Allosteric regulation of U1 snRNP by splicing regulatory proteins controls spliceosomal assembly

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ABSTRACT
Alternative splicing is responsible for much of the transcriptomic and proteomic diversity observed in eukaryotes and involves combinatorial regulation by many cis-acting elements and trans-acting factors. SR and hnRNP splicing regulatory proteins often have opposing effects on splicing efficiency depending on where they bind the pre-mRNA relative to the splice site. Position-dependent splicing repression occurs at spliceosomal E-complex, suggesting that U1 snRNP binds but cannot facilitate higher order spliceosomal assembly. To test the hypothesis that the structure of U1 snRNA changes during activation or repression, we developed a method to structure-probe native U1 snRNP in enriched conformations that mimic activated or repressed spliceosomal E-complexes. While the core of U1 snRNA is highly structured, the 5′ end of U1 snRNA shows different SHAPE reactivities and psoralen crosslinking efficiencies depending on where splicing regulatory elements are located relative to the 5′ splice site. A motif within the 5′ splice site binding region of U1 snRNA is more reactive toward SHAPE electrophiles when repressors are bound, suggesting U1 snRNA is bound, but less base-paired. These observations demonstrate that splicing regulators modulate splice site selection allosterically.

Keywords: allosteric regulation; splice site selection; spliceosomal assembly; splicing regulatory proteins; U1 snRNP

INTRODUCTION
Pre-mRNA splicing is a fundamental RNA processing step that is common to all eukaryotes (Matera and Wang 2014). The combinatorial control that drives alternative pre-mRNA splicing can generate over 200,000 mRNA transcripts from roughly 20,000 human genes and accounts for much of the transcriptomic and proteomic diversity observed in humans and other higher eukaryotes (Hertel 2008; Nilsen and Graveley 2010; Park et al. 2018; Shenasa and Hertel 2019). There are many factors that influence alternative splice site usage which include the strength of splice sites, RNA secondary structure and the presence or absence of splicing regulatory elements (SREs) that recruit trans-acting splicing activators and repressors (Hertel and Maniatis 1998; Buratti and Baralle 2004; Shepard and Hertel 2008; Warf and Berglund 2010; McManus and Graveley 2011; Shepard et al. 2011; Erkelenz et al. 2013; Fu and Ares 2014). All of these features affect which splice sites are selected in any given transcript. There are also examples of each of these factors being the primary drivers of alternative splicing in different cellular and experimental contexts (Shenasa and Hertel 2019). The spliceosome is a large dynamic macromolecular machine that is composed of small nuclear ribonucleoprotein (snRNP) components, which assemble onto the pre-mRNA in a stepwise fashion, along with other protein factors (Wahl et al. 2009). U1 snRNP recognizes the pre-mRNA through base-pairing interactions between the 5′ end of U1 snRNA and the 9 nt consensus sequence that defines the 5′ splice site. This RNA-RNA base-pairing is the basis for 5′ splice site selection (Wang and Cooper 2007; Wahl et al. 2009; Matera and Wang 2014). The 3′ splice site is initially recognized by the heterodimer U2AF, which binds the polypyrimidine tract and contacts the AG dinucleotide at the 3′ splice site (Wu et al. 1999). Subsequent ATP-dependent rearrangements lead to higher order spliceosomal complex formation, intron excision and exon ligation (Moore et al. 1993; Wahl et al. 2009).

It is known that U1 snRNA participates in splice site pairing and can be the target of splicing regulators. For example, stem–loop IV of U1 snRNA interacts with the U2 snRNP component SF3A1 or the polypyrimidine tract-binding protein to modulate splicing (Sharma et al. 2011, 2014).
and mutagenesis of stem–loop IV causes exon skipping in vivo (Sharma et al. 2014). Furthermore, hnRNP L and A1 mediated splicing repression can promote the formation of extended U1 snRNA base-pairing with an exonic pre-mRNA sequence, which is predicted to block U6 snRNA base-pairing (Chiou et al. 2013).

