Pseudouridine-mediated translation control of mRNA by methionine aminoacyl tRNA synthetase

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ABSTRACT

Modification of nucleotides within an mRNA emerges as a key path for gene expression regulation. Pseudouridine is one of the most common RNA modifications; however, only a few mRNA modifiers have been identified to date, and no one mRNA pseudouridine reader is known. Here, we applied a novel genome-wide approach to identify mRNA regions that are bound by yeast methionine aminoacyl tRNA\textsubscript{Met} synthetase (MetRS). We found a clear enrichment to regions that were previously described to contain pseudouridine (\(\Psi\)). Follow-up \textit{in vitro} and \textit{in vivo} analyses on a prime target (position 1074 within \textit{YEF3} mRNA) demonstrated the importance of pseudouridine for MetRS binding. Furthermore, polysomal and protein analyses revealed that \(\Psi\) 1074 mediates translation. Modification of this site occurs presumably by Pus6, a pseudouridine synthetase known to modify MetRS binding. Consistently, the deletion of Pus6 leads to a decrease in MetRS association with both tRNA\textsubscript{Met} and \textit{YEF3} mRNA. Furthermore, while global protein synthesis decreases in \textit{pus6\Delta}, translation of \textit{YEF3} increases. Together, our data imply that Pus6 ‘writess’ modifications on tRNA and mRNA, and both types of RNAs are ‘read’ by MetRS for translation regulation purposes. This represents a novel integrated path for writing and reading modifications on both tRNA and mRNA, which may lead to coordination between global and gene-specific translational responses.

GRAPHICAL ABSTRACT

INTRODUCTION

Over the past decades, >100 post-transcriptional RNA modifications have been identified in all kingdoms of life (1). One of the most common RNA modifications in living cells is pseudouridine (\(\Psi\)) (2,3). The isomerization of uridine to pseudouridine has a similar base pairing as uridine, yet it enables stabilization of RNA structure by an extra hydrogen bond (4,5). In yeast, non-coding RNAs such as rRNA, snoRNA, snRNA and tRNA are highly enriched with pseudouridine at dozens of specific positions (6,7).

The isomerization of pseudouridine is catalyzed by two groups of enzymes (‘writers’). The first group, operating as an RNP complex, uses guide-RNA-based pseudouridine isomerization in which the essential Cbf5 enzyme is guided by H/ACA box snoRNA to its target tRNA sites (7–9). The second pseudouridine synthase (PUS) group is the stand-alone enzyme family that catalyzes the reaction without additional proteins or RNAs (10). Pseudouridine isomerization of cytosolic and mitochondrial tRNAs (11–16) is catalyzed by these enzymes.

Recently, high-throughput pseudouridine-seq studies revealed hundreds of pseudouridine sites in 509 different mRNAs. Most of this mRNA pseudouridylation activity was catalyzed by the stand-alone tRNA PUS family. Moreover, those studies indicated that pseudouridylation may be reg-
ulated in response to environmental changes (17–19). While in vitro analysis revealed an impact on translation (20), the in vivo impact of this post-transcriptional modification on mRNA activity remains largely unknown.

Modifications in RNA molecules usually serve as recognition sites to proteins (‘readers’) that exert a regulatory function. To date, the only identified pseudouridine reader is the yeast Prp5 RNA helicase. Prp5 interacts with Ψ42 and Ψ44 within U2 snRNA and promotes pre-mRNA splicing (21). Surprisingly, no mRNA or tRNA pseudouridylase readers have been identified thus far.

The family of aminoacyl tRNA synthetases (aaRS) sustains extensive interactions with pseudouridylated RNAs (i.e. tRNAs). aