Dual nature of pseudouridylation in U2 snRNA: Pus1p-dependent and Pus1p-independent activities in yeasts and higher eukaryotes

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ABSTRACT

The pseudouridine at position 43 in vertebrate U2 snRNA is one of the most conserved post-transcriptional modifications of spliceosomal snRNAs; the equivalent position is pseudouridylated in U2 snRNAs in different phyla including fungi, insects, and worms. Pseudouridine synthase Pus1p acts alone on U2 snRNA to form this pseudouridine in yeast Saccharomyces cerevisiae and mouse. Furthermore, in S. cerevisiae, Pus1p is the only pseudouridine synthase for this position. Using an in vivo yeast cell system, we tested enzymatic activity of Pus1p from the fission yeast Schizosaccharomyces pombe, the worm Caenorhabditis elegans, the fruit fly Drosophila melanogaster, and the frog Xenopus tropicalis. We demonstrated that Pus1p from C. elegans has no enzymatic activity on U2 snRNA when expressed in yeast cells, whereas in similar experiments, position 44 in yeast U2 snRNA (equivalent to position 43 in vertebrates) is a genuine substrate for Pus1p from S. cerevisiae, S. pombe, Drosophila, Xenopus, and mouse. However, when we analyzed U2 snRNAs from Pus1 knockout mice and the pus1Δ S. pombe strain, we could not detect any changes in their modification patterns when compared to wild-type U2 snRNAs. In S. pombe, we found a novel box H/ACA RNA encoded downstream from the RPC10 gene and experimentally verified its guide RNA activity for positioning Ψ43 and Ψ44 in U2 snRNA. In vertebrates, we showed that SCARNA8 (also known as U92 scaRNA) is a guide for U2-Ψ43 in addition to its previously established targets U2-Ψ34/Ψ44.

Keywords: modification guide RNA; pseudouridine; Pus1p; U2 snRNA

INTRODUCTION

The isomerization of uridine to pseudouridine is the most abundant post-transcriptional modification found in stable RNA molecules. When pseudouridine was discovered in early RNA studies, it was even designated as a “fifth” nucleoside (Cohn and Volkin 1951; Davis and Allen 1957). First identified in abundant RNAs, such as rRNAs, tRNAs, and U snRNAs, pseudouridine has since been found in almost all types of RNAs, including mRNAs, snoRNAs, long noncoding RNAs, and telomerase RNA (Zhao et al. 2007; Carlile et al. 2014; Lovejoy et al. 2014; Schwartz et al. 2014). Pseudouridine synthases are enzymes that catalyze pseudouridine formation. Based on sequence similarities, pseudouridine synthases have been classified into six distinct families (Mueller and Ferré-D’Amaré 2009; Spenkuch et al. 2014). Pseudouridine synthases act either alone or with a guide RNA, known as a box H/ACA snoRNA. In the latter case, target recognition relies on specific base-pairing between the substrate and the guide RNA. Thus, a single pseudouridine synthase, Cbf5/NAP57/dyskerin, functions at different positions and in different RNAs (for review, see Yu and Meier 2014).

Eukaryotic spliceosomal snRNAs are extensively modified (Karijolich and Yu 2010). For instance, numerous pseudouridines are found in all vertebrate snRNAs and at least 14 pseudouridines were detected in human U2 snRNA alone (Deryusheva et al. 2012). Most snRNA modifications in higher eukaryotes depend on guide RNAs (Karijolich and Yu 2010; Deryusheva and Gall 2013; Adachi and Yu 2014). However, in budding yeast (Saccharomyces cerevisiae), snRNAs contain only nine pseudouridines: In addition to six constitutive pseudouridines, two in U1 snRNA, three in U2 snRNA, and one in U5 snRNAs (Massenet et al. 1999), there are three inducible pseudouridines, two in U2 snRNA (Wu et al. 2011) and one in U6 snRNA (Basak and Query 2014). Modification mechanisms have been identified for all five pseudouridines in U2 snRNA, both constitutive (Massenet et al. 1999; Ma et al. 2003, 2005) and inducible (Wu et al. 2011).
2011), and for the inducible pseudouridine in U6 snRNA (Basak and Query 2014). Four of these six positions are modified by two stand-alone pseudouridine synthases, Pus1p and Pus7p.

