotranscriptional structure probing of the SRP RNA folding pathway<sup>35</sup> with computational structure prediction algorithms<sup>32</sup> to reconstruct possible free energy landscapes that the SRP RNA may traverse during transcription (Figure 1). In this reconstruction, we observe that the elongating transcript traverses vastly changing landscapes, where every transcribed nucleotide transforms the conformational ensemble space. At shorter RNA lengths the SRP landscape is shallow, with few minima creating many possible folds with similar free energies. However, as the RNA chain grows longer, deeper minima emerge in the free energy landscape, reflecting the resolution of folding possibilities into fewer states. In between these two regimes the SRP RNA dramatically rearranges its fold from a tri-helix intermediate at 109 nt to an elongated helix at 110 nt, though it is not clear from the energy landscape reconstructions alone how this may happen. Here a deeper understanding of the mechanisms by which RNAs may interconvert between states may be important for understanding how RNAs can traverse rugged and changing free energy landscapes.

As a field, we appear poised to significantly elaborate our biophysical framework of RNA folding by incorporating the physics of cotranscriptional RNA folding as it traverses free energy landscapes. In this view, the RNA fold does not move along a fixed free energy landscape, but instead navigates a shifting landscape that transforms with each transcribed nucleotide<sup>19</sup> (Figure 1). Each new nucleotide could reveal new peaks and troughs in the landscape, allowing dynamical switching between conformations that on a fixed landscape may appear to be separated by insurmountable free energy barriers. Deeper research from a range of approaches is needed to confirm and refine this model. Applying this framework to a diverse set of RNA sequences may help us understand general principles of rapid dynamical conformational switching at the heart of many RNA functions.

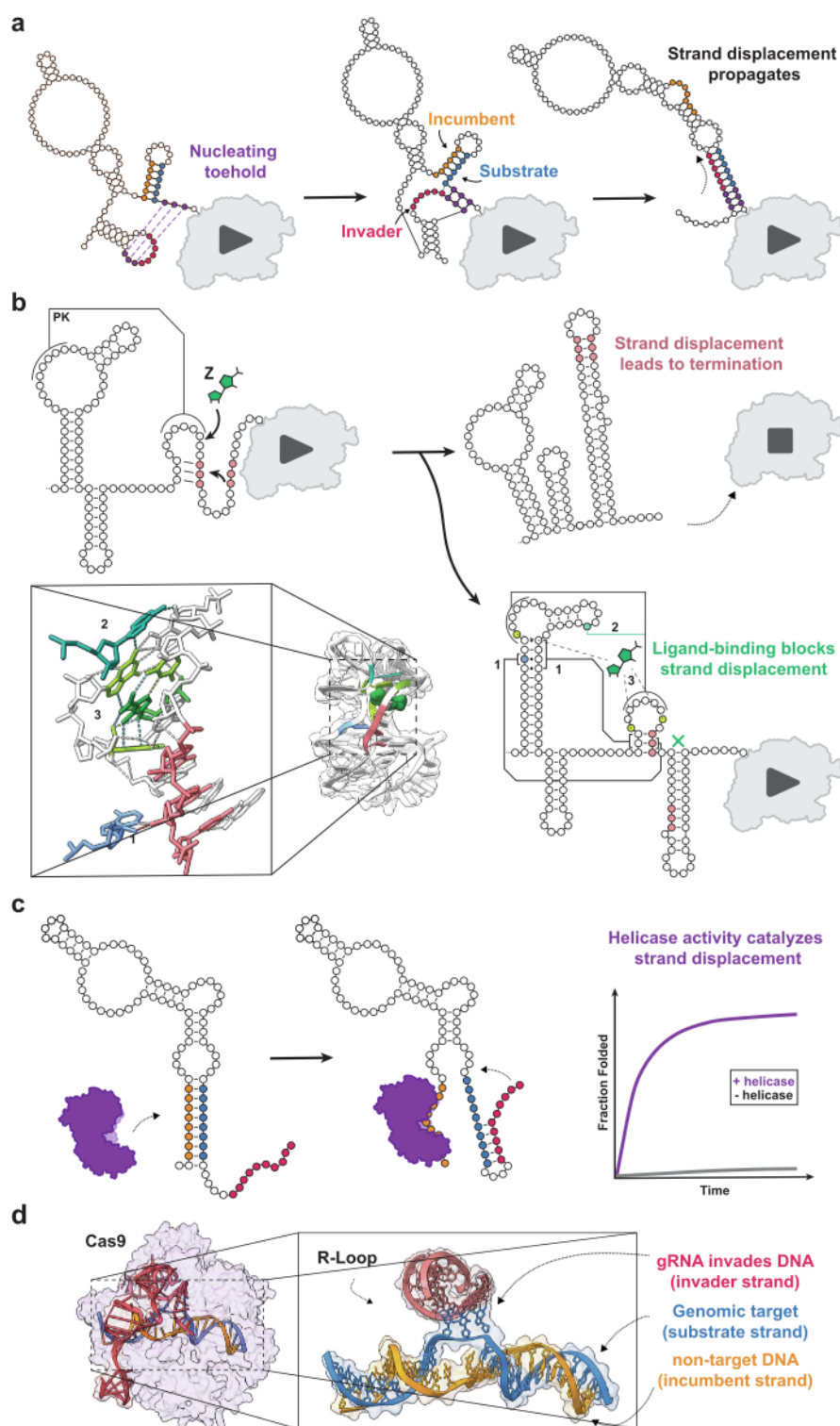
## How Widespread is Strand Displacement as a General Feature of RNA Conformational Switching?

Large RNA conformational changes can present significant free energy barriers to structural rearrangement, requiring milliseconds to hours to



occur – a time window that may limit cellular functions.<sup>19</sup> An intriguing solution to this challenge is ‘strand displacement’, a process by which RNA structures can efficiently rearrange through small nucleotide fluctuations that facilitate base pair exchange.<sup>58</sup> ‘Strand displacement’ or ‘strand exchange’ refers to a process whereby a double-stranded region comprising a substrate strand and

incumbent strand is disrupted by a third invader strand, resulting in the formation of a new invader-substrate duplex (Figure 2(a)).<sup>58</sup> While strand displacement can occur with three separate nucleic acid strands, in the context of RNA conformational switching, the incumbent, substrate, and invader strands may all be regions of the same RNA molecule.



Strand displacement was first identified by LeCuyer and Crothers, who proposed that this mechanism would allow mutually exclusive helices with similar free energy to quickly interconvert in a 'break-one-form-one' fashion comparable to branch migration.<sup>59,60</sup> Since then, strand displacement has been observed in two major areas of RNA function: i) dynamically during cotranscriptional folding,<sup>32,35,56,57</sup> and ii) within some post-transcriptional rearrangements and cellular factor binding events<sup>61,62</sup> (Figure 2).

Since cotranscriptional folding favors local structures that can form immediately after synthesis, there is a potential for kinetic traps that consist of metastable RNA structures that must be rearranged into more thermodynamically stable or functional folds. In cases of conformational switching during the timescale of transcription, RNAs appear to utilize strand displacement mechanisms. This is especially true for transcriptional riboswitches, *cis*-acting non-coding RNAs that regulate transcription elongation through ligand-mediated conformational switching.<sup>25</sup> Several studies of transcriptional riboswitches reveal that, in these cases, the genetic decision is the result of a competition between two mutually exclusive structures that exchange via a strand displacement mechanism (Figure 2(b)). Ligand binding can then bias the folding pathway by stabilizing certain structures that prevent strand displacement by a downstream intrinsic terminator structure, allowing ligand binding to directly lead to the enactment of a genetic regulatory decision (Figure 2(b)).<sup>35,57</sup> Furthermore, for some transcriptional riboswitches, the presence of an additional competing strand can flip this logic, with strand displacement leading to the retention of a strand that competes with the terminator structure.<sup>56</sup>

Moreover, cotranscriptional strand displacement can be guided by higher order transcriptional dynamics. For example, while the *E. coli* SRP RNA must form a long hairpin structure for proper function, it was shown that efficient folding of this structure utilizes RNA polymerase (RNAP) pausing to promote the formation of a labile 5' hairpin structure that then rearranges to form the extended functional fold.<sup>63</sup> Recent studies strongly suggest that this rearrangement proceeds via a strand displacement pathway<sup>31,32</sup> (Figure 2(a)).

In addition to mediating structural rearrangement during transcription, strand displacement could play a role in many post-transcriptional cellular contexts where RNA conformational switching is functionally important. For example, structure-probing studies showed that the NEAT1 lncRNA has two functionally distinct folds – one with many short helices that has unknown biological function, and another with a long-range helix that facilitates paraspeckle formation to regulate gene expression.<sup>64</sup> Here, strand displacement could be an advantageous mechanism of conformational switching. Such a mechanism may offer a much faster route to rearranging RNA structures that occur on rugged free energy landscapes (Figure 1), because rather than having to melt local structures within two complementary regions prior to refolding, strand displacement could allow local structural interconversions that would each have relatively smaller energy barriers to traverse.<sup>61</sup> In fact, DNA nanotechnology studies have found that strand displacement kinetics can occur on the microseconds timescale,<sup>65</sup> similar to the timescales of local nucleotide fluctuations.<sup>66</sup> Moreover, because strand displacement can rely on only a few nucleating base pairs for initiation (Figure 2(a)), conformational switching could be tightly regulated by controlling the formation of these base



**Figure 2. Strand displacement is a widely observed mechanism of RNA conformational switching.** Illustrations of several well-studied strand displacement processes. (A) Cotranscriptional strand displacement for structural rearrangement during transcription of the *E. coli* SRP RNA. After transcription of 110 nucleotides, three nucleating base pairs (purple) can form between the 3' end and the loop of a 5' hairpin structure. This interaction seeds a strand displacement reaction in which an invading strand (red) outcompetes an incumbent strand (orange) for binding to a substrate strand (blue), causing a global conformational change that remodels the nascent RNA from a tri-helix structure to a single elongated helix.<sup>32</sup> (B) Strand displacement in RNA genetic switches. Secondary structure diagram of the cotranscriptional folding pathway of the *Clostridium beijerinckii* pfl ZTP/ZMP riboswitch. In the absence of ligand, a strand displacement event seeds the formation of a strong intrinsic terminator. In the presence of ligand, numerous tertiary structural interactions, including: (1) ribose zippers, (2) an A-minor triple, and (3) hydrogen bonds between ligand and the binding pocket stabilize the holo-aptamer structure such that strand displacement is unable to efficiently proceed. This results in the formation of an alternative anti-terminating structure<sup>57,101</sup> (PDB ID: 4ZNP). (C) Protein-catalyzed strand displacement. The Mss116p non-processive RNA helicase has been shown to catalyze strand displacement by capturing the incumbent strand (orange), freeing the substrate strand (blue) to anneal to the invading strand (red)<sup>68,102</sup> (PDB ID: 3I5X). (D) CRISPR gene editing. Cryo-EM structure model of target-bound Cas9 illustrates the molecular basis of strand displacement in Cas9-gRNA-DNA recognition<sup>55</sup> (PDB ID: 7S38). The gRNA (red) forms a heteroduplex with the DNA target sequence substrate (blue) by invading the dsDNA, forcing the incumbent non-target strand (orange) to dissociate.



pairs.<sup>58</sup> For example, large-scale conformational changes that proceed via strand displacement can be catalyzed by nonprocessive RNA helicases like DEAD-box RNA helicases<sup>67,68</sup> (Figure 2(c)). Thus, strand displacement may underlie the regulation of some RNAs with post-transcriptional function.

Strand displacement also appears to play a role in biological mechanisms that utilize RNA for sequence recognition such as CRISPR-Cas9. CRISPR-Cas9 is a bipartite system composed of the Cas9 endonuclease and a guide RNA (gRNA) which targets the complex to complementary genomic loci. After the Cas9-gRNA complex binds tightly to a protospacer-adjacent motif, the single-stranded seed region of the gRNA is positioned to invade the target genomic site, opening up the dsDNA duplex and forming a RNA:DNA heteroduplex via strand displacement<sup>55,62,69</sup> (Figure 2(d)). In this way, Cas9 facilitates strand displacement, which can typically only disrupt double-stranded structures if there are transient fluctuations in base pairing at the ends. Structural studies of other Cas enzymes have shown similar dynamics.<sup>70</sup> While different from Cas9, argonaute proteins also pre-organize the 5' end of their miRNA (seed region), allowing increased discrimination between sites with and without a single mismatch in this region at rates much higher than would be expected from simple nearest-neighbor thermodynamic rules.<sup>71</sup> Thus, while strand displacement was first identified for RNA helical switching, it appears to have also been inherited in RNA-protein binding and recognition mechanisms.

Strand displacement allows RNAs to efficiently traverse large energetic barriers to enact conformational switching.<sup>61</sup> The process can be tightly regulated by controlling the spatial proximity of the three participating strands, and by controlling the formation of nucleating base pairs. This raises the intriguing question: How broadly-distributed is strand displacement as a mechanism for RNA conformational switching? An evolutionary study of bacterial P RNA, SRP RNA, the *trp* operator leader, hepatitis delta virus ribozyme, *Levivirus* maturation gene, and a class of S-adenosylmethionine riboswitches, found evidence that the cotranscriptional folding pathways of these RNAs likely feature evolutionarily conserved transient metastable structures that guide the RNA to adopt its final fold, potentially by helping the RNA avoid kinetic traps.<sup>72</sup> Further bioinformatic analysis could be used to investigate other RNA systems that have sequences that could facilitate strand displacement, with experimental techniques used to corroborate these possibilities. As techniques develop to understand how RNA dynamically switches conformations in different contexts, we predict strand displacement will be shown to be a general-purpose strategy for guiding RNA structural transitions throughout biology.

