

RNA-DNA Triplex Formation by Long Noncoding RNAs

Yue Li,¹ Junetha Syed,¹ and Hiroshi Sugiyama^{1,2,*}

¹Department of Chemistry, Graduate School of Science, Kyoto University, Kitashirakawa-oiwakecho, Sakyo-ku, Kyoto 606-8502, Japan

²Institute for Integrated Cell-Material Sciences (WPI-iCeMS), Kyoto University, Yoshida-ushinomiya-cho, Sakyo-ku, Kyoto 606-8501, Japan

*Correspondence: hs@kuchem.kyoto-u.ac.jp

<http://dx.doi.org/10.1016/j.chembiol.2016.09.011>

Long noncoding RNAs (lncRNAs) play a pivotal role in the regulation of biological processes through various mechanisms that are not fully understood. Proposed mechanisms include regulation based on RNA-protein interactions, as well as RNA-RNA interactions and RNA-DNA interactions. Here, we focus on one possible mechanism that lncRNA might be using to impact biological function, the RNA-DNA triplex formation. We summarize currently available examples of lncRNA triplex formation and discuss the details surrounding orientation of triplex formation as one of the key properties guiding this process. We propose that symmetrical triplex-forming motifs, especially those in *cis*-acting lncRNAs, favor triplex formation. We also consider the effects of lncRNA structures, protein or ligand binding, and chromatin structures on the lncRNAs triplex formation.

Introduction

Over the past decade, advances in the field of genome sequencing technologies have provided a deeper understanding of the complex genomic architecture (Kapranov et al., 2007; ENCODE Project Consortium, 2004, 2012). The most surprising insights that genome sequencing has revealed is that only a small fraction of the mammalian DNA encodes protein-coding mRNA. However, this does not mean that the rest of the DNA is not transcribed, and we now appreciate that a large proportion of transcribed RNA corresponds to noncoding RNAs (ncRNAs), which are thought to play roles in important biological processes, particularly epigenetic regulation (Wilusz et al., 2009; Ponting et al., 2009; Guttman et al., 2009; Kung et al., 2013; Morris and Mattick, 2014).

Over the last several years, we have seen a steady increase in the interest in identifying ncRNAs and understanding their regulatory functions (Quinn and Chang, 2016; Bonasio and Shiekhattar, 2014; Batista and Chang, 2013; Lee, 2012). ncRNAs include a wide class of regulatory RNAs that can be classified into small ncRNAs such as microRNAs (miRNAs), small interfering RNAs (siRNAs), PIWI-interacting RNAs, and long noncoding RNAs (lncRNAs), which are more than 200 nt in length (Cech and Steitz, 2014; Kowalczyk et al., 2012). The majority of mammalian ncRNAs, including lncRNAs, are highly conserved, strongly supporting the view that these RNAs play important *in vivo* roles (Struhl, 2007). Several recent studies have provided evidence for biological functions of lncRNAs, especially in the regulation of gene expression at various levels (Rinn and Chang, 2012; Mattick, 2004). These findings are consistent with the observation that a large number of lncRNAs are transcribed from loci with proximity to protein-coding genes, such as enhancer RNAs and upstream antisense RNAs (Bonasio and Shiekhattar, 2014).

lncRNAs regulate gene expression through complex mechanisms that are not fully understood (Guttman and Rinn, 2012; Hung and Chang, 2010). It has been suggested that lncRNAs fulfill their regulatory functions by serving as molecular signals, decoys, guides, and scaffolds (Wang and Chang, 2011). The molecular mechanisms of lncRNAs can be simplified into these four

main archetypes, but an individual lncRNA is not confined to one archetype and may function using several different mechanisms. For example, during X chromosome inactivation, Xist RNA acts as both a molecular signal for the chromosome-wide repression of gene expression and a guide for the recruitment of polycomb repressive complex 2 (PRC2) (Van Bommel et al., 2016; Maclary et al., 2013; Galupa and Heard, 2015; Zhao et al., 2008). Telomeric repeat-containing RNA can bind to the human telomere-binding protein TLS/FUS, recruit histone-modifying enzymes, and regulate telomere length through histone modifications (Takahama and Oyoshi, 2013; Takahama et al., 2013). As these two examples suggest, lncRNA biological mechanisms converge on establishing intermolecular interactions with other biomolecules, such as proteins, DNA, and other RNAs. Therefore, one practical approach to expand our understanding of the lncRNA molecular mechanisms is to elucidate the RNA-protein and RNA-DNA interactions they engage in. lncRNAs can form RNA-protein complexes, which most commonly mediate the correct localization of chromatin regulatory proteins (Xist RNA) or recruitment of functional proteins with distinct protein interaction domains (HOTAIR) (Chu et al., 2015; Sarma et al., 2014; Tsai et al., 2010). In addition, lncRNAs can target specific DNA sequences by forming structures such as RNA-DNA hybrid duplexes or the RNA-DNA triplexes (Wang and Chang, 2011; Li et al., 2016). Given the structural complexity of lncRNAs, several different interactions might be involved in the molecular mechanisms relating to each individual lncRNA.

Several excellent reviews have covered the biological roles of RNA-DNA triplexes in general (Duca et al., 2008; Bacolla et al., 2015; Buske et al., 2011; Jain et al., 2008) and the molecular mechanisms of lncRNAs (Guttman and Rinn, 2012; Hung and Chang, 2010; Wang and Chang, 2011). Here, we focus on the current knowledge about RNA-DNA triplex formation by lncRNAs. We start by discussing the structure of RNA-DNA triplexes and summarizing recently reported examples of RNA-DNA triplex formation by lncRNAs. We then discuss the importance of the orientation of triplex formation to the mechanism of lncRNA actions and the possible ways proteins and ligands may regulate triplex formation.

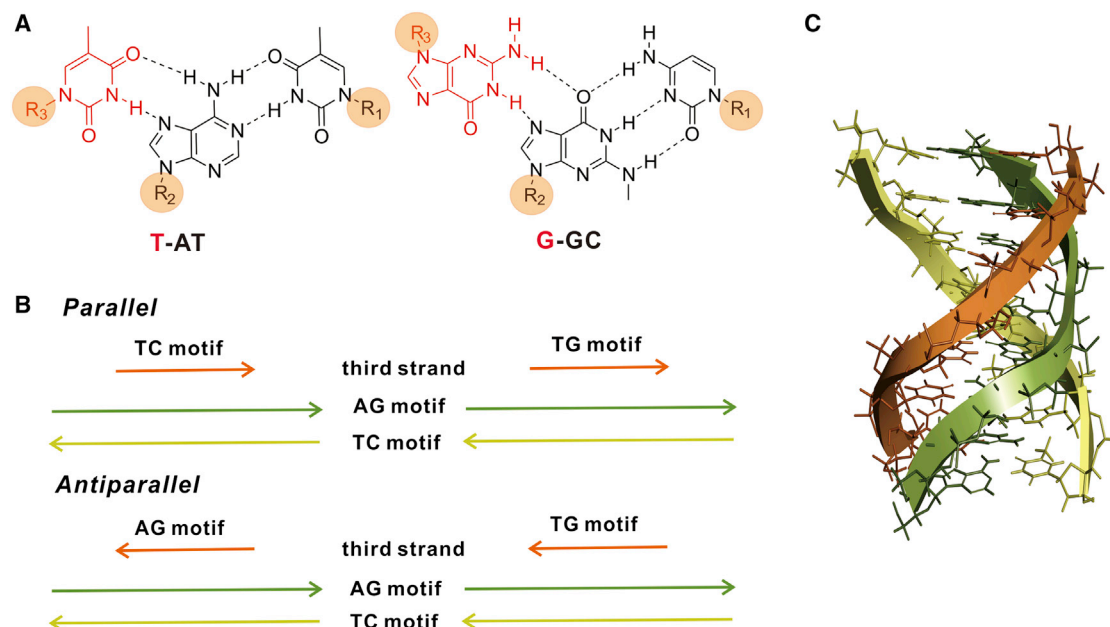


Figure 1. Triplex Formation between the Third-Strand Oligonucleotide and dsDNA

(A) Examples of Hoogsteen and reverse Hoogsteen hydrogen bonding in the triplex formation.

(B) Illustration of parallel and antiparallel triplex formation. The arrow indicates 5' to 3' direction.

(C) Structure of RNA-DNA triplex. The third-strand RNA (brown) binds to the major groove of the DNA duplex. The picture was generated by the reported RNA-DNA triplex structure (PDB: 1R3X) (Gottfredsen et al., 1998).

The Discovery and Nature of Triplex Structures

Not long after the discovery of the DNA double-helix structure in 1953 (Watson and Crick, 1953), Felsenfeld et al. (1957) suggested the possibility of triplex formation, whereby two pyrimidine strands and one purine strand would form a more complex structure. Over the decades, the evidence of triplex formation went from a scientific curiosity (Morgan and Wells, 1968; Beal and Dervan, 1991), to inspiring researchers to apply the concept of the triple helix to develop various sequence-specific triplex-forming oligonucleotides (TFOs) as DNA cleavage tools with major biological applications in chromosome mapping (Moser and Dervan, 1987; Le Doan et al., 1987; Broitman et al., 1987; Wells et al., 1988).

In the triplex structure, a third strand can insert into the major groove of the duplex structure with sequence specificity (Hoogsteen, 1959; De los Santos et al., 1989; Koshlap et al., 1997; Pasternack et al., 2002; Radhakrishnan and Patel, 1994a, 1994b). As shown in Figure 1, each triplet can be formed by either Hoogsteen hydrogen bonding or reverse Hoogsteen hydrogen bonding between the third base and the Watson-Crick base pair. Two types of triplexes can be formed based on the orientation of the third strand. Parallel triplex structures are formed by TA·T, CG·C⁺, and CG·G triplets, and antiparallel triplexes are formed by TA·A, TA·T, and CG·G. In the case of CG·C⁺, an acidic condition is favorable for triplex formation because the protonation of cytosine is required for effective triplet formation. Using these principles, one can determine the orientation of the third strand from its sequence (Figure 1B). The third strand with T(U)C motifs prefers to form parallel triplexes, whereas AG motifs prefer antiparallel triplexes, and GT(U) motifs can form either parallel or antiparallel triplexes.

The Hoogsteen hydrogen bonding by the third strand is weaker than the Watson-Crick hydrogen bonding in the duplex structure. Therefore, there are some limitations to the formation of the triplex structure, especially under physiological conditions (Duca et al., 2008). As mentioned above, an acidic condition is required for the protonation of cytosine during the formation of the CG·C⁺ triplet (De los Santos et al., 1989). Multivalent cations such as Mg²⁺ are frequently used in vitro to neutralize the unfavorable charge repulsion among three negatively charged nucleic acid strands (Floris et al., 1999). Molecular crowding conditions that mimic the cell environment are suggested to increase the melting temperature of triplex structures by the hydration effect (Nakano et al., 2014; Miyoshi and Sugimoto, 2008). Another study has shown that molecular crowding induces the formation of the i-motif structure by triplet repeat DNA oligomers under neutral pH, which implies an effect similar to the protonation of cytosine in triplex formation (Rajendran et al., 2010). To overcome these limitations, chemical modifications, including base, sugar, and backbone modifications, are used widely in the design of TFOs (Torigoe et al., 2001; Lacoste et al., 1997; Hojland et al., 2007; Roberts and Crothers, 1992; Lacroix et al., 2000). These modifications may facilitate triplex formation by increasing Hoogsteen hydrogen bonding or by restricting the third strand at the desired conformation by blocking the secondary structure formation, or other mechanisms.

Given that triplex formation requires base-pairing that is weaker than Watson-Crick or unusual protonation states for the bases involved, it is a valid and an important question to ask whether triplex structures exist in vivo. Many bioinformatics tools have been developed to analyze the potential for the

presence of triplex-forming motifs in the genome (Goñi et al., 2004; Buske et al., 2012). Bioinformatics analyses have suggested that a large population of triplex-forming motifs is present across the genome. Interestingly, these motifs tend to accumulate in the gene-regulatory regions, particularly in promoter regions, leading to a proposal that triplex formation might play significant regulatory roles in vivo. More direct evidence of triplex formation in vivo comes from using triplex-binding antibodies, as well as fluorescent molecules that selectively recognize triplex structures (Lee et al., 1987; Agazie et al., 1994, 1996; Lubitz et al., 2010). The results of studies using immunofluorescence of triplex-specific antibodies strongly suggest the formation of the triplex structure in the nucleus. Most importantly, these antibodies exhibit a stronger affinity for RNA-DNA triplexes than for DNA triplexes, which indicates the formation of RNA-DNA triplexes in vivo. Recent studies have also demonstrated the presence of specific proteins that can interact with triplexes in vivo. For instance, helicases such as RecQ have been shown to unwind triplex structures, which strongly suggests a potential regulatory role of triplexes in vivo (Jain et al., 2010; Maine and Kodadek, 1994; Dixon et al., 2008; Brosh et al., 2001).

Consistent with the RNA-DNA triplex, studies have suggested that RNA, as the third strand, forms more stable triplexes than its DNA counterparts (Roberts and Crothers, 1992). The first example of triplex formation between RNA polymers and the formation of intramolecular triplexes of RNA is well characterized. A 2'-methoxy group has also been introduced to stabilize the C3'-endo conformation of the sugar and to promote RNA-mediated triplex formation (Shimizu et al., 1992; Asensio et al., 1999). Therefore, third-strand RNA is favorable for the formation of the triplex. Recent studies have provided examples of lncRNA-mediated RNA-DNA triplex structures. In the following section, we discuss the reported lncRNAs that have been shown to form triplex structures.

lncRNA:DNA:DNA Triplexes In Vivo

Despite being recognized as a structure that RNA and DNA molecules can form in vitro very early on in the history of structural biology of nucleic acids, it took another 50 years to begin describing triplex structures in vivo and documenting their physiological relevance. In 2007 Martianov et al. (2007) reported triplex formation by a noncoding RNA that was transcribed as an interfering transcript from the minor promoter of the human gene encoding dihydrofolate reductase (*DHFR*). This interfering RNA transcript had a repressive effect on the transcription of *DHFR* mRNA from the major *DHFR* promoter located downstream. This effect was observed only when the sequence of interfering transcript overlapped with the sequence of the core major *DHFR* promoter. Subsequent studies have shown that the noncoding *DHFR* minor transcripts can bind to the transcription factor TFIIB, form a stable triplex with the G-track sequence in the major promoter, and impede the formation of the transcription preinitiation complex. The formation of RNA-DNA triplexes was confirmed by an electrophoresis mobility shift assay (EMSA) in vitro using either long RNA transcripts or short RNA oligonucleotides. The potential triplex-forming motif found in noncoding *DHFR* minor transcripts was a particularly AG-rich sequence, which suggests an antiparallel triplex formation between lncRNAs and the DNA sequence.

Promoter-associated RNAs (pRNAs) regulate the transcription of *rRNA* genes by forming stable RNA-DNA triplex with the promoter sequence (Schmitz et al., 2010). pRNAs form triple helical structures with the transcription factor binding site TTF-1 in the *rRNA* promoter. Most importantly, the formed RNA-DNA triplex specifically recognizes DNA methyltransferase DNMT3b, the enzyme that promotes heterochromatin formation in the *rDNA* promoter by DNA hypermethylation and subsequently silences the *rRNA* genes. The triplex-forming motif of pRNA contains mainly G, U, and C, and may form a parallel triplex with the promoter DNA based on the principle of triplex orientation.

Another interesting lncRNA, Fendrr, plays an important role in the development of the mammalian heart and body wall and has been suggested to regulate neighboring and distant genes by recruiting either PRC2 or the trithorax group/mixed lineage leukemia (TrxG/Mll) protein complexes in the target promoter (Grote et al., 2013; Grote and Herrmann, 2013). Recent studies have reported that Fendrr can form RNA-DNA triplexes in the promoter region of developmental genes such as *Foxf1* and *Pitx2*. The RNA-DNA triplex serves as an anchor to recruit PRC2, which in turn leaves the repressive histone marks in the above-mentioned gene promoters, thereby facilitating proper tissue differentiation. Triplex formation of Fendrr with target DNA sequences was confirmed by in vitro pull-down experiments using a UC-rich triplex-forming motif. Although acidic conditions and protonation of cytosine are believed to be required for formation of triplex of UC-rich motifs, this study suggested the possibility that UC-rich motifs do form RNA-DNA triplex in vivo.

This possibility of RNA-DNA triplex formation by UC-rich motifs under physiological conditions was supported by a study of the antisense lncRNA Khps1 (Postepska-Igielska et al., 2015). Khps1 is transcribed in the antisense direction to the proto-oncogene *SPHK1* and recruits the histone acetyltransferase p300/CBP to the *SPHK1* promoter. The recruited p300/CBP acetylates the histones, which results in an active chromatin confirmation in the *SPHK1* promoter, thereby providing access to the E2F1 transcription factor, and activates *SPHK1* expression. The experimental evidence reported in that study suggested that the resultant RNA-DNA triplex could be formed by the UC-rich motif of the third-strand lncRNA Khps1. Interestingly, the gel shift indicating triplex formation was observed with EMSA with 10 mM Mg^{2+} at pH 7.5 without the need for an acidic condition.

Recent studies of two lncRNAs have provided considerable evidence of the formation of RNA-DNA triple helices through AG-rich motifs of third-strand lncRNAs. One study reported that the antisense lncRNA PARTICLE is transcribed from the promoter of *MAT2A* when cells are exposed to low-dose radiation (O'Leary et al., 2015). Surface plasmon resonance experiments suggested that the AG motif of PARTICLE forms a triple helix with the *MAT2A* promoter to recruit PRC2, which in turn methylates the CpGs of the respective promoter, thereby negatively regulating the expression of *MAT2A* after irradiation. The next lncRNA to be reported was MEG3, which is transcribed from human maternally expressed gene 3 (*MEG3*) (Mondal et al., 2015). In an attempt to characterize the targeting mechanism of lncRNAs in chromatin, the authors used a modified chromatin oligo affinity precipitation method to perform genome-wide mapping of MEG3 binding sites; they found that these binding

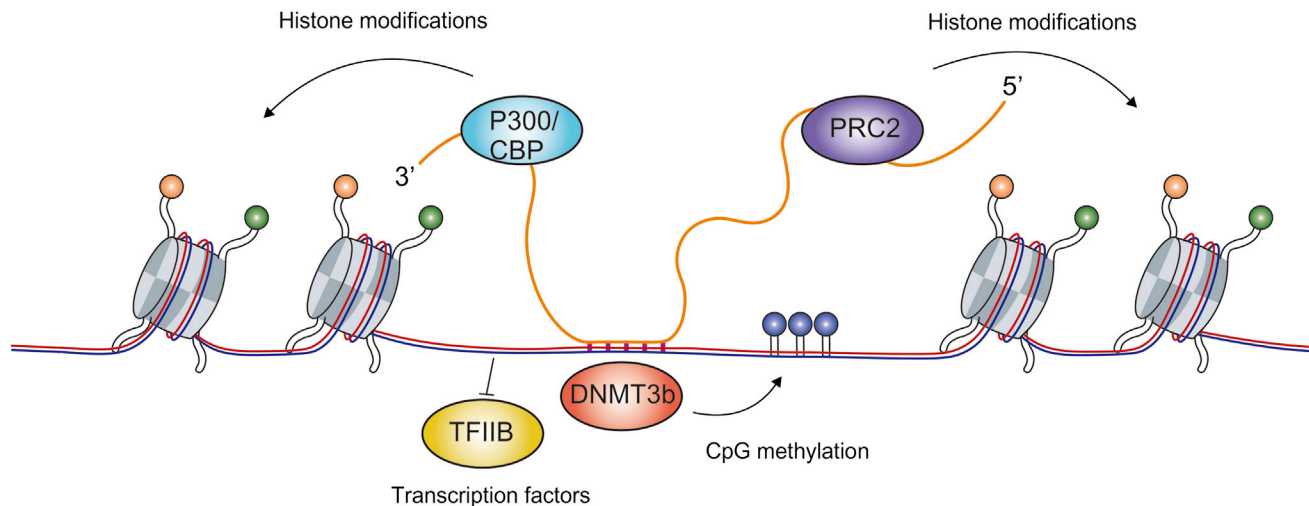


Figure 2. Examples of lncRNAs that Are Reported to Form RNA-DNA Triplexes

lncRNAs can form a triplex structure with a DNA sequence in either neighboring or distant genes and regulate their expressions. lncRNAs also interact with various regulatory proteins such as transcription factors and chromatin modifiers.

sites were enriched with AG-rich motifs. Subsequently, they performed various experiments using EMSA, circular dichroism spectra analysis, immunostaining with anti-triplex dA.2rU antibody, and pull-down experiments, and found that AG-rich motifs guide MEG3 to its target genes through the formation of RNA-DNA triplexes. MEG3 can also interact with PRC2 and facilitate recruitment of PRC2 to target sites, such as the distal regulatory elements of transforming growth factor β (*TGF- β*) genes. The systematic studies reported by Mondal et al. (2015) strongly suggested that RNA-DNA triplex formation may be a general mechanism of target-site recognition by lncRNAs.

In addition to the direct evidence of triplex formation by lncRNAs, other reports, especially those focusing on the genome-wide mapping of target binding sites of lncRNAs, have provided indirect evidence of the possibility of triplex formation by AG-rich motifs. For example, the method of chromatin isolation by RNA purification has been developed for the deep sequencing of lncRNA occupancy sites in the genome (Chu et al., 2011; Simon et al., 2011). HOTAIR and *Drosophila roX2* lncRNAs were found to preferentially occupy AG-rich DNA motifs across the genome. Considering the number of AG-rich motifs in these lncRNAs, it is likely that they target the AG-rich DNA motifs across the genome through triplex formation. A very recent report suggested lncRNA HOTAIR could function on mesenchymal stem cells by forming a triplex with the AG-rich motifs in the promoter of target genes (Kalwa et al., 2016). The authors predicted the triplex-forming motifs and analyzed the triplex formation by gel electrophoresis. The triplex formation by HOTAIR was thought to recruit PRC2 and LSD1, which are involved in gene silencing.

Although GT(U) motifs could form both parallel and antiparallel triplexes with the DNA duplex containing AG motifs according to the principle of triplex formation, so far there are no reports of triplex formation by lncRNAs with GU motifs. Actually, a previous report excluded the possibility of triplex formation between RNA with a GU motif and DNA with AG motifs by gel electrophoresis (Semerad and Maher, 1994). By testing the different combina-

tions of an RNA-DNA duplex and the third strand, the report suggested that only the DNA third strand with a GT motif could form a stable triplex with the DNA duplex with an AG motif but not the RNA. These results might explain why lncRNAs with GU motifs are not favorable in the triplex formation with the DNA duplex.

From these examples of RNA-DNA triplex formation by lncRNAs, one may conclude that triplex formation plays an important role in lncRNA function (Figure 2). lncRNAs inhibit the binding of transcription factors and dissociate the preinitiation complex; they also serve as guides for the recruitment of histone-modifying proteins or DNA methylation enzymes to specific sites in chromatin. For example, lncRNAs recruit PRC2 for histone modification or DNMT3b for DNA methylation in promoter regions. It has been suggested that during these epigenetic processes, triplex formation provides the general mechanism of action used by lncRNAs for the sequence-specific targeting of DNA sequences to chromatin modifiers in the genome.