Tissue specific alternative splicing often depends on SREs that recruit differentially expressed splicing regulatory proteins (Zahler et al. 1993; Kamma et al. 1995). These splicing regulators can be divided into two large families that include serine/arginine-rich proteins (SR proteins) and heterogeneous nuclear ribonucleoproteins (hnRNPs). Although SR proteins are best known for splicing activation through binding exonic splicing enhancers (ESEs) and hnRNPs have traditionally been associated with exonic splicing silencers (ESSs), both SR proteins and hnRNPs have been shown to act as activators and repressors in different splicing contexts (Wang and Cooper 2007; Wahl et al. 2009; Matera and Wang 2014). A systematic study showed that SR proteins and hnRNPs display a position-dependent activity at the 5′ splice site (Erkelenz et al. 2013). SR proteins activate splicing when bound upstream of the 5′ splice site, but repress splicing when bound intronic, downstream from the 5′ splice site. An opposite effect was observed for hnRNPs, which repress splicing when bound upstream of the 5′ splice site, but activate splicing when bound intronic, downstream from the 5′ splice site (Fig. 1A). The molecular mechanism for this position-dependent and antagonistic activity by SR proteins and hnRNPs has yet to be elucidated (Erkelenz et al. 2013).

Three related hypotheses can be proposed to explain position-dependent splicing regulation by SREs. It can be postulated that the binding of SR proteins and hnRNPs at activating or repressing positions leads to the recruitment of additional proteins that subsequently cause activation or repression. Alternatively, the interaction of splicing regulators at different positions relative to the 5′ splice site may change the binding affinity of U1 snRNP or U1snRNP-specific proteins with the 5′ splice site of the pre-mRNA. Lastly, it can be hypothesized that splicing regulators change the structure of U1 snRNA in such a way that U1 snRNP binding at the 5′ splice site can occur, but splice site selection or splice site pairing, which is associated with spliceosomal A-complex formation, cannot occur (Lim and Hertel 2004). This last hypothesis is the focus of the present study.

To test whether splicing regulators change the structure of U1 snRNA, we developed a U1 snRNP enrichment and SHAPE (selective 2′-hydroxyl acylation analyzed by primer extension) structure-probing approach (Wilkinson et al. 2006; Spitale et al. 2013). The method reliably identifies the binding of the U1 snRNP-specific proteins U1-70K and U1-A. We found the core of U1 snRNA (nucleotides 26–108) to be highly structured across all conditions tested. However, we discovered a motif within the 5′ end of U1 snRNA, referred to as the junction motif, which is predicted to base-pair with the intron–exon junction and is less reactive to SHAPE electrophiles when SRSF7 or TIA-1 bind the pre-mRNA from activating positions. These results suggest that activating splicing regulators promote U1 snRNA base-pairing with the 5′ splice site of the pre-mRNA, a proposal supported by psoralen crosslinking experiments. In contrast, splicing regulators bound at repressive positions prevent base-pairing with the 5′ splice site, thus providing a mechanistic framework for position-dependent splicing activation and repression.

RESULTS

U1 snRNA structure-probing design

We developed an approach to enrich and structure-probe native U1 snRNPs from HeLa cell nuclear extract. Substrate RNAs were derived from pre-mRNAs that were shown to undergo activated or repressed splicing (Erkelenz et al. 2013). Each RNA contained a functional 5′ splice site in addition to verified binding sites for SRSF7 or TIA-1, either upstream or downstream from the 5′ splice site (Table 1; Fig. 1B). The designed substrates were short, excluded a 3′ splice site and a downstream exon to ensure that the 5′ splice site was the only U1 snRNP binding site present and to stall spliceosomal assembly at E-complex without the need to deplete nuclear extract of ATP or use spliceosomal inhibitors (Erkelenz et al. 2013). To confirm that the pull-down approach enriches fully assembled U1 snRNP, western blot analysis was used to verify the presence of
U1-specific proteins U1-70K and U1-C (Fig. 2A). Additionally, enrichment of SRSF7 and TIA-1 binding on substrate RNAs that contain their respective binding sites was also verified (Table 1; Fig. 2B).

Comparison between in vitro transcribed U1 snRNA and nuclear extract derived U1 snRNA

To verify that the structure-probing protocol allows for the identification of U1 snRNA regions that interact with U1-specific proteins, we compared the SHAPE reactivity of in vitro transcribed U1 snRNA (in vitro U1 snRNA) with the reactivity of U1 snRNA present in nuclear extract (nuclear extract U1 snRNA) (Fig. 3A lane 1 vs. 3, B,C). Loop regions of stem–loop I and II, which are the known interaction sites of U1-70K and U1-A, should be more reactive for in vitro U1 snRNA when compared to nuclear extract U1 snRNA, which is bound by U1-specific proteins (Kondo et al. 2015). In agreement with these expectations, stem–loop I nucleotides U27–G38 of nuclear extract U1 snRNA show no SHAPE reactivity, whereas many of the same nucleotides of in vitro U1 snRNA display increased reactivity (Fig. 3A–C).

