REPORT

Pseudouridine and N^6 -methyladenosine modifications weaken PUF protein/RNA interactions

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ABSTRACT

RNA modifications are ubiquitous in biology, with over 100 distinct modifications. While the vast majority were identified and characterized on abundant noncoding RNA such as tRNA and rRNA, the advent of sensitive sequencing-based approaches has led to the discovery of extensive and regulated modification of eukaryotic messenger RNAs as well. The two most abundant mRNA modifications—pseudouridine (Ψ) and N^6 -methyladenosine (m^6A)—affect diverse cellular processes including mRNA splicing, localization, translation, and decay and modulate RNA structure. Here, we test the hypothesis that RNA modifications directly affect interactions between RNA-binding proteins and target RNA. We show that Ψ and m^6A weaken the binding of the human single-stranded RNA binding protein Pumilio 2 (hPUM2) to its consensus motif, with individual modifications having effects up to approximately threefold and multiple modifications giving larger effects. While there are likely to be some cases where RNA modifications essentially fully ablate protein binding, here we see modest responses that may be more common. Such modest effects could nevertheless profoundly alter the complex landscape of RNA:protein interactions, and the quantitative rather than qualitative nature of these effects underscores the need for quantitative, systems-level accounting of RNA:protein interactions to understand post-transcriptional regulation.

Keywords: epitranscriptomics; pseudouridine; N^6 -methyladenosine; PUMILIO; RNA binding proteins; RNA–protein interactions

INTRODUCTION

Over a hundred distinct chemical modifications of RNA have been described previously (Cantara et al. 2011; Machnicka et al. 2013; Karijolich et al. 2015; Gilbert et al. 2016; Schwartz 2016). While a majority of these modifications were identified from chemical analysis of abundant noncoding RNA (snoRNA, tRNA, and rRNA), novel and sensitive sequencing-based approaches have revealed the widespread prevalence of modified bases such as pseudouridine (ψ) (Fig. 1A), N^6 -methyladenosine (m^6A) (Fig. 1B), N^1 -methyladenosine (m^1A), and 5-methylcytosine (m^5C) in eukaryotic messenger RNAs, collectively termed the epitranscriptome (Carlile et al. 2014; Lovejoy et al. 2014; Meyer and Jaffrey 2014; Schwartz et al. 2014; Liu and Pan 2016 and references therein; Squires et al. 2012; Khoddami and Cairns 2013; Dominissini et al. 2016; Li et al. 2016b).

ψ and m^6A are the most abundant mRNA modifications known, with 0.2%–0.5% of uridine and ~0.5% of adenosine

residues converted to ψ and m^6A , respectively (Dominissini et al. 2016). Ψ s in mRNA are made by ψ synthases that use a guide RNA-dependent (Cbf5, Dyskerin) or -independent (PUS family proteins) mechanism to recognize and target specific uridines for modification in a manner dependent on nutrient availability and environmental stress conditions (Charette and Gray 2000; Gilbert et al. 2016). m^6A modifications are catalyzed by “writer” methyltransferases (METTL3, METTL14) that target the adenosine within an RRACH (R = G, A; H = A, C, U) consensus motif for modification. Unlike pseudouridines, m^6A modifications are reversible and can be removed by “eraser” demethylase proteins in a regulated manner (FTO, ALKBH5) (Meyer and Jaffrey 2014; Liu and Pan 2016; Zhao et al. 2017).

The abundance and broad distribution of mRNA modifications along with the high degree of conservation of the modifying enzymes suggest that these modifications may

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FIGURE 1. Pseudouridine (Ψ) and N⁶-methyladenosine (m⁶A) are the two most abundant mRNA modifications in eukaryotes. (A) Ψ, the 5' ribosyl isomer of uridine, is made by pseudouridine synthases via guide-RNA dependent and independent mechanisms. (B) N⁶-methyl modifications to adenosines are made by m⁶A “writers” and removed by “erasers.” (C) Cartoon representation of a structural model of human Pumilio2 in complex with an 8mer RNA whose sequence is shown below. PDB ID: 3Q0Q (adapted with permission from Lu and Hall 2011). The Pus7 recognition motif (UGUAR) in the sequence is boxed in blue.

have important functional roles in gene expression (Machnicka et al. 2013; Dominissini et al. 2016; Hoernes et al. 2016). Indeed, pseudouridine (ψ) has been found to cluster in functionally important domains in transfer RNA (tRNA), ribosomal RNA (rRNA), and U2 small nuclear RNA (snRNA), and shown to affect protein translation and pre-mRNA splicing (Charette and Gray 2000; Ge and Yu 2013; Karijolic et al. 2015). ψ was also observed to increase nonsense suppression of mRNA stop codons resulting in altered protein products (Karijolic and Yu 2011). Biochemical and genetic studies have revealed regulatory roles for m⁶A modifications in a wide variety of cellular processes including, but not limited to, pre-mRNA splicing, and mRNA transport, translation, and decay (Meyer and Jaffrey 2014; Liu and Pan 2016). m⁶A has also been implicated in germline development and fertility, maintenance of the cell cycle and circadian clocks, and in a wide variety of human dis-

eases including cancer (Meyer and Jaffrey 2014; Liu and Pan 2015, 2016). Despite these advances, we have a limited understanding of how these modifications exert their functions.

RNA modifications have been proposed to regulate gene expression via multiple molecular mechanisms (Zhao et al. 2017). The first and most obvious mechanism is direct recognition of a modification, and indeed the mammalian YTH domain-containing “reader” proteins specifically bind single-stranded m⁶A RNAs and can regulate their function and fate (Meyer and Jaffrey 2014; Wang et al. 2014; Liu and Pan 2016; Zhao et al. 2017). The m⁶A modification position and level in mRNA varies as a function of a complex and dynamic interplay between these “writers” and “erasers,” as expected for a regulatory phenomenon (Meyer and Jaffrey 2014; Liu and Pan 2016). In contrast, specific ψ reader or eraser proteins have not yet been identified, perhaps reflecting difficulties in identifying the more subtle modification, and in breaking the new ψ C–C bond (Zhao and He 2015). Second, RNA modifications can affect gene expression by altering the stability of RNA structure (Roost et al. 2015) and thereby access to RNA binding proteins (RBPs) and the translation or processing machinery (Liu et al. 2015; Roost et al. 2015). For instance, ψ has been shown to stabilize local secondary structure in RNA via increased base stacking and the ability to coordinate a water molecule through the N1 hydrogen (Davis 1995; Charette and Gray 2000; Jiang et al. 2013). Such local structural alterations have been shown to be important for efficient codon–anticodon recognition (Charette and Gray 2000; Ge and Yu 2013 and references therein) and for spliceosome assembly and function (Wu et al. 2016). m⁶A, on the other hand, destabilizes RNA duplexes (Liu et al. 2015; Roost et al. 2015). Sites with m⁶A tend to be in less structured regions in vivo (Spitale et al. 2015), suggesting that these modifications could behave as structural “switches” to regulate the accessibility of the nearby sequences (Liu and Pan 2016).

Here, we address a third likely model of regulation via these and other RNA modifications—altering the interactions between RNA and RNA binding proteins (RBPs). Between 3%–8% of eukaryotic genomes encode proteins dedicated to binding RNA (Glisovic et al. 2008). There is now compelling evidence that many RBPs bind to large sets of cytoplasmically and functionally related RNAs, presumably to coordinate and control function (Ule et al. 2003; Gerber et al. 2004; Morris et al. 2010). RNA modifications may alter these RBP•RNA pools, by either eliminating certain interacting partners or by subtly altering the balance and identity of bound RNAs and thereby altering gene expression and cellular states.

To test this hypothesis, we determined the effects of ψ and m⁶A on the binding of human Pumilio 2 (hPUM2) RNA binding protein to modified RNA. hPUM2 is a member of the highly conserved PUF family of sequence-specific, single-stranded RNA binding proteins (Fig. 1C), binds over 750 unique mRNA targets in vivo and plays critical roles in brain and germline development and stem cell maintenance (Galgano

et al. 2008; Morris et al. 2008). To incisively test for direct effects on RNA/protein interactions from these substitutions we systematically probed binding to simple, single-stranded RNAs. We found that ψ and m^6A modifications modestly weaken binding of hPUM2 to target RNA suggesting that there are additional functional roles of RNA modifications and a greater complexity in cellular post-transcriptional regulatory networks.

RESULTS AND DISCUSSION

We set out to ask the general question of whether and to what extent mRNA modifications might alter protein binding, using the RNA binding domain of human Pumilio2 as a model RBP (herein referred to as hPUM2). hPUM2 is composed of 8 PUF repeats each of which recognizes and makes specific interactions with bases in the RNA (Fig. 1C; Lu and Hall 2011). Analysis of crystal structures of human Pumilio proteins in complex with various RNAs (Lu and Hall 2011) suggests that ψ is not expected to directly disrupt hydrogen bond interactions with the protein but that the methyl group of m^6A and the *syn* conformational preferences of the N^6 methylamino group (Roost et al. 2015) may disrupt interactions. In addition, these modifications could have more subtle effects on stacking, van der Waals interactions, or local sterics and solvation that weaken affinity. To test this hypothesis, we determined the quantitative effects on affinity from introducing Ψ or m^6A residues at each U or A position within the hPUM2 consensus sequence.

RNA modifications weaken hPUM2 binding to RNA

Pseudouridine (Ψ) effects on hPUM2 binding

We first used a direct gel-shift binding experiment to measure binding affinities of hPUM2 to eight Ψ -containing 10mer RNA oligos (CCUGUAAAUA) (Table 1; Fig. 2A; Supplemental Fig. S1A–D). While weakened binding from these substitutions was observed and direct gel-shift experiments have the apparent advantage of direct observation of the complex of interest at equilibrium, this assay is subject to effects from alteration in equilibrium conditions upon addition of loading buffers and upon placement in the gel well and entry into the gel. These effects are expected to be most pronounced for weaker binders (Batey et al. 2001). We therefore also determined affinities by competition, using binding to a radiolabeled unmodified 13mer

TABLE 1. Thermodynamic and kinetic measurements of hPUM2 binding of modified RNAs

RNA Oligo	25 °C				0 °C
	K_D direct ^a (nM)	K_D comp ^{a,b} (nM)	k_{off} ($\times 10^{-2}$ s ⁻¹)	k_{on} (calc) ($\times 10^7$ M ⁻¹ s ⁻¹) ^d	k_{off} ($\times 10^{-5}$ s ⁻¹)
UCUUGUAUAUA (RU ₁₃)	0.14 (0.11, 0.18) ^c	0.34 (0.15, 0.66) ^c	-	-	-
CCUGUAAAUA	0.34 ± 0.01	0.66 ± 0.05	2.3 ± 0.3	3.5	1.2
CC Ψ GUAAAUA	0.70 ± 0.1	1.15 ± 0.05	3.8	3.3	2.8
CCUG Ψ AAAUA	1.54 ± 0.05	1.5 ± 0.3	6.2	4.1	5.2
CCUGUAAA Ψ A	1.3 ± 0.1	1.8 ± 0.2	5.3	2.9	2.7
CC Ψ G Ψ AAAUA	4.9 ± 0.6	3.2 ± 0.5	>13.6	>4.3	15.6
CC Ψ GUAAA Ψ A	9 ± 4	3.9 ± 0.2	>14.2	>3.6	10.8
CCUG Ψ AAA Ψ A	2.9 ± 0.2	3.30 ± 0.05	>8.9	>2.7	7.7
CC Ψ G Ψ AAA Ψ A	129 ± 31 (791 ± 9*)	13.5 ± 1.9	>7.9	>0.6	23.7
CCUGUAUAUAU	0.14 ± 0.0	0.22 ± 0.03	1.9 ± 0.3	8.6	1.1
CCUGUAUAUAU	0.19 ± 0.05	0.26 ± 0.02	2.5	9.6	1.5
CCUGUAUAUAU	0.40 ± 0.11	0.57 ± 0.03	6.2	10.9	3.2
CCUGUAUAUAU	0.42 ± 0.0	0.62 ± 0.27	5.0	8.1	2.6
CCUGUAUAUAU	5.3 ± 0.6	6.5 ± 1.9	-	-	13.9

^a K_D values shown represent mean ± standard deviations from two independent replicates unless otherwise noted.

^b K_D comp derived by competition against bound labeled RU₁₃ RNA.

^cMean and 95% confidence intervals (in parentheses).

^d k_{on} values were calculated from the equation $k_{on} = \frac{k_{off}}{K_D \text{ comp}}$.

^e K_D calculated by forcing the fit through a maximum fraction RNA bound of one; an $R^2 \geq 0.96$ was obtained, compared to an R^2 value of 0.99 with the maximum fraction

bound treated as a variable $\left(\text{Fraction bound}(Fb) = \left(Fb_{\max} \times \frac{[P]}{[P] + K_D} \right) + \text{background} \right)$.

This value is expected to represent an upper bound.

RNA (*RU₁₃) as a common signal (Supplemental Fig. S1E, F) and the relative ability of each unlabeled modified RNA to reduce binding of *RU₁₃. Absolute values of dissociation constants for the modified RNAs are then readily calculated using the well-established dissociation constant for *RU₁₃ (Fig. 2B; Supplemental Fig. S1; also see Materials and Methods).

The dissociation constants measured by the two methods are in good agreement with each other, with the exception of the weakest binding, triply modified oligo (3× Ψ), whose binding was 10-fold stronger by competition (Fig. 2C). The weaker apparent affinity observed in the direct assay presumably reflects labeled RNA dissociation that occurs prior to or during gel entry that may be stemmed at higher protein concentrations as nearby free protein may “rescue” binding. Thus, we adopt the value from the competition assay that compares the binding of each RNA with a common readout, and this assignment was supported by agreement with kinetic measurements (see below).

Our results indicate that changing any of the single uridines to Ψ weakens hPUM2 binding by two- to threefold, relative to the unmodified control oligo (Fig. 2C; Table 1). Replacing an additional uridine by pseudouridine resulted in a further two- to threefold decrease in affinity, in agreement with simple energetic additivity (Fig. 2D). Changing all three uridines to Ψ s reduced hPUM2 binding

approximately twofold more than predicted by an additive model (Fig. 2D), an effect that might represent cumulative error, increased base-stacking from the presence of three Ψ s in the free RNA that is disrupted upon binding, or effects within the bound complex.

*N*⁶-methyadenosine (*m*⁶A) effects on hPUM2 binding

*m*⁶A modifications have been shown to indirectly facilitate protein binding by destabilizing RNA secondary structure (Liu et al. 2015; Roost et al. 2015). Here we tested whether these modifications could directly alter the stability of an RNA:protein complex, distinct from indirect structural effects that involve flanking sequences. Using gel-shift assays as described above, we measured hPUM2 binding to a series of 11mer RNA oligos in which *m*⁶A replaced zero, one, or three of the adenosines in the PUM2 recognition sequence (CCUGUAUAUAU) (Fig. 3A,B; Supplemental Fig. S2A,B). In all cases good agreement was observed between the direct and competition gel-shift methods (Fig. 3C).

FIGURE 2. Ψ weakens hPUM2 binding affinity to target RNA. Binding isotherms (A) and inhibition curves (B) measuring hPUM2 affinities to RNA oligos (CCUGUAAUAUA) containing 0 (green circles), 1 (blue symbols), and 3 (red circles) Ψ modifications. Representative curves from two independent replicates shown. (C) Bar graph comparing mean hPUM2 K_D values measured directly (gray) or by competition (black). Error bars represent standard deviations. (D) Additivity of the effects of individual Ψ modifications. Bar graph showing the observed (black) mean K_D competition values for RNA oligos containing two or more Ψ modifications compared to values predicted based on energetic additivity from the individual modifications (gray). Error bars represent standard deviations. The predicted K_D (K_D pred) of an oligo containing multiple Ψ s was calculated as follows. The observed K_D s (K_D obs) of the singly modified oligos were first normalized by the K_D of the unmodified oligo (0x Ψ) to yield the corresponding K_D rel and K_D pred for an oligo with Ψ s at positions *i* and *j* was calculated as: K_D pred(*i, j*) = K_D rel(*i*) $\times$ K_D rel(*j*) $\times$ K_D obs(0x Ψ).

FIGURE 3. *m*⁶A weakens hPUM2 binding affinity to target RNA. Binding isotherms (A) and inhibition curves (B) measuring hPUM2 affinities to RNA oligos (CCUGUAUAUAU) containing 0 (green circles), 1 (blue symbols), and 3 (red circles) *m*⁶A modifications. Representative curves from two independent replicates are shown. (C) Bar graph comparing mean hPUM2 K_D values measured directly (gray) or by competition (black). Error bars represent standard deviations.

Replacing a single A with *m*⁶A at position 4 did not affect binding, but the modification at position 6 or 8 decreased binding affinity to RNA by ~2.5-fold each (Fig. 3A; Table 1). The oligo containing three *m*⁶As (3x *m*⁶A) bound ~30-fold weaker than the unmodified oligo (Fig. 3A; Table 1), and about three- to fourfold weaker than predicted from an energetic additivity model (Supplemental Fig. S2C, and see below).

Ψ and *m*⁶A modifications increase the rate of dissociation of the protein–RNA complex

RNA modifications can weaken binding affinity between protein and RNA by reducing their rate of productive association (k_{on}) or by increasing the rate of dissociation of the bound

complex (k_{off}). Measurement of these kinetic rate constants can provide valuable information about the physical processes underlying the formation or dissociation of the protein–RNA complex. We measured the dissociation rate constants of the 13 unmodified and modified RNA complexes, using a pulse-chase gel shift assay. The protein–RNA complexes were each first allowed to form by mixing a high concentration of protein and trace amounts of the radiolabeled RNA. A large excess of unlabeled “chase” RNA was then added to the incubation mix to sequester the protein that dissociated from the radiolabeled RNA. Aliquots were removed at regular intervals and monitored for the amount of undissociated radiolabeled complex remaining by loading onto a native gel. As the only variable in a dissociation experiment is the amount of time the complex spends with unlabeled chase RNA before being loaded on a gel, this experiment is not subject to the potential artifacts from direct gel-shift equilibrium measurements (e.g., if, say, 50% of the protein dissociates upon entering the gel, it does so for each timepoint and the observed rate constant for loss of protein-bound RNA is unaffected).

We initially carried out these measurements at 25°C, the temperature of the equilibrium binding measurements (Supplemental Fig. S3A). At this temperature, k_{off} for hPUM2 from unmodified oligos ($0\times\Psi$ and $0\times m^6A$) is $\sim 0.02\text{ sec}^{-1}$ ($t_{1/2} \sim 35\text{ sec}$). While we were able to measure reliable k_{off} values for oligos with 0 or 1 modifications (8/13 oligos), only limits could be obtained for oligos with multiple base modifications because dissociation was too fast (Table 1). We therefore also measured dissociation rates at 0°C where dissociation is slowed ~ 2000 -fold, allowing all k_{off} values to be quantitatively compared (Table 1; Supplemental Fig. S3B). The relative dissociation rates (k_{rel}) of the modified oligos at the two temperatures are in excellent agreement (Supplemental Fig. S3C).

The relative dissociation rates highly correlate with the relative binding affinities between the oligos (Supplemental Fig. S3D), indicating that the modified bases weaken binding affinity by speeding dissociation of the protein from the RNA and supporting the veracity of the competition K_D measurements (Table 1). Indeed, the only outlier to a linear relationship with slope 1 is the $3\times m^6A$ oligo, which has a threefold smaller effect on k_{off} than K_D (Supplemental Fig. S3D) and thus likely binds to hPUM2 threefold slower. The threefold effect on the binding rate constant matches the threefold weaker binding of the $3\times m^6A$ oligo relative to the predicted affinity from an energetic additivity model (Supplemental Fig. S3D), consistent with formation of a single-stranded structure that is not binding-competent, such as a more stacked state, or less favorable partitioning of a partially bound state such that dissociation is favored over formation of the fully bound complex.

Physical models for the effects of RNA modifications on protein binding

NMR and crystallographic studies have revealed that the unique N^1 proton of Ψ forms a hydrogen bond with the

phosphate backbone in single-stranded RNA (Davis 1995; Jiang et al. 2013), and Ψ and m^6A may stabilize base stacking (Davis 1995; Liu and Pan 2016). These effects would weaken protein binding if the stabilizing interactions in free RNA are disrupted in the protein bound complex. Conversely, although not observed in our experiments, modifications could increase protein affinity if there is increased base stacking in bound complexes or enhanced interactions with amino acid side chains.

RNA modifications: an additional layer of complexity in the regulation of gene expression

Pseudouridines and N^6 -methyladenosines are the most abundant mRNA modifications in eukaryotes (ranging from 100–1000 Ψ sites [Li et al. 2016a] and $\sim 12,000$ m^6A sites [Yue et al. 2015] in humans). While we used hPUM2 as a convenient in vitro test case to study the effects of these modifications, it is likely that Ψ and m^6A disrupt hPUM2:RNA interactions in vivo as well.

Human Pumilio proteins, like other PUF family RBPs, recognize a highly conserved core motif, typified by the sequence UGUA(N)_nAUA (Wickens et al. 2002; Gerber et al. 2004; Galgano et al. 2008; Morris et al. 2008). Notably, the recognition motif (UGUAR) of the conserved PUS7 Ψ synthase (Carlile et al. 2014; Schwartz et al. 2014) overlaps with the 5′ half of the PUF consensus sequence (Fig. 1C). Prompted by this observation, we used the RMBase database (Sun et al. 2015) and identified a number of in vivo targets of hPUM2 that appear to harbor these modifications at various positions near or within the binding site (see Supplemental Note S1 for examples).

Modifications near RBP binding sites can play an indirect role by enhancing the accessibility of these sites to RBPs (Spitale et al. 2015). For example, we have shown that m^6A can significantly destabilize RNA secondary structure to increase the accessibility of an occluded hPUM2 site (Supplemental Fig. S4). Conversely, modification-induced structural rearrangements could also lead to decreased accessibility.

Here, we have shown that m^6A and Ψ modifications within hPUM2 binding sites can directly affect the binding affinity of hPUM2, weakening it by two- to threefold per modification on average. In principle, such effects can occur for many other RNA binding proteins. Indeed, RMBase searches show that RNA modifications appear to be widely prevalent in the binding sites of many different RBPs including hnRNP C, Ago2, Lin28b, and TIAL1 among others (Sun et al. 2015). While it is possible that there are RBPs that are affected more profoundly by RNA modifications, modest affinity changes such as those observed herein nevertheless have the potential to substantially alter gene expression.

The effects of RNA modifications on protein binding may thus be modest; nevertheless, other modest but widespread effects profoundly influence biology, such as the less than two-fold effects on mRNA translation and abundance typically

observed to arise from microRNA regulation (Hendrickson et al. 2008, 2009; Bartel 2009). In this regard, RNA modifications may function analogously to microRNAs, as “rheostats” to fine tune gene expression (Baek et al. 2008; Hendrickson et al. 2008, 2009; Bartel 2009). Our results therefore, reveal an additional layer of complexity in RNA:protein interaction networks and underscore the need for genome-wide quantitative data and modeling to unravel their regulatory logic.

MATERIALS AND METHODS

Protein expression and purification

The RNA-binding domain (residues 706–1059) of *H. sapiens* PUM2 (a kind gift from Andre Gerber, University of Surrey, UK) was cloned into a custom pET28a-based expression vector in frame with an N-terminal His-tag and SNAP tag. The construct was transformed into *E. coli* protein expression strain BL21 (DE3). Protein expression was induced at an OD₆₀₀ of between 0.6 and 0.8 with 1 mM IPTG at 18°C for ~20 h. Cells were harvested by centrifugation at 4500g for 20 min. Cell pellets were resuspended in Buffer A (20 mM HEPES-Na, pH 7.4, 500 mM potassium acetate [KOAc], 5% glycerol, 0.2% Tween-20, 10 mM Imidazole, 2 mM DTT, 1 mM PMSF and Complete Mini protease inhibitor cocktail [Roche]) and lysed four times using an Emulsiflex (Avestin). The lysate was clarified by centrifugation at 20,000g for 20 min, nucleic acids were then precipitated with dropwise addition of Polyethylene Imine (Sigma-Aldrich) to a final concentration of 0.21% with constant stirring at 4°C and pelleted by centrifugation at 20,000g for 20 min. The supernatant was loaded on a Nickel-chelating HisTrap HP column (GE) and washed extensively over a shallow 10–25 mM imidazole gradient. Protein was eluted over a linear 25–500 mM gradient of imidazole. Peak protein fractions were pooled and desalted into Buffer B (20 mM HEPES-Na, pH 7.4, 50 mM KOAc, 5% glycerol, 0.1% Tween-20, 2 mM DTT) using a desalting column. The His-tag was removed by incubation with TEV protease for 13–16 h at 4°C, and the protein solution was loaded again on the HisTrap HP column. The flow-through containing cleaved protein was collected, desalted into Buffer B, and loaded on a Heparin column and washed extensively to remove any bound RNA. Protein was eluted over a linear gradient of potassium acetate from 50 to 1000 mM. Fractions were pooled and desalted into Buffer C (20 mM HEPES-Na, pH 7.4, 100 mM KOAc, 5% glycerol, 0.1% Tween-20, and 2 mM DTT), concentrated and diluted twofold with Buffer C containing 80% glycerol for final storage at –20°C.

RNA oligos used for binding experiments

Pseudouridine-containing 10mer RNA oligos (CCUGUAAAUA) were synthesized by the Protein and Nucleic Acid (PAN) facility at Stanford University on an ABI 394 DNA synthesizer using standard RNA protocols and commercially available pseudouridine (Glen Research). After deprotection, RNA oligos were purified using Glen-Pak RNA cartridges (Glen Research) following manufacturer's recommendations.

N⁶-methyladenosine 11mer oligos (CCUGUAUAUAU) were synthesized with a MerMade 6 DNA/RNA synthesizer (Bioautomation)

utilizing standard phosphoramidite chemistry and commercially available N⁶-methyladenosine phosphoramidite (Glen Research). The short m⁶A oligos contained an additional 3' uridine in order to simplify their synthesis (m⁶A is available as phosphoramidite but not 3'-coupled to a synthesis column). Standard deprotection protocols for base and 2'-hydroxyl deprotection were used, with subsequent purification using Glen-Pak RNA purifications cartridges following manufacturer's recommendations (www.glenresearch.com). Oligos were then ethanol precipitated and dissolved in TE buffer (10 mM Tris-Na, pH 8, 1 mM Na-EDTA). All synthesized RNAs were observed to migrate as a single band on a native polyacrylamide gel. In addition, we observed >95% purity for three m⁶A hairpin oligos (Supplemental Fig. S5), which were tested by reverse-phase HPLC.

RNA oligonucleotides (25 pmol) were radiolabeled using [γ -³²P] ATP (Perkin-Elmer) and T4 Polynucleotide Kinase (NEB) for 30 min at 37°C and purified from nondenaturing 20% acrylamide gels run in 0.5× TBE (50 mM Tris-Na, 41.5 mM Boric Acid, 0.5 mM Na-EDTA).

Electrophoretic mobility shift assays (EMSA)

For equilibrium measurements, serially diluted protein amounts were incubated with trace amounts (<40 pM) of radiolabeled RNA in 20 μ L reaction volumes for ≥ 30 min at 25°C in buffer containing 20 mM HEPES-Na, pH 7.4, 100 mM KOAc, 0.1% Tween-20, 5% glycerol, 0.1 mg/mL BSA, 2 mM MgCl₂ and 2 mM DTT. Equilibrium conditions were verified by varying incubation time (2–510 min) and the amount of radiolabeled RNA (threefold), as well as by measuring the dissociation kinetics of the protein–RNA complex (see below). An aliquot of each reaction (7.5 μ L) was mixed with 5 μ L of ice-cold quench solution containing a 2.5 μ M unlabeled version of the same RNA, 6.25% Ficoll 400, and 0.075% bromophenol blue. A specified aliquot of that mixture (7.5 μ L) was immediately loaded on a preequilibrated, native 20% acrylamide gel in 0.5× TBE. Gels were run at 750 V for ~1 h before drying and exposing to phosphorimager screens. Screens were scanned on a Molecular Dynamics Typhoon instrument (GE Healthcare). Bound and free RNA were quantified using TotalLab Quant and fit using the hyperbolic equation

$$\left(\text{Fraction bound}(F_b) = \left(F_{b_{\max}} \times \frac{[P]}{[P] + K_D} \right) + \text{background} \right)$$

in Kaleidagraph (Synergy Software). We observed nonunity endpoints in our direct binding experiments. These can arise from a fraction of the material being damaged and/or from dissociation of the bound complex prior to or during gel entry, neither of which should affect measured K_D values.

For competition experiments, serially diluted, unlabeled modified RNAs were incubated at 25°C for ≥ 30 min with hPUM2 and radiolabeled RNA (*UCUUGUAUAUAUA; RU₁₃; ≤ 0.05 nM) in 20 μ L reaction volumes at a concentration of protein (0.5 nM) at which 50%–70% of the p*RNA was bound. A 30 min incubation was determined to be sufficient for equilibration based on the controls in the direct binding experiment and measurements of the dissociation kinetics. Reactions were quenched in ice-cold quench solution containing 1 μ M unlabeled RU₁₃ RNA and immediately loaded on a preequilibrated, native 20% acrylamide gel in 0.5× TBE. Gels were

run, dried, and quantified as above. Competition data were fit to linear and quadratic models (Ryder et al. 2008) for one-site specific inhibition in Kaleidagraph (Synergy Software) with excellent agreement between the K_D values obtained (within 9%). All binding measurements with modified RNAs were carried out in parallel with unmodified RNA to provide the most accurate relative affinities.

Dissociation rate constants were monitored by gel-shift pulse chase experiments as follows. A saturating amount of hPUM2 protein (typically 10–50 nM for 25°C experiments and 100–500 nM for 0°C experiments) was incubated with radiolabeled RNA for ≥ 15 min. A large excess of unlabeled RNA (10–20 times the protein concentration) was then added to the incubation mix. Aliquots were removed at regular intervals and mixed with ice-cold gel loading buffer (as above without unlabeled RNA); 7.5 μ L of this mix was loaded on a pre-equilibrated, native 20% acrylamide gel in 0.5 $\times$ TBE; and gels were run, dried, and quantified as above. Dissociation rate constants were extracted from a fit to a model for single exponential decay.

The active fraction for hPUM2 was determined by titration experiments. Briefly, varying protein amounts were incubated with a fixed, saturating concentration (26 nM) of RU₁₃ RNA and reactions were incubated at 25°C for ≥ 30 min. An aliquot of each reaction (7.5 μ L) was mixed with 5 μ L ice-cold quench solution containing 2.5 μ M unlabeled RU₁₃ and electrophoresed as above. The active fraction of the hPUM2 preparation used in these experiments was $\sim 55\%$ and the dissociation constants reported here have been corrected appropriately by the amount of active protein.

UV melting experiments

Absorbance versus temperature profiles for the structured m⁶A-containing RNA oligos were collected on a Shimadzu 1800 UV-vis spectrophotometer using an eight-chamber quartz microcuvette. RNA oligos were diluted twofold in buffer containing 40 mM sodium cacodylate, pH 7, 2 M NaCl, and 1 mM Na-EDTA. Melt profiles were collected at 260 nm in duplicate for at least three different concentrations of each RNA with a constant heating rate of 1°C/min and subsequently analyzed using LIFFT program (<https://github.com/DasLab/LIFFT>). The Nupack web server was used to predict secondary structures of the RNA oligos used in these experiments (Zadeh et al. 2011).

SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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