Cis- and Trans-acting Mechanisms

As described above, lncRNAs can form triplexes and regulate gene expression in either a *cis*-acting manner (on neighboring genes) or a *trans*-acting manner (on distant genes) (Guttman and Rinn, 2012; Guil and Esteller, 2012). In the *cis*-acting manner, lncRNAs are usually transcribed from the promoter regions of target genes, form a triplex with promoter DNA sequences, and regulate gene expression. In the *trans*-acting manner, lncRNAs specifically target binding sites through triplex formation with distant DNA sequences. This suggests that triplex formation may be a general mechanism to explain why *trans*-acting lncRNAs can target specific genes in remote regions (Mondal et al., 2015). How the lncRNAs form triplexes with DNA sequences and whether there is any requirement for triplex formation by lncRNAs are important questions that need to be answered in order to understand the mechanism of triplex formation by both *cis*-acting and *trans*-acting lncRNAs.

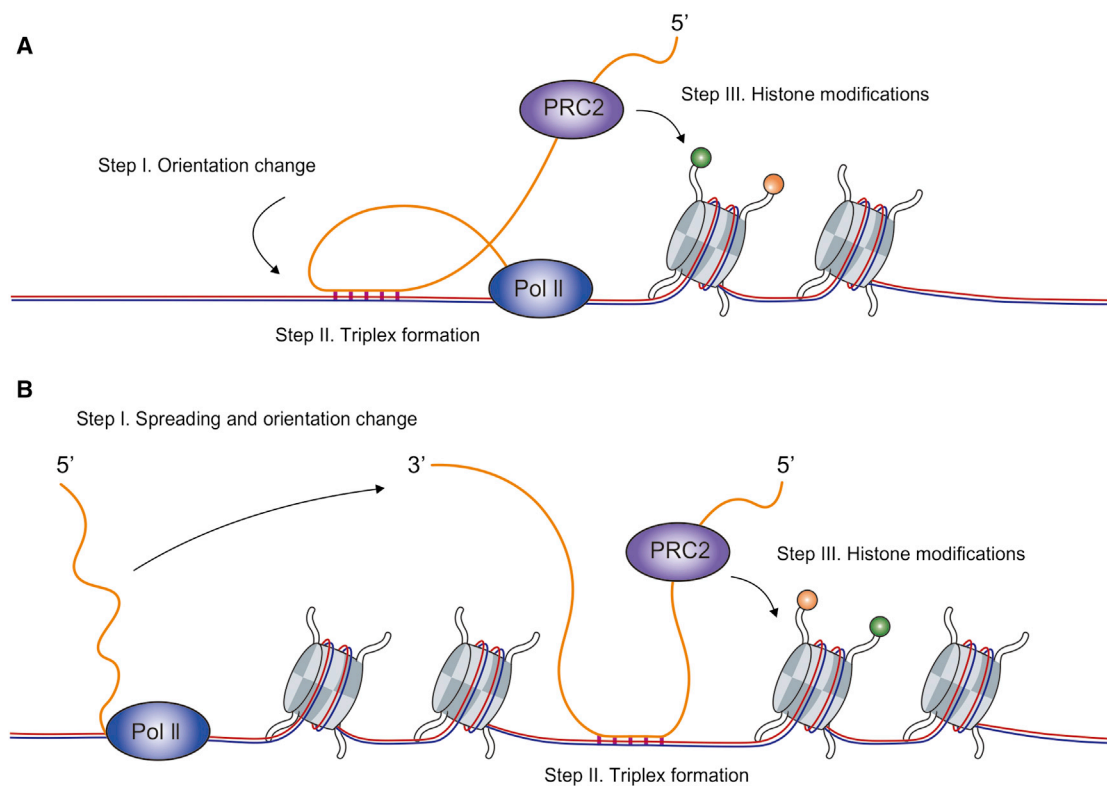


Figure 3. Orientation of Triplex Formation by lncRNAs

(A) If a transcribed lncRNA forms a triplex with a DNA sequence in the same region, it must change its orientation; in this case, the symmetrical triplex-forming motif is favorable.

(B) In the case of the *trans*-acting mechanism, a change in orientation is also needed for triplex formation. The lncRNA can form a triplex with the same symmetrical sequence or a reversed asymmetrical sequence.

With respect to the orientation of triplex formation, it is interesting that the sequence of the third strand is not the same, but rather, in the reverse orientation to target DNA sequences in both the parallel and antiparallel triplex structures (Figure 1B) (Yamagata et al., 2016). For example, in the parallel triplex, the direction of the UC motif in the RNA is 5' to 3', whereas the direction of the TC motif in the DNA is 3' to 5'. In the antiparallel triplex, the direction of the AG motif in the third-strand RNA is 3' to 5', whereas the direction of the AG motif in the DNA is 5' to 3'. This means that the third-strand RNA cannot sequence-specifically target double-stranded DNA (dsDNA) of the same sequence, and it can bind only if the sequence is in the reverse orientation. This interesting observation is based on the fact that the triplex-forming motif of lncRNA can only form a triplex with the corresponding DNA sequence when it is symmetrical, because the symmetrical sequence in the reverse orientation is exactly the same as the original sequence. Triplex-forming RNAs harboring such a symmetrical sequence were coined as palindrome RNA, which has the ability to form intermolecular triple helix structures with the DNA duplex at the transcription site (Buske et al., 2011).

The requirement for symmetrical motifs is important when considering *cis*-acting lncRNAs because they are transcribed and bind to DNA sequences in the same or neighboring regions (Figure 3A). The transcribed lncRNA with symmetrical triplex-forming motifs must change orientation and form a triplex struc-

ture with the corresponding DNA sequence. We propose that the change in orientation and symmetrical motifs are important for triplex formation by *cis*-acting lncRNAs. As shown in Table 1, the triplex-forming motifs of *cis*-acting lncRNAs are either fully or partially symmetrical. This supports our hypothesis of the need for symmetrical motifs for the formation of RNA-DNA triplex structures during gene regulation.

In the case of *trans*-acting lncRNAs, if the triplex-forming motif is symmetrical it will target dsDNA with the same sequence; if not, it will target dsDNA in the reverse direction (Figure 3B). Therefore, symmetrical motifs are also important for the identification of target sites of *trans*-acting lncRNAs. Based on the above discussion, we propose the model of the effects of symmetrical motifs on triplex formation by lncRNAs shown in Figure 3. RNA-DNA hybrid duplex formation has also been suggested as one of the potential mechanisms for lncRNA actions (Wang and Chang, 2011). The major difference between the formation of the RNA-DNA hybrid duplex and the RNA-DNA triplex is that, for the RNA-DNA hybrid duplex, the sequence and orientation are exactly the same as the corresponding DNA sequence.

Concluding Remarks

Despite the large numbers of lncRNAs reported, their structures have received little attention and their structure-function relationships remain unclear (Novikova et al., 2013). Many methods have been developed for the detection and determination of RNA 2D

Table 1. Examples of Triplex Formation by lncRNAs

lncRNA	Triplex-Forming Motifs (5' to 3')	<i>cis</i> versus <i>trans</i>	Associated Proteins
pRNA of <i>DHFR</i>	ACAAAUGGGGACGAGGGGGG CGGGGCGGCC	<i>cis</i> on <i>DHFR</i> gene	interact with TFIIB and dissociate the preinitiation complex (Martianov et al., 2007)
pRNA of <i>rRNA</i> gene	GUCGACCAGUUGUCCUUUG	<i>cis</i> on <i>rRNA</i> gene	recruit DNMT3b for DNA methylation (Schmitz et al., 2010)
Fendrr	UCCCCUCCAUCUCUCCUU CUCCUCCUCCUUCUUU	<i>cis</i> on <i>Foxf1</i> gene; <i>trans</i> on <i>Pitx2</i> gene	bind to PRC2 and TrxG/MLL (Grote et al., 2013)
Khps1	UCCCCUUUUUUUCCUCCU	<i>cis</i> on <i>SPHK1</i> gene	recruit p300/CBP to the <i>SPHK1</i> promoter (Postepska-Igielska et al., 2015)
PARTICLE	AAGGGGGGGGGAA	<i>cis</i> on <i>MAT2A</i> gene	interact with G9a and SUZ12 (subunit of PRC2) (O'Leary et al., 2015)
MEG3	CGGAGAGCAGAGAGGGAGCG	<i>trans</i> on TGF- β pathway genes	recruit PRC2 to target site (Mondal et al., 2015)
HOTAIR	AGAGGAGGGAAGAGAG	<i>trans</i> on <i>PCDH7</i> , <i>HOXB2</i> , etc.	bind to PRC2 and LSD1 (Kalwa et al., 2016)

and 3D structures, such as chemical or enzymatic probing, UV crosslinking, nuclear magnetic resonance (NMR), X-ray, and various software predictions (Mortimer et al., 2014; Zahran et al., 2015; Kubota et al., 2015; Magnus et al., 2014; Tijerina et al., 2007). However, when it comes to triplex formation by lncRNAs many important questions remain open. For example, given the potential structural complexity of lncRNAs, we are yet to understand how the RNA-DNA triplex forms and what regulates triplex formation. For example, if the triplex-forming motifs are located in the loop structure, is it possible for the loop sequence to bind directly to the DNA sequence? By contrast, if the triplex-forming motifs are involved in other secondary structures, such as hairpin structures, the structural equilibrium between RNA structures and RNA-DNA triplex formation must be considered. As shown in Table 1, some AG motifs are potentially G-rich sequences, suggesting possible competition between RNA G-quadruplex and triplex structures. Structural competition has been reported between RNA G-quadruplex and hairpin structures, as has the fact that different ions or small molecules can regulate these preferences (Bugaut et al., 2012).