Based on a published high-resolution substructure of U1 snRNA bound by the RRMs of U1-70K and U1-A, the first 7 loop nucleotides of stem–loop II are known to be ordered by the RNA recognition motif (RRM) of U1-A (Kondo et al. 2015). In agreement with U1-A binding, we observe reduced SHAPE reactivity for nucleotides G68 and C69 when comparing nuclear extract U1 snRNA with in vitro transcribed U1 snRNA and increased reactivity for C73 and C74 as they are shown to be single stranded and not interacting with the RRM of U1-A (Fig. 3A–C; Kondo et al. 2015). Interestingly, U66 is the only nucleotide that is reactive in the first 7 loop nucleotides of stem–loop II in nuclear extract U1 snRNA. This observation is plausible when considering the crystal structure, because the 2′ hydroxyl of U66 participates in the coordination of a water molecule between the amide carbonyl oxygen of leucine 49 and N3 of G68, which may activate it toward SHAPE electrophiles (Kondo et al. 2015).

Nucleotides U60, U62, and C86 display slightly elevated SHAPE reactivity in nuclear extract U1 snRNA compared to in vitro U1 snRNA (Fig. 3B,C). Several considerations could provide an explanation for these observations. It is possible that the bulged nucleotide preceding U60 adopts a unique conformation that enhances the SHAPE reactivity of its immediately downstream nucleotide. Nucleotide C86 is predicted to be bulged out and it is possible that protein interactions stabilize a conformation that also enhances its SHAPE reactivity in nuclear extract. Nucleotides C86 and U60 are adjacent to strong SHAPE-independent stops in the nuclear extract lanes. Thus, it is also plausible that the quantitation of nucleotides C86 and U60 is artificially influenced by these SHAPE-independent stops.

Loop nucleotides of stem–loop III, which are predicted to be single stranded and unbound by proteins across all conditions, are highly reactive toward 2-methylnicotinic acid imidazolide (NAI). The results obtained by SHAPE structure-probing are consistent with the crystallographic data for protein-associated U1 snRNA and they demonstrate that the use of SHAPE electrophiles, such as NAI (Spitale et al. 2013), can be used in conjunction with our pull-down approach to detect changes in the physical state of U1 snRNA.

### TABLE 1. Substrate RNAs used in pull-down experiments

| Substrate | 5’ ss sequence | SRE sequence | Length (nt) |
|-----------|----------------|--------------|-------------|
| N         | CUG/GUGAGU     | CCAAAACA (3×) | 107         |
| SR U      | CUG/GUGAGU     | AGACAACGAUUGAUCAU (2×) | 94         |
| SR D      | CUG/GUGAGU     | AGACAACGAUUGAUCAU (2×) | 118        |
| TIA U     | CUG/GUGAGU     | UCUUUUUAAGUCGUACCUEU (2×) | 94         |
| TIA D     | CUG/GUGAGU     | UCUUUUUAAGUCGUACCUEU (2×) | 118        |

### FIGURE 2. Confirmation of U1 snRNP assembly and splicing regulatory protein enrichment on substrate RNAs. (A) U1 snRNPs bound to substrate RNAs contain both U1-70K and U1-C; hnRNP A1 was used as a loading control. Beads sample contains no RNA and β-globin RNA was used as a positive control. (B) SRSF7 and TIA-1 are enriched on substrate RNAs containing their respective binding sites.
Comparison between nuclear extract U1 snRNA versus enriched U1 snRNA

To determine whether the interaction of U1 snRNP with a 5’ splice site changes the structure or dynamics of U1 snRNA, we compared the SHAPE reactivity of enriched U1 snRNAs, defined as those bound to substrate RNAs, to nuclear extract U1 snRNA, defined as those unbound to RNA. The majority of U1 snRNPs in nuclear extract are assumed to be in an unbound state, with a minority of U1 snRNPs bound to genuine 5’ splice sites and other noncanonical sequences. The SHAPE analysis of nucleotides 26–108 showed minimal differences between nuclear extract U1 snRNA and enriched U1 snRNAs, suggestive of minimal changes in the protein bound core of U1 snRNA upon 5’ splice site binding (Fig. 4: compare lanes 5, 7, 9, 11, and 13 to lane 17, Fig. 5 nuclear extract compared to enriched conditions).