## How do Proteins Mediate Cotranscriptional RNA Conformational Switching?

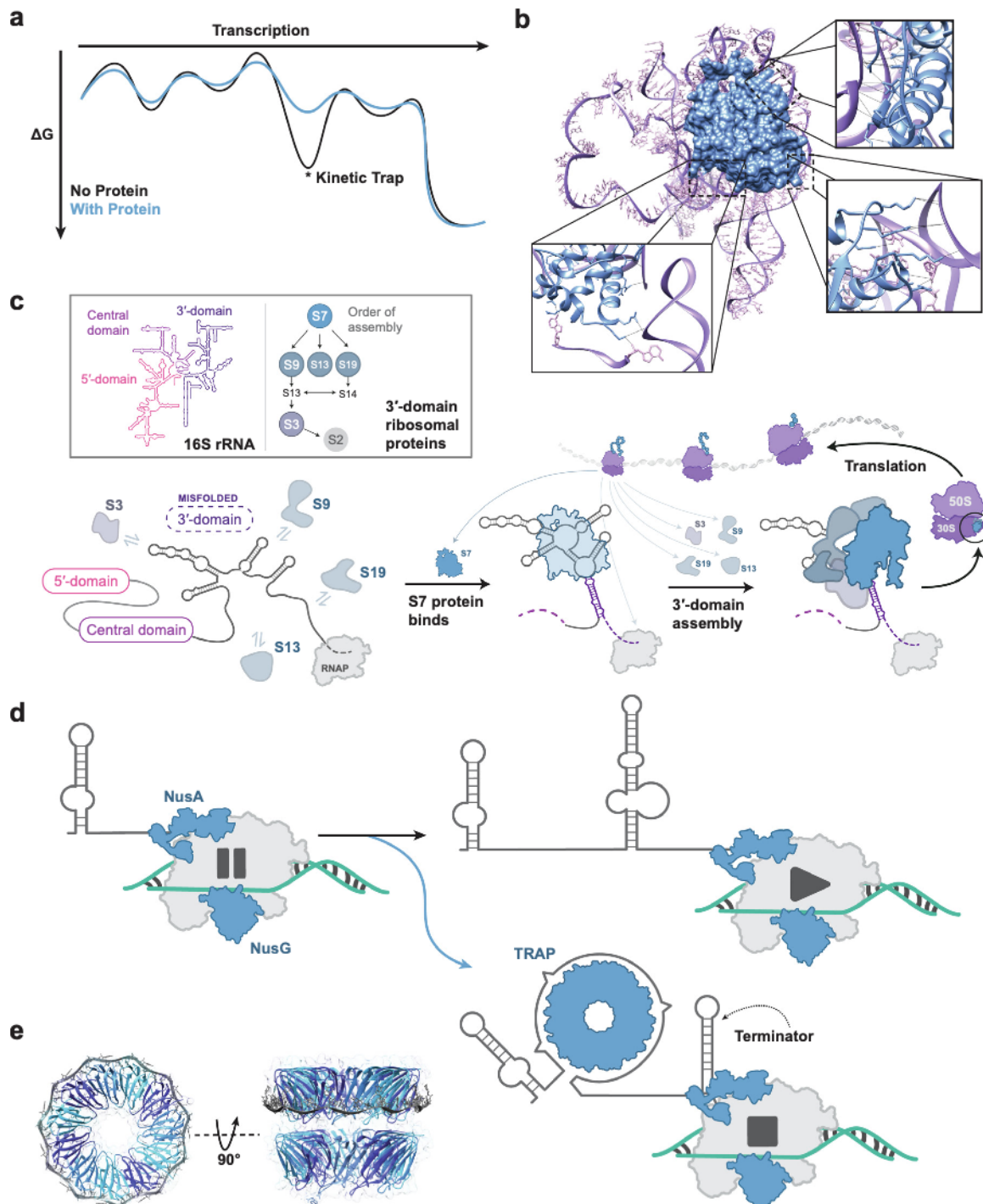
The biophysics of cotranscriptional folding creates a key challenge for cellular RNAs – the propensity of nascent RNAs to fold into non-functional kinetically trapped states at local minima in a rugged and changing free energy landscape (Figure 1). However, cellular RNA folding does not occur within a vacuum, but in a complex milieu filled with other cellular components such as proteins that can bind to RNA, influencing its folding and function.<sup>73</sup> For example, RNA folding chaperones can prevent RNAs from folding into non-functional states by mitigating kinetic folding traps<sup>74</sup> (Figure 3(a)). This raises the fundamental biological question: How do proteins mediate dynamic RNA folding through RNA conformational switching?

A confluence of biophysical and biochemical methods are now revealing the details of how cellular factors associate with transient intermediate structures of nascent RNAs to help the RNA navigate its complex conformational landscape towards its functional form.<sup>75</sup> Several aspects of biological processes appear to be particularly influenced by RNA-protein interactions: the assembly of ribonucleoprotein complexes (RNPs) (Figure 3(b), (c)), the feedback between nascent RNAs and the elongation complex (Figure 3(d)), eukaryotic RNA processing and gene expression, and RNA-driven formation of nuclear compartments.

RNPs can assemble cotranscriptionally from their molecular components and involve many RNA-protein interactions, including those that influence the fold of the RNA molecule as part of the assembly process<sup>76</sup> (Figure 3(c)). Bacterial ribosomal biogenesis is among the most well-studied systems exemplifying the importance of timely incorporation of cellular factors during RNP assembly. Ribosomal assembly begins during transcription, and is intimately coupled with processing and maturation of the pre-rRNA.<sup>77,78</sup> The hierarchical addition of ribosomal proteins (RPs) to ribosomal subunits can lead to structural changes in the pre-rRNA, with each newly incorporated RP stabilizing the folded structure of its immediate RNA binding site, in turn stimulating structural changes in adjacent pre-rRNA residues that help recruit other proteins to the complex.<sup>29,30</sup> For example, the nascent 16S pre-rRNA 3' domain initially misfolds, but the binding of the S7 RP structurally remodels 16S to enable subsequent hierarchical assembly of the 30S ribosomal subunit<sup>79</sup> (Figure 3(b), (c)). Overall, ribosome biogenesis is illustrative of how proteins can smooth the RNA folding landscape by preferentially stabilizing productive folding intermediates.<sup>80</sup>

Another key area where RNAs and proteins intimately interact is within the transcription elongation complex itself. Transcriptional dynamics arising from the two-way interplay between nascent RNA folding and the transcription elongation complex have been shown to play important roles in the temporal coupling of RNA synthesis and conformational switching.<sup>54,81</sup> Furthermore, RNAP pausing has been shown to influence RNA folding pathways, either by giving time for the RNA to rearrange into

more thermodynamically stable structures or by selecting for certain metastable structures that prevent downstream sequence from interfering with labile folds.<sup>63</sup> Elongation factors that affect the rate of transcription or bind to nascent RNA structures themselves can also play roles in guiding nascent RNA through its free energy landscape. For example, the elongation factor NusA forms transient contacts with the elongating RNA chain, allowing this protein to modulate transcriptional dynamics in response to the formation of certain RNA structures





and vice versa.<sup>63,82–85</sup> This points to broader roles for elongation factors such as NusA, NusG and others in facilitating cotranscriptional RNA folding (Figure 3(d)).

Principles from RNP assembly and protein-mediated modulation of transcriptional dynamics can both be seen in the intricate *trp* operator leader system, which employs RNAP pauses and binding of the 11-mer *trp* RNA-binding attenuation protein (TRAP, Figure 3(e)) to nascent RNA for the regulation of gene expression.<sup>86,87</sup> During transcription of the *trp* leader, NusA- and NusG-stimulated RNAP pauses allow enough time for holo-TRAP to bind the elongating transcript at repeating tryptophan codons, conformationally switching the RNA to form a terminator hairpin that attenuates transcription of the downstream tryptophan biosynthetic operons<sup>86–89</sup> (Figure 3(d), (e)). These examples reveal the importance of a tight interplay between protein binding and dynamic RNA folding, as the timely addition of proteins can aid other segments of RNA to fold into place, carving a direct pathway towards functionality through an otherwise incredibly rugged energy landscape.

Eukaryotes have evolved additional mechanisms that leverage RNA-protein interactions to influence RNA folding. In eukaryotes, nascent transcripts bind a diverse set of proteins throughout their lifetimes in the nucleus, forming heterogeneous ribonucleoprotein complexes (hnRNPs).<sup>90,91</sup> Due to their ability to both recognize DNA and RNA structures and recruit necessary protein components for particular processes, hnRNPs play crucial roles in almost every aspect of RNA biogenesis, from transcription and splicing to nuclear export and translation.<sup>92–95</sup> In addition, there is now evidence that nascent RNA structures themselves may play an equally valuable role by driving the formation of macromolecular assemblies within the nucleus, for example acting as seeds to drive spatial localization of otherwise diffusive non-coding RNA and protein molecules.<sup>96</sup> Ultimately, much

work remains to elucidate the precise mechanisms by which hnRNPs promote, suppress, and guide nascent RNA folding dynamics, as well as the role of RNA structure and conformational switching in these processes.

Regardless of the environment, the propensity of RNA to fold into local kinetically trapped states that prevent proper folding and function creates a need for mechanisms to resolve misfolding. While conformational switching mechanisms like strand displacement are nucleic acid-centric, the mechanisms described above demonstrate how proteins have evolved to also address nucleic acid folding challenges. Utilizing proteins as dynamic RNA folding chaperones could allow advantageous modularity, allowing the RNA sequence to develop into highly specialized final structural states that ultimately influence cellular events. And while this question is limited in scope to proteins, as techniques develop, the same questions could be taken further to understand how chemical modifications,<sup>97</sup> temperature,<sup>98</sup> small molecules,<sup>22</sup> and even other RNAs<sup>99</sup> influence RNA conformational dynamics. Ultimately, more work investigating a connection between proteins and the dynamic folding free energy landscape of RNAs could reveal an exciting new area of RNA biology to explore.