Pus1p, a member of the TruA pseudouridine synthase family, acts on several types of RNA substrates. Besides U2 and U6 snRNAs in S. cerevisiae (Massenet et al. 1999; Basak and Query 2014), it modifies uridines in many different tRNAs (Motorin et al. 1998; Chen and Patton 1999; Hellmuth et al. 2000; Patton and Padgett 2003; Patton et al. 2005; Behm-Ansmant et al. 2006), in the mouse steroid receptor RNA activator (Zhao et al. 2007), and in a set of mRNAs (Carlile et al. 2014; Lovejoy et al. 2014; Schwartz et al. 2014). In humans, mutations in the PUS1 gene are associated with mitochondrial myopathy and sideroblastic anemia (MLASA). Although Pus1p is a site-specific enzyme, it lacks strict sequence recognition requirements (Sibert and Patton 2012). Pus1p’s from different species, although highly conserved structurally, differ in their ability to catalyze pseudouridylation of certain conserved positions in tRNAs (Behm-Ansmant et al. 2006). As for U2 snRNA, in S. cerevisiae Pus1p is the only pseudouridylation machinery that modifies position 44 at the branch point recognition region; in vertebrates the equivalent position is 43, and this pseudouridine is one of the most conserved pseudouridines across all eukaryotic species. Whereas mouse Pus1p can pseudouridylate U2 snRNA when expressed in the pus1Δ yeast strain (Behm-Ansmant et al. 2006), the activity of Pus1p from other species is rather ambiguous (Hellmuth et al. 2000; Patton and Padgett 2003). Using an in vivo yeast cell system, we test here Pus1p from different phyla: a mammal (mouse), an amphibian (Xenopus tropicalis), an insect (Drosophila melanogaster), a worm (Caenorhabditis elegans), and two fungi (S. cerevisiae and Schizosaccharomyces pombe). We show species-specific differences in the ability of Pus1p to modify U2 snRNA. For instance, C. elegans Pus1p appears to lack activity on U2 snRNA when expressed in yeast cells. At the same time, our data demonstrate redundant Pus1p-dependent and guide RNA-mediated pseudouridylation of U2 snRNA at position 43 in vertebrates and in S. pombe.

RESULTS AND DISCUSSION

Genes encoding Pus1p have been annotated in many species. Based on data from yeast S. cerevisiae and mouse (Massenet et al. 1999; Behm-Ansmant et al. 2006), Pus1p is generally assumed to modify U2 snRNA, yet Pus1p from other species has never been tested directly for enzymatic activity on U2 snRNA. In fact, C. elegans U2 snRNA has not been considered a cePus1p substrate, because in mutant worms, the absence of Pus1p activity on tRNAs had no effect on U2 snRNA modification (Patton and Padgett 2003). Based on complementation analysis and sequence similarity, PUS1 has been earlier identified in S. pombe (Hellmuth et al. 2000); more recently, the pus1Δ strain has become available from the S. pombe gene deletion collection (Kim et al. 2010). Pus1p is the only pseudouridine synthase in S. cerevisiae that is active on U2 snRNA at position 44 (Fig. 1A). However, the pus1Δ strain of S. pombe shows a U2 snRNA modification pattern identical to its parental ED666 “wild-type” strain (Fig. 1B). It is worth noting here that we verified the PUS1 deletion by PCR and sequencing the amplified fragments. These observations suggest that U2 snRNA can be modified in C. elegans and S. pombe by a Pus1p-independent mechanism.

Pus1 knockout mice have been recently generated. In these mice, Pus1p-dependent modification of tRNAs was shown to
be missing, but U2 snRNA was not tested (Mangum et al. 2016). We analyzed U2 snRNA pseudouridylation in the Pus1 knockout mice and their wild-type siblings using RNA samples kindly provided by Jeffrey Patton and Diego Altomare (University of South Carolina). Again we did not detect any differences between wild-type and mutant mice in their U2 snRNA modification patterns (Fig. 1C). While spPus1p and cPus1p enzymatic activity on U2 snRNA has not yet been verified (Hellmuth et al. 2000; Patton and Padgett 2003), U2 snRNA was identified as a genuine substrate for mpus1p (Behm-Ansmant et al. 2006).

We decided to test the enzymatic activity of Pus1p from S. pombe and several higher eukaryote species in the pus1Δ S. cerevisiae strain. We chose X. tropicalis, D. melanogaster, and C. elegans Pus1p. Mouse and S. cerevisiae Pus1p served as positive controls in these experiments. The activity on U2 snRNA was evident for Pus1p from all examined species, except for C. elegans (Fig. 1A). However, in contrast to yeast Pus1p, mouse, Xenopus, and Drosophila enzymes were less efficient. To restore Ψ44 in U2 snRNA, they needed to be overexpressed, whereas both spPus1p and spPus1p could rescue pseudouridylation of this position in the pus1Δ S. cerevisiae strain even when expressed at a low level from a leaky Gal5 promoter. Expression levels of exogenous Pus1p were verified by Western blots using HA-tagged versions of the Pus1p expression constructs; beforehand, we tested that the HA-tag did not affect Pus1p enzymatic activity.

Intriguingly, the predicted PUS1 gene in Drosophila, CG4159, produces two alternatively spliced variants, of which only the shorter isoform, CG4159-PA, can modify U2 snRNA (Fig. 1A). The difference between the two isoforms of Drosophila Pus1p is small—a short 32 aa extension at the N terminus. Nevertheless, the web server IUPred (http://iupred.enzim.hu), which predicts intrinsically unstructured regions (Dosztányi et al. 2005), suggests that the N terminus of the longer isoform no longer forms a typical disordered tail (Czudnochowski et al. 2013), but instead is incorporated in a large globular domain. In mammals, short and long isoforms of Pus1p have also been reported, but these variants do not change the unstructured nature of the N terminus. For now, we do not know if both isoforms of Pus1p are expressed in Drosophila, even though the two variants of PUS1 mRNA are equally abundant. However, it is tempting to speculate that expression of the inactive, alternatively spliced isoform of Pus1p could represent a mechanism for fine-tuning U2 snRNA pseudouridylation at the branch point recognition region and consequently regulating pre-mRNA splicing efficiency.

We should emphasize here that we consider only positive results as meaningful in this study; one could find many reasons why a protein lacks enzymatic activity when expressed in a heterologous system. Our data clearly demonstrate that Pus1p is an enzyme for U2 snRNA pseudouridylation in the yeasts S. cerevisiae and S. pombe, the fruit fly Drosophila, as well as mammals and amphibians. At the same time, pseudouridylation of U2 snRNA at position 43 can occur independently of Pus1p, at least in S. pombe and mouse (Fig. 1B,C). Redundant modification activity has been previously observed for pseudouridylation of position 34 in Xenopus U2 snRNA, when the guide RNA pugU2-34/44 was tested in the Xenopus oocyte system (Zhao et al. 2002). The corresponding guide RNA (SCARNA8 or U92) is highly conserved across species (Darzacq et al. 2002; Lestrade and Weber 2006; Deryusheva and Gall 2013). Later, the guide RNA-independent activity on U2 snRNA at position 34 was identified for the single-protein pseudouridine synthase Pus7p from Xenopus, human, and Drosophila (YT Yu laboratory; S Deryusheva and JG Gall, unpubl.). Thus, we were not surprised that position 43 also could be modified by two independent mechanisms. The remaining problem was to identify the Pus1p-independent modification machinery at that position.

In our previous study of Xenopus oocytes, we found that depletion of SCARNA8 (also called pugU2-34/44 and U92) resulted in depletion of pseudouridylation not only at position 44, but at position 43 as well (Deryusheva and Gall 2013). However, when we tested the orthologous Drosophila guide RNA in the yeast cell system, we could only detect modification activity on yeast U2 snRNA at positions 35 and 45 (equivalent to positions 34 and 44 in vertebrates). With the new evidence for redundant pseudouridylation activity on U2 snRNA, we reanalyzed potential base-pairing of Drosophila scRNA:ΨU2-35.45 and vertebrate SCARNA8 with U2 snRNA (Fig. 2A). In fact, the ability of vertebrate SCARNA8 to direct modification at positions 43 and 44 is reasonably stronger than Drosophila scRNA:ΨU2-35.45. Thus, as predicted, mouse SCARNA8 induced the