As stated above, we observe that nucleotides U60, U62, and C86 are reactive in nuclear extract U1 snRNA. Interestingly, however, these nucleotides do not display SHAPE reactivity in U1 snRNA bound to the 5’ splice site sequence (Fig. 5, compare nuclear extract to enriched conditions). The loss of SHAPE reactivity for U60, U62, and C86 may therefore be an indication of 5’ splice site binding. Despite these small differences in SHAPE reactivity upon 5’ splice site binding, the overall lack of differences is largely consistent with expectations given that U1 snRNA is a highly structured RNA that forms its secondary structure even in the absence of U1 specific proteins. Therefore, 5’ splice site binding is not expected to significantly change the overall structure of U1 snRNA. We conclude that the core of U1 snRNA maintains a near uniform SHAPE reactivity regardless of its interactions with pre-mRNAs.

We also compared the 5’ splice site binding region of U1 snRNA, defined as nucleotides 2–11, between nuclear extract U1 snRNA and enriched U1 snRNAs. We identified a 6 nt junction motif (5’-UACCUG-3’), which is more reactive to NAI in nuclear extract U1 snRNA compared to U1 snRNAs bound adjacent to activators (Fig. 6). This result shows that the SHAPE reactivity of the junction motif correlates with enhanced 5’ splice site binding.
Impact of splicing activation and repression on SHAPE reactivity of U1 snRNA

A detailed analysis of the SHAPE reactivities of U1 snRNA bound to different substrate RNAs (activated versus repressed) was performed (Fig. 4, compare lanes 5, 7, 9, 11, and 13; Fig. 5, compare between enriched conditions). The core of U1 snRNA (nucleotides 26–108) has a uniform SHAPE reactivity regardless of whether it is in an activated or repressed state, suggesting that U1 snRNA structural changes that promote spliceosomal assembly are not medi-
ated through this region. However, the junction motif is consistently observed to be less reactive toward NAI when splicing regulators are bound at activating positions (Fig. 7A–C). This result suggests that SRSF7 and TIA-1 promote more rigid base-pairing between U1 snRNA and the 5′ splice site when bound at activating positions (Fig. 7A–C; Supplemental Table 1). The increased SHAPE reactivity of the junction motif when splicing regulatory proteins are bound at repressive positions suggests that repressor/U1 snRNP interactions interfere with efficient base-pairing between the 5′ splice site and U1 snRNA. Interestingly, the average SHAPE reactivity of U1 snRNA bound to a neutral control substrate RNA lies between those observed for the activated and repressed substrates, suggesting that neutral sequences adopt an intermediate base-pairing state (Fig. 7C).

To determine whether the observed differences in SHAPE reactivity are due to a lack of base-pairing between U1 snRNA and the 5′ splice site in repressed conditions or differences in base-pairing efficiency, we carried out psoralen crosslinking experiments of U1 snRNPs to activated or repressed substrates (Fig. 8A, B). The results show that U1 snRNA base-pairs with the 5′ splice site in all substrate configurations, however at varying levels of crosslinking efficiency. Activated substrate RNAs crosslink to U1 snRNA more efficiently than repressed or the neutral substrate RNAs (Fig. 8A, compare lanes 3 and 9 to lanes 1, 5 and 7, Fig. 8B; Supplemental Table 2). One notable anomaly is the increased crosslinking efficiency of the SRSF7 bound downstream (SR D) substrate relative to the neutral substrate (N). Overall, the crosslinking results converge with the SHAPE structure-probing data, providing additional evidence that activators and repressors modulate the efficiency of 5′ splice site base-pairing interactions. U1 snRNP binds the 5′ splice site in various contexts, but splicing activators increase base-pairing, whereas strong splicing repressors such as TIA-U decrease base-pairing.