Some proteins interact directly with RNA-DNA triplex structures or regulate RNA-DNA triplex structures. For example, it has been suggested that Argonaute proteins may stabilize the miRNA-DNA triplex structure (Toscano-Garibay and Aquino-Jarquin, 2014). Some helicases have also been found to unwind the triplex formation or remodel RNA structures. It has been suggested that the helicase domain of the chromatin remodeler ATRX can remodel the structure of the repeat A region of Xist RNA to facilitate PRC2 binding (Sarma et al., 2014). These studies suggested the possible structural regulation by proteins in the formation or unwinding of the RNA-DNA triplex structure. The fact that RNA binds to the major groove of duplex DNA also suggests competition between RNA-DNA triplexes and protein binding. Moreover, in the case of pRNAs mentioned above, the RNA-DNA triplex structure may recruit DNMT3b and methylate the promoter CpGs, which suggests functional interactions between the triplex and proteins (Schmitz et al., 2010).

Small molecules that interact selectively with the triplex structure may also regulate or target the triplex structures formed by

lncRNAs. For example, amino sugars such as neomycin are groove binders of triplex structure and other molecules with a planar structure such as quinoxaline derivatives are selective intercalators of triplex structures (Arya, 2011; Arya and Coffee, 2000; Arya et al., 2001a; Zain et al., 1999; Duval-Valentin et al., 1995; Escudé et al., 1998). Small ligands targeting miRNA triplex structures were also reported (Watkins et al., 2015; Nahar et al., 2015). These ligands might regulate the functions of lncRNAs by directly targeting the triplex structure. However, selectivity toward the RNA-DNA triplex is quite important for the design of small molecules as RNA-DNA hybrid duplex, RNA triplex, DNA triplex, and other hybrid triplex structures could also form in vivo. For example, neomycin was suggested to induce and stabilize the RNA-DNA triplex formation in vitro (Arya et al., 2001b). The design of selective ligands toward the RNA-DNA hybrid duplex may shed light on the development of RNA-DNA triplex-specific ligands (Shaw and Arya, 2008; Shaw et al., 2008). Overall, the emergence of triplex formation by lncRNAs may provide an important direction for the design of RNA-DNA triplex-targeted ligands.

Accumulating evidence shows that lncRNAs may also control chromatin structure, including nucleosome positioning and chromosome looping (Böhmendorfer and Wierzbicki, 2015). The examples of triplex formation included in Table 1 also suggest that triplex formation is often associated with the recruitment of histone-modifying proteins such as PRC2. Therefore, triplex formation at the chromatin level should be considered, especially for the *trans*-acting lncRNAs that target distant genes. Previous studies have suggested that nucleosome structures can affect or even inhibit triplex formation (Brown and Fox, 1996, 1998a, 1998b, 1999). Therefore, nucleosome and chromatin structures may affect RNA-DNA triplex formation in a manner similar to that of sequence-specific DNA binders such as polyamides (Dudouet et al., 2003; Suto et al., 2003; Gottesfeld et al., 2001; Han et al., 2014).

In summary, we have described the current evidence that supports the idea that triplexes of lncRNAs exist and play significant biological roles. We also discussed the orientation of triplex formation based on the *cis*- and *trans*-acting mechanisms, and

explored the possibility that proteins or small molecules can regulate triplex structures. We suggest that triplex formation represents a general mechanism for targeting DNA sequences by lncRNAs, and that triplexes may directly interact with functional proteins. However, the major concern is the need for direct evidence that supports triplex formation as the common mechanism followed by lncRNAs to target DNA sequencing particularly in gene regulation. Until now, the triplex-forming motifs of lncRNAs have been manually designated or predicted by software. The exact crystal or NMR structures of triplex formation in the context of lncRNA and its associated proteins are yet to be reported. Moreover, the design of selective RNA-DNA ligands that could regulate the functions of lncRNAs offers new opportunities for the development of drugs targeting lncRNAs. Given the increasing numbers of lncRNAs recently reported, one may anticipate that more examples of lncRNAs involved in triplex formation will be reported and these questions could be addressed in the near future.

AUTHOR CONTRIBUTIONS

Y.L. wrote the manuscript supervised by H.S. J.S. and H.S. revised the manuscript.

ACKNOWLEDGMENTS

We thank Professor Peter A. Beal from Department of Chemistry, University of California (Davis), for the critical reading and valuable comments on the manuscript. This work was supported by JSPS KAKENHI (grant number 24225005).

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