## Discussion

Innovative computational and experimental techniques are shedding new light on the complexities of the cellular RNA folding problem. An increased appreciation for the predominance of kinetically-driven processes necessitated by RNA biogenesis is causing a re-thinking of classic RNA folding paradigms.

Previous conceptualizations of the RNA folding problem posited that the minimum free energy structure was likely the final folded state of most RNAs.<sup>1</sup> However, later developments have estab-

**Figure 3. Protein-RNA interactions guide cotranscriptional folding, influence transcription dynamics, and determine gene expression outcomes.** (A) Cartoon representation of how a protein can alter an RNA folding free energy landscape, allowing alternate folding pathways to, for example, avoid kinetic traps. (B) Three-dimensional structure of the *E. coli* ribosomal protein S7 binding to 16S rRNA<sup>79</sup> (PDB ID: **4V9P**). Hydrogen bonding interactions with the rRNA are shown in the insets. (C) Cartoon of ribosomal protein S7 binding to nascent 16S rRNA during transcription as a portion of ribosome biogenesis leading to a fully functional ribosome. The ribosome can then perform its function of protein translation. In some cases, such as for S7 depicted, the translated protein can bind to nascent RNA during transcription, participating in hierarchical ribosome assembly pathways. (D) Cartoon example of transcription attenuation through the 11-mer *trp* RNA-binding attenuation protein (TRAP) mechanism. As the 5' UTR of the *trp* attenuator is transcribed, RNA polymerase reaches a NusA-NusG mediated pause site. In excess tryptophan conditions, tryptophan cooperatively binds between adjacent subunits of the 11-mer TRAP complex, which can bind the 5' UTR allowing an intrinsic terminator to form. In limiting tryptophan conditions, TRAP does not form nor bind the 5' UTR. Instead, an anti-terminating stem forms, allowing the RNA polymerase to continue transcribing the downstream operon.<sup>87,88</sup> (E) Three-dimensional structure of the *Geobacillus stearothermophilus trp* RNA-Binding Attenuation Protein bound to single-stranded RNA (PDB ID: **1C9S**).

lished that RNAs are more accurately described as adopting an ensemble of structures occupying multiple local minima of a rugged energy landscape.<sup>19</sup> Detailed studies of specific RNA systems are revealing how the process of transcription guides structural ensembles through ever-shifting energy landscapes, prompting us to revisit our conceptual framework for understanding the principles of RNA folding, and how these principles manifest in RNA function. Interestingly, cotranslational protein folding may serve as a guide for a deeper understanding of cotranscriptional RNA folding. For example, cotranslational folding is commonly described as a nested energy landscape, where the volume of conformational space is proportional to the length of the nascent polypeptide.<sup>100</sup> How a similar conceptualization of cotranscriptional folding can be woven in with the inherent ruggedness of RNA folding landscapes will be interesting to develop further.

Ultimately, the dynamic aspects of cellular RNA folding provide nature with an expanded complexity of RNA folds beyond those attainable in equilibrium. It is our hope that the three central questions posed above will inspire the next wave of research that probes a deeper understanding of this RNA folding regime and how it facilitates myriad cellular functions. We anticipate answers to these questions will drive fundamental understanding and accelerate developments in RNA-based biotechnologies.

## Author Contributions

All authors conceptualized, wrote, reviewed, and edited the manuscript.

## CRedit authorship contribution statement

**David Z. Bushhouse:** Conceptualization, Writing – original draft, Writing – review & editing. **Edric K. Choi:** Conceptualization, Writing – original draft, Writing – review & editing. **Laura M. Hertz:** Conceptualization, Writing – original draft, Writing – review & editing. **Julius B. Lucks:** Conceptualization, Writing – original draft, Writing – review & editing.

## DECLARATION OF COMPETING INTEREST

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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† These authors contributed equally to this article.

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