DISCUSSION

SR proteins and hnRNP have dual and antagonistic activities depending on where they bind relative to a regulated 5′ splice site. During splicing repression, U1 snRNP is recruited to the pre-mRNA and forms E-complex, however the formation of higher order complexes is severely hindered (Erkelenz et al. 2013). U1 snRNP is the only stably associated snRNP at E-complex (Hong et al. 1997; Das et al. 2000). Here, we carried out a thorough investigation of the U1 snRNA secondary structure in different contexts of splicing regulation. Importantly, we show that the junction motif of U1 snRNA is more reactive for U1 snRNPs not engaged in RNA binding and repressed U1 snRNPs when compared to activated U1 snRNPs bound to RNA (Fig. 6). While this may seem intuitive, it highlights two important points. First, the enriched SHAPE method is sensitive
enough to detect changes in base-pairing efficiency, such as splice site activation by splicing regulatory proteins. Secondly, the method used herein can be a useful tool for researchers probing dynamic RNAs or RNPs. While the use of cell permeable structure probes such as DMS, NAI and NAz is extremely useful and permits RNA structure-probing in native environments, RNPs such as the spliceosome are known to be highly dynamic (Spitale et al. 2013; Rouskin et al. 2014; Zubradt et al. 2017; Feng et al. 2018). In specific instances it may be essential to probe an RNP in one unique context. For example, this study focuses on U1 snRNP’s functional role in the context of inhibiting premature polyadenylation (Gunderson et al. 1998; Kaida et al. 2010). SHAPE carried out on whole cells would not be able to distinguish between U1 snRNAs that are bound to canonical 5’ splice sites, those that are preventing pre-mature polyadenylation or those that are not bound to RNA. It would also be unable to distinguish between U1 snRNPs in activated versus repressed states. As a result, an average SHAPE reactivity would be collected for in cell SHAPE. Our findings also add to previous U1 snRNP structure-probing studies, which include chemical and enzymatic probing of deproteinized U1 snRNA, hydroxyl radical probing of U1 snRNP in E and A complex and SHAPE chemistry performed on in vitro transcribed U1 snRNA as a validation method for the SHAPE probe FAI-N3 (Krol et al. 1990; Dönmez et al. 2007; Chan et al. 2018). The structure-probing performed with FAI-N3 is largely consistent with our SHAPE data. However, because FAI-N3 can undergo a click chemistry-mediated biotin-streptavidin enrichment step, it has a higher signal to noise ratio and shows all nucleotides within loop regions to be reactive toward SHAPE electrophiles (Chan et al. 2018). The hydroxyl radical probing data shows that U1 snRNA does not undergo any major structural rearrangements that lead to differences in solvent accessibility, from E to A complex (Dönmez et al. 2007). We also observe that the core of U1 snRNP maintains a near uniform SHAPE reactivity when not associated with RNA or when bound to different substrate RNAs that mimic activated or repressed states. Both of these observations suggest that U1 snRNP maintains a uniform structure and protein composition in the early spliceosome.

The most striking differences in SHAPE reactivity that correlate with splicing activation and repression were detected when we compared SHAPE profiles of the U1 snRNA 5’ splice site binding region, in particular the intron–exon spanning junction motif (Fig. 7A–C). These results provide evidence that splicing activation promotes base-pairing and strong splicing repression, such as the proven repressor TIA-1 bound exonically, interferes with base-pairing between the junction motif and the 5’ splice site pre-mRNA.

FIGURE 5. Reactivity plots of enriched U1 snRNAs and nuclear extract U1 snRNA. Black bars (0–0.3) signify background reactivity, orange bars (0.3–0.7) signify intermediate reactivity, and red bars (≥0.7) indicate highly reactive nucleotides. The y-axis represents the normalized SHAPE reactivity and the x-axis indicates the nucleotide position from 5’ to 3’ in the U1 snRNA sequence. Loop nucleotides for stem–loops I and II are indicated in addition to nucleotides that display different SHAPE reactivities between enriched conditions and the NE condition. The reactive nucleotides for stem–loop III are also indicated.
nucleotides. This proposal is supported by psoralen crosslinking (Fig. 8A,B) and consistent with functional splicing data for these RNAs (Erkelenz et al. 2013).

It is plausible that the conformation of the junction motif acts as a switch where increased base-pairing with the 5′ splice site promotes splice site pairing, while U1 snRNAs with less base-paired junction motifs cannot promote splice site pairing. The fact that the junction motif shows increased SHAPE reactivity and decreased psoralen crosslinking when both SRSF7 and TIA-1 are bound at repressive positions further suggests that its base-pairing state can be modulated by inputs from both faces of U1 snRNP (Fig. 1A).