![FIGURE 1. Pus1p enzymatic activity on U2 snRNA in different species.](image-url)
pseudouridylation of yeast U2 snRNA at positions 35 and 45 when expressed in the pus7 Δ strain (Fig. 2B, top two traces), and pseudouridylation of yeast U2 snRNA at positions 45 and 44 when expressed in the pus1 Δ strain (positions 45, 44, and 35 correspond to positions 44, 43, and 34 in vertebrates) (Fig. 2B, magenta trace). The yeast U2 snRNA sequence at the branch point recognition region is slightly different from that of vertebrates, which makes base-pairing weaker (the mismatch is depicted by an arrow in Fig. 2A). Thus the modification of endogenous yeast U2 snRNA is probably less efficient. Pseudouridylation of the target positions became more prominent when an artificial RNA was tested, which contained a sequence identical to vertebrate U2 snRNA (Fig. 2C, magenta trace). In the same optimized substrate, Drosophila scaRNA:ΨU2-35.45 could not induce modification at the position corresponding to position 43 in vertebrate U2 snRNA (Fig. 2C, black trace). Thus we positively identified SCARNA8 as a guide for U2 pseudouridylation at position 43 in vertebrates, which makes SCARNA8 a triple guide for the highly conserved positions 34, 43, and 44 in the branch point recognition region. It is puzzling that Drosophila scaRNA:ΨU2-35.45 has the same base-pairing with yeast U2 snRNA for positioning U2-Ψ44 as a vertebrate SCARNA8 (Fig. 2A), yet the Drosophila RNA is not functional at this position (Fig. 2B, black trace). The efficiency might depend on the binding affinity with the substrate as a whole, not with each modifying position separately; thus, in vertebrate SCARNA8, stronger binding in the alternative configuration to position U2-Ψ45 might facilitate efficient sliding to position U2-Ψ44. We still need to learn more about guide RNA functionality, especially in the cases of complex multitarget pseudouridylation pockets.

Multiple guide RNAs are typically assigned to the most important modified positions. Pseudouridines at the branch point recognition region of U2 snRNA represent such functionally crucial modifications (Yu et al. 1998; Zhao and Yu 2004, 2007; Yang et al. 2005). Yet SCARNA8 is encoded by a single-copy gene in most vertebrate genomes. To find other potential guide RNAs for pseudouridylation of position 43 in U2 snRNA, we screened known vertebrate box H/ACA RNAs for antisense elements specific to the U2 snRNA branch point recognition region. SNORA71 stood out in this analysis because its 5′-terminal pseudouridylation pocket could base pair with U2 snRNA (Fig. 2A). Since both positions 43 and 41 can be modified by base-pairing within the same pseudouridylation pocket, we used the S. cerevisiae double mutant pus1Δsnr81Δ to test SNORA71 guide activity on U2 snRNA. When SNORA71 was expressed in this mutant strain, it could not restore pseudouridylation of the endogenous yeast U2 snRNA (Fig. 2B, two bottom traces). Moreover, it did not support modification of an artificial substrate identical to the branch point recognition region of vertebrate U2 snRNA. Even though we did not demonstrate SNORA71 modification activity on U2 snRNA, we could not rule it out by our experiments.

Another candidate for pugU2-43 guide RNA was identified in our recent bioinformatic search for novel snoRNAs in Xenopus oocytes. A novel box H/ACA RNA was found in intron 2 of the annotated transcript Xetov90007238m
and (iii) pseudouridylation of U2 snRNA at position 45, equivalent to position 44 in *S. pombe* U2 snRNA (Fig. 4D). This demonstrated that the newly identified *S. pombe* RNA is a genuine guide RNA for U2 snRNA pseudouridylation. More importantly, expression of this guide RNA in the *pus1Δ* *S. cerevisiae* strain induced modification of both position 45 and 44, although position 44 was modified less efficiently than in the wild-type strain (Fig. 4D). The pseudouridylation pocket of *S. pombe* pugU2-43/44 RNA has a one-nucleotide bulge in the upper stem when base paired with *S. pombe* U2 snRNA for positioning Ψ43 (Fig. 4B, star). It has been noticed previously in an *S. cerevisiae* cell system that stability of the upper stem structure in a pseudouridylation pocket-bearing hairpin might significantly influence guide RNA functionality (Xiao et al. 2009). We deleted this one nucleotide and expected to see a slight increase in guide activity on Ψ44, when the mutated guide RNA was expressed in the *pus1Δ S. cerevisiae* strain. Strikingly, the one-nucleotide deletion, which stabilized the configuration for Ψ44 positioning, completely abolished Ψ45 formation (Fig. 4D, black trace). These observations indicate that preexisting pseudouridylation at position 44 inhibits the formation of pseudouridine at position 45. Similar inhibition was observed previously when we tested *Drosophila* scaRNA:ΨU2-35.45 in the yeast cell system (Deryusheva and Gall 2013). If crosstalk between different positions within heavily modified regions is a general phenomenon, a mechanism for coordinated positioning of post-transcriptional modifications deserves special investigation.

**Conclusions**

We found two types of pseudouridine synthase that catalyze the formation of U2-Ψ43 (or 44) in yeasts, *Drosophila*, and vertebrates. One is the stand-alone enzyme Pus1p, which acts on U2 snRNA in yeasts, *Drosophila*, and vertebrates. At the same time, another pseudouridine synthase Cbf5/Dyskerin and a guide RNA are found in *S. pombe* and vertebrates. Based on this redundancy, we postulate that alterations in U2 snRNA modification do not contribute to the mitochondrial myopathy and sideroblastic anemia (MLASA) phenotype associated with *PUS1* mutations.

**MATERIALS AND METHODS**

**Guide RNA sequence analysis**

To identify the box H/ACA guide RNA that modifies U2 snRNA at positions 44 and 43 in *S. pombe*, we used the short sequence BLASTN search on PomBase (http://genomebrowser.pombase.org/Schizosaccharomyces_pombe/blastview); the query sequence was ggACAGATACgg (3’-side of the pseudouridylation pocket, highly conserved across species). Hits were screened for the TGAAAA sequence (expected yeast-specific 5’-side of the pocket) upstream of the query sequence and the ACA box downstream from the query.

![FIGURE 3. Expression of the novel Xenopus pugU2-43 RNA detected by RNA deep-sequencing of oocyte nuclear and cytoplasmic RNA (A) and by Northern blot analysis of *X. tropicalis* liver RNA (B). (A) IGV browser view of RNA deep-sequencing reads (PRJNA302326) previously generated in our laboratory (Talhouarne and Gall 2014) and aligned to the *X. tropicalis* 9.0 genome. The black bar in the gene model panel indicates pugU2-43 RNA newly identified in the nuclear fraction.](image-url)
sequence. The computationally identified guide RNA was verified by Northern blot analysis and 5’ and 3’ RACE.

Expression in yeast cells

Yeast strains used in this study were the following: S. cerevisiae BY4741, which served as the wild-type strain; mutant strains pus7Δ, pus1Δ, and pus1Δsnr81Δ were provided by Yi-Tao Yu (University of Rochester Medical Center); S. pombe wild-type strain 972, the mutant strain pus1::KanMX4, and its parental strain ED666 were provided by Peter Espenshade (Johns Hopkins University School of Medicine).

The construct for expression of mouse Pus1p in yeast (Behm-Ansantz et al. 2006) was kindly provided by Dr. Yuri Motorin (CNRS-Université de Lorraine). The insert originally cloned into p416GAL vector was cut with XbaI and XhoI and sub-cloned into p416ADH vector. To clone PUS1 ORFs from yeast, the annotated ORFs of S. cerevisiae PUS1/YPL212C and S. pombe PUS1/SPCC126.03 were amplified from genomic DNA of BY4741 and 972 yeast strains using proofreading DNA polymerase Q5 (New England Biolabs). RT-PCR was used to obtain PUS1 ORFs from yeast, the annotated ORFs of S. cerevisiae PUS1/YPL212C and S. pombe PUS1/SPCC126.03 were amplified from genomic DNA of BY4741 and 972 yeast strains using proofreading DNA polymerase Q5 (New England Biolabs). RT-PCR was used to obtain PUS1 ORFs from X. tropicalis, C. elegans, and D. melanogaster. Total RNA isolated from different tissues (oocytes and liver of X. tropicalis) or from whole organisms (wild-type strain of C. elegans and y w strain of D. melanogaster) was used as the templates for RT-PCR. Appropriate restriction sites were added with primers to clone the PUS1 coding sequences into various yeast expression vectors: p416GAL, p416ADH, p416TEF. Two annotated isoforms of CG4159 mRNA predicted to encode Pus1p in Drosophila (http://flybase.org) were amplified and cloned: One isoform produces a 410 aa protein (CG4159-PA), and the other gives rise to a 442 aa protein with an extra 32 aa at the N terminus (CG4159-PB). The C. elegans Pus1p coding sequence was as previously identified (Patton and Padgett 2003). Since disordered N- and C-terminal residues of human Pus1p were found not essential for enzymatic activity (Czudnochowski et al. 2013), we additionally made HA-tagged versions of the Pus1p expression constructs. This allowed us to monitor expression of the exogenous Pus1p. To add one copy of HA-tag to the C terminus of Pus1p, the corresponding in-frame sequence was included in the reverse primers.