The “bind and activate” mode of 5′ splice site selection could be a safeguard mechanism to differentiate between U1 snRNPs functional roles in spliceosomal recruitment and prevention of premature polyadenylation. It is possible that U1 snRNPs binds the pre-mRNA in an inactive conformation that is allosterically activated by splicing regulators bound at activating positions. Such a mechanism may be necessary because U1 snRNA is known to not only interact with genuine 5′ splice sites, but also with cryptic 5′ splice sites that frequently inhibit premature polyadenylation (Kaida et al. 2010). Thus, enhanced 5′ splice site base-pairing with U1 snRNP may act as an allosteric switch for spliceosomal assembly. U1 snRNP binding to a 5′ splice site triggers E-complex formation regardless of where splicing regulatory proteins are bound (Erkelenz et al. 2013). However, locking in base-pairing between U1 snRNA and the 5′ splice site occurs only in the presence of splicing activators. This activated conformation may then allow U1 snRNP to engage in cross intron activities, which include U1 snRNA stem–loop IV-mediated contacts with the U2 snRNP component SF3A1 during A complex formation (Sharma et al. 2014).

The proposed allosteric mechanism for the initiation of splice site pairing does not rule out any of the other potential mechanisms of position-dependent splicing activation/repression. The conformation of U1 snRNA could be regulated through U1- specific proteins such as U1-C, which is known to enhance 5′ splice site binding (Kondo et al. 2015). It is possible that TIA-1, which has been demonstrated to contact U1-C and recruit U1 snRNP to the 5′ splice site when bound intronically, can disrupt U1 snRNP’s structural integrity when bound exonically (Förch et al. 2002). A similar mode of action could be envisioned for SRSF7, which is likely to mediate its effect on the junction motif SHAPE reactivity of U1 snRNA junction motif (U6 to G11) upon RNA binding. The SHAPE reactivity of U1 snRNA enriched on substrate SR U is compared with the SHAPE reactivity of nuclear extract U1 snRNA (NE), which is not bound to substrate RNA. The y-axis represents the normalized SHAPE reactivity and the x-axis indicates the nucleotide position from 5′ to 3′ in the U1 snRNA sequence.

![FIGURE 6. SHAPE reactivity of U1 snRNA junction motif (U6 to G11) upon RNA binding. The SHAPE reactivity of U1 snRNA enriched on substrate SR U is compared with the SHAPE reactivity of nuclear extract U1 snRNA (NE), which is not bound to substrate RNA. The y-axis represents the normalized SHAPE reactivity and the x-axis indicates the nucleotide position from 5′ to 3′ in the U1 snRNA sequence.](image)

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through U1-70K. Lastly, the recruitment of additional proteins cannot be ruled out. It is possible that splicing regulators recruit different proteins to the two different faces of U1 snRNP depending on where they bind the pre-mRNA.

We have developed a targeted SHAPE methodology, which allows for probing U1 snRNP in the context of position-dependent splicing activation or repression. This method can be adapted to probe other snRNPs/spliceosomal complexes and possibly other dynamic RNPs. Importantly, we have demonstrated that the SHAPE reactivity of the U1 snRNA junction motif is altered during splicing activation and repression. These results provide molecular insights into the mechanism of position-dependent splicing activation and repression.

MATERIALS AND METHODS

In vitro transcription of substrate RNAs

Sequences for substrate RNAs were simultaneously linearized and amplified using standard PCR in 50 µL reactions. Subsequently 40 µL of PCR product was transcribed in vitro using the RiboMax RNA transcription kit (Promega). DNase treatment was performed according to the RiboMax protocol and RNAs were purified with the RNA Clean and Concentrator Kit (Zymo Research). A Nanodrop 2000 instrument was used to obtain RNA concentration values.

U1 snRNP enrichment approach

U1 snRNP was enriched through a pull-down assay using various substrate RNAs that contained verified binding sites for SRSF7 or TIA-1 at activating or repressing positions (Fig. 1B). Enriched U1 snRNPs within the context of activated or repressed E-complex were then probed using NAI, a SHAPE reagent (Spitale et al. 2013). SHAPE electrophiles acylate the 2′OH of RNA nucleotides based on their flexibility and dynamics (Wilkinson et al. 2006; McGinnis et al. 2012). Unstructured regions of RNA are more likely to be modified by SHAPE reagents, which cause reverse transcription to terminate one nucleotide before the acylated nucleotide. To detect structural changes within U1 snRNA including stem-loop IV, an LNA modified primer was designed to bind the 3′ end of U1 snRNA stem-loop IV. SHAPE reactivities of stem-loop IV were poor due to structure stops. Despite the poor SHAPE reactivities obtained for stem-loop IV, the LNA primer provided excellent reverse transcription (RT) signal and high-resolution data for the rest of U1 snRNA. Dideoxy sequencing was performed on enriched U1 snRNAs and nuclear extract U1 snRNA to confirm that the reverse transcription signal obtained from RNAs was generated from U1 snRNA (Supplemental Fig. 1).

Oxidation of substrate RNAs and RNA affinity pull-down

An amount of 300 picomoles of each substrate RNA was incubated with 5 mM NaOAc (sodium [meta]periodate) and 0.1 M NaOAc (sodium acetate) for 1 h in the dark. Substrate RNAs were subsequently precipitated with 100% ethanol (EtOH). Adipic acid dihydrazide agarose beads were prepared by washing 125 µL of beads per reaction four times with two volumes of 0.1 M NaOAc. After the final wash, the bead volume was adjusted to 250 µL beads per reaction and 250 µL of beads were added to each substrate RNA pellet. Beads were subsequently incubated at 4°C on a rotator overnight. Beads were washed two times with 1 mL 2 M NaCl and three times with Buffer D (20 mM HEPES-KOH [pH 7.6], 20%, glycerol, 0.1 M KCl, 0.2 mM EDTA, and 0.5 mM DTT). 200 µL of 30% nuclear extract diluted with Buffer D containing 1 M MgCl₂ was added to each sample and incubated on a rotator for 15 min at 30°C. All samples were subsequently washed five times with 1 mL Buffer D containing 1 M MgCl₂.

RNA acylation and isolation

After the final wash, 5 µL of 1 M NAI or DMSO was added to 45 µL of evenly mixed beads and samples were incubated at 30°C for 15 min. After incubation with NAI, 150 µL of H₂O was added followed by the addition of 200 µL of formamide. Samples were mixed by pipetting and then heated at 95°C for 1 min. RNA was separated from beads using 0.45 µm cellulose acetate Corning Costar SpinX columns, spun at 6000 rpm for 8 min. Final volumes were adjusted to 600 µL with H₂O and subsequently 600 µL of premixed phenol:chloroform isopropanol alcohol (25:24:1) pH 6.6 (Ambion) was added. The aqueous phase was extracted and washed with chloroform. For nuclear extract U1 snRNA, 5 µL of 1 M NAI or DMSO was added to 45 µL of 30% nuclear extract and incubated for 15 min at 30°C. Samples were directly phenol:chloroform extracted as described above. For in vitro U1 snRNA, U1 snRNA was folded and probed as described by Spitale et al. (2013). In vitro U1 snRNA was phenol:chloroform extracted as described above. All samples were precipitated with 300 mM NaOAc, 3 volumes of 100% EtOH and 2 µL of GlycoBlue.
(Invitrogen). All samples were washed twice with 70% EtOH and resuspended in 5 µL of water.

Reverse transcription

Four picomoles of a gene specific LNA modified primer were added to each reaction. Primers were 5’ end phosphorylated with T4 polynucleotide kinase (New England Biolabs [NEB]) using standard procedures from NEB and yielded 9000–10,000 CPM/µL. To anneal the primer, samples were heated to 95°C for 2 min and then cooled to 50°C. A master mix containing dNTPs, DTT, water (to 20 µL), 5× FS Buffer (250 mM Tris-HCl [pH 8.3], 375 mM KCl, 15 mM MgCl$_2$) and Superscript III Reverse Transcriptase (Invitrogen) was added according to the superscript III protocol. For dideoxy (dd) sequencing samples, 2 µL of 10 mM ddNTPs (TriLink Biotechnologies) were added to the RNA and primer mixture before annealing. All samples were reverse transcribed for 50 min at 50°C. The RNA template was hydrolyzed with addition of 2 µL (one tenth volume) of 4 M NaOH and incubation at 95°C for 5 min. Samples were ethanol precipitated as previously described and washed two times as previously described. Samples were resuspended in 95% formamide, 18 mM EDTA, 0.025% xylene cyanol and bromophenol blue.