Drosophila scaRNA:ΨU2-35.45 expression constructs were previously generated (Deryusheva and Gall 2013). To express mouse U92/SCARNA8, Xenopus pugU2-43, and S. pombe pugU2-43/44 guide RNAs, the corresponding coding sequences were amplified by PCR from genomic DNA and cloned into YEplac181 [LEU2 2μ] vector containing a GPD promoter, an RNT1 cleavage site, and an snR13 terminator; the vector was constructed in the Yi-Tao Yu Laboratory (University of Rochester Medical Center) (Huang et al. 2011). Additionally, a fragment of S. pombe genomic DNA containing most of the RPC10 protein coding sequence, the 3’-UTR and a downstream 232-bp region was cloned into YEplac181 between EcoRI and Hind-III restriction sites (see Fig. 4A). The Xenopus SNORA71 coding
sequence was introduced by replacement of snR191 in the fragment of the yeast NOG2 gene cloned into p415GAL1 vector. To make an artificial RNA substrate identical to the branch point recognition region of vertebrate U2 snRNA, a fragment of human U2 snRNA (nucleotides 29–52) was inserted into the human U87 scaRNA and a yeast expression construct was generated as previously described (Deryusheva and Gall 2013). PCR-based mutagenesis and overlap-extension PCR were used to make mutant and chimeric RNA-expressing constructs.

Junction regions and entire inserts were sequenced in all recombinant plasmids. Plasmids were introduced into yeast cells using standard lithium acetate methods. Selective SC and EMM media were used to grow S. cerevisiae and S. pombe, respectively, with either glucose or galactose as a source of sugar.

**RNA extraction**

Hot acid phenol extraction was used to isolate RNA from yeast cells. TRIzol reagent was used to extract RNA from X. tropicalis oocytes and liver, while D. melanogaster adults, and whole C. elegans worms. After phase separation, RNA was purified using the Direct-zol RNA MiniPrep kit (Zymo Research). All manipulations with X. tropicalis were performed according to established and approved IACUC protocols.

**Northern blot**

Total RNA was separated on 8% polyacrylamide–8 M urea gels and transferred onto a nylon membrane (Zeta Probe, Bio-Rad). Digoxigenin (Dig)-labeled PCR probes were used to detect endogenous and overexpressed guide RNAs. Hybridization was performed at 42°C in DIG Easy Hyb buffer (Roche). The hybridized probes were visualized using an anti-Dig antibody conjugated with alkaline phosphatase and the chemiluminescent substrate CDP-Star (Roche) according to the manufacturer’s protocols.

**Western blot analysis**

For protein extraction, yeast cells were centrifuged, resuspended in 0.1 N sodium hydroxide, and incubated for 5 min at room temperature. Samples were pelleted, resuspended in the 2× sample buffer, and boiled for 5–6 min. Proteins were separated on Tris–SDS acrylamide gels, transferred onto a PVDF membrane, and probed with rat anti-HA mAb 3F10 (Roche) at a final concentration of 40 ng/ml in 5% nonfat dry milk. A secondary anti-rat antibody conjugated with horseradish peroxidase (Jackson ImmunoResearch) was used at a dilution of 1:40,000. SuperSignal West Dura chemiluminescent substrate (Thermo Scientific) was used to detect HRP.

**RNA modification analysis**

A primer extension-based modification mapping method was used to analyze pseudouridylation of yeast and mouse U2 snRNA (Kiss and Jädy 2004; Deryusheva and Gall 2009). Fluorescently labeled oligonucleotides specific for vertebrate and S. cerevisiae U2 snRNA and for U87-based artificial substrate RNA were previously described (Deryusheva and Gall 2013; Deryusheva et al. 2012). To analyze S. pombe U2 snRNA, a new 6-FAM-labeled oligonucleotide was designed: [106–129] AAGCCAGAGGCTTTCCA0CT0AAA. To detect pseudouridines, test RNA samples were treated first with CMC [N-cyclohexyl-N’-(2-morpholinoethyl) carbodiimide metho-p-toluene sulfonate] (Sigma-Aldrich) for 20–30 min and then with pH 10.4 sodium carbonate buffer for 3–4 h at 37°C. Primer extension was performed using either AMV-RT (New England Biolabs) or EpiScript RT (Epicentre) at 0.5 mM dNTP concentration. Reaction products were purified by ethanol precipitation; dry pellets were dissolved in formamide, mixed with the GeneScan-500 LIZ Size Standard and separated on an ABI3730xl capillary electrophoresis instrument (Applied Biosystems).

**COMPETING INTEREST STATEMENT**

The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health. J.G.G. is an American Cancer Society Professor of Developmental Genetics.

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