Gel electrophoresis, visualization, and quantification

Samples were run on 10% polyacrylamide (19:1 acrylamide:bisacrylamide) 7 M urea gels, see reference for detailed gel casting protocol (Movassat et al. 2017). Gels were prerun at 20 watts for 1 h, followed by 50 watts for 4–6 h depending on the desired resolution. Gels were dried for 1 h under vacuum at 80°C and exposed to a phosphor screen overnight. Images were obtained using GE Typhoon imager and peaks were integrated using SAFA software (Das et al. 2005). Results from junction motif analysis were independently verified using ImageJ software (Schneider et al. 2012). For the reactivity plots, the counts at each position in the NAI lane were subtracted by counts in the corresponding DMSO control lane. The RNA samples were divided by the top 92%–98% most reactive nucleotides within that lane to obtain reactivity plots. The value obtained for each nucleotide was averaged across three biological replicates. For the fold change of the junction motif (Fig. 7C) the SHAPE reactivity of the junction motif was normalized to the SHAPE reactivity of nucleotide U22, which stays constant across all conditions. Normalized SHAPE reactivities were divided by the TIA D SHAPE reactivity to obtain a fold change. Fold changes were averaged across three biological replicates. One tailed t-tests were performed in a pairwise manner to determine statistical significance. Comparisons to the crystal structure reference PDB code=4PKD. Each experiment was performed in triplicate.

Western blot assay

Protein samples from RNA pull-down experiments were separated on precast 12% Tris-Glycine SDS-PAGE gels (Bio-Rad) at 125 V for 50–60 min at room temperature. Gels were transferred to methanol activated Immun-Blot PVDF membrane (Bio-Rad) using Tris-Glycine transfer buffer at 60 V, 2 h, 4°C, with an ice pack and stirring of transfer buffer. The membrane was blocked for 1 h, at room temperature with 5% nonfat milk in tris-buffered saline with Tween-20 (TBS-T), followed by an overnight incubation with primary antibody at 4°C. Primary antibody was diluted in 2.5% nonfat milk in TBS-T. Primary antibody conditions in this work were the following: U1-70K at 1:100 (Santa Cruz Biotechnology, clone C-18 or Millipore mouse monoclonal, clone 9C4.1), U1-C at 1:200 (Sigma, rat monoclonal, clone 4H12), hhnRNP A1 at 1:250 (Santa Cruz Biotechnology, mouse monoclonal, clone 9H10), SRSF7 at 1:1000 (MBL, rabbit polyclonal), and TIA-1 at 1:100 (Santa Cruz Biotechnology, goat polyclonal, clone C-20). The membranes was washed with TBS-T prior to incubation with the secondary antibody. Secondary antibodies were diluted in 2.5% nonfat milk in TBS-T and incubated for 1 h at room temperature. The membrane was washed further with TBS-T and then developed for 1 min using SuperSignal West Pico Chemiluminescent Substrate (ThermoFisher Scientific). Blots were imaged on a GE Typhoon imaging station.

Psoralen crosslinking

An amount of 300 picomoles of the five different substrate RNAs were dephosphorylated using NEB Quick Dephosphorylation Kit in 20 µL reactions. Reactions contained 20 units NEB quick CIP (calf intestinal phosphatase), 1× NEB CutSmart Buffer (50 mM postassium acetate, 20 mM Tris-acetate, 10 mM Magnesium acetate, 100 µg/mL BSA, pH 7.9) and were incubated at 37°C for 2 h. The CIP enzyme was heat inactivated at 80°C for 5 min. Substrate RNAs were 5’ end labeled with the addition of 20 units NEB T4 PNK (polynucleotide kinase), 2 µL gamma-ATP, and DTT (final concentration 5 mM). Kinase reactions were incubated at 37°C for 3 h and heat inactivated at 65°C for 20 min. Substrate RNAs were purified using ZYMO RNA Clean and Concentrator kits, resuspended in 25 µL and yielded a specific activity of ~600,000 cpm/µL. Crosslinking reactions were carried out using 1 µL of radiolabeled substrate RNA, 30% nuclear extract and monitored the conditions used in SHAPE experiments. Each reaction contained 20 µg/mL 4’-aminomethyltrioxsalen (AMT) and was exposed to 365 nm UV for 10 min or kept in the dark as described by Tan and Steitz (1994). Samples were then digested with proteinase K, phenol chloroform extracted and resolved on 6% denaturing polyacrylamide gels as described previously (Movassat et al. 2017). Crosslinking efficiencies were obtained using ImageJ. Briefly, a box was drawn around each band. The intensity in the +UV lane was subtracted by the intensity in the −UV lane and the resulting value was divided by the intensity of the band corresponding to the full-length substrate RNA in the +UV lane. The crosslinking efficiencies were averaged across three biological replicates and two tailed t-tests were performed in a pairwise manner to determine statistical significance.

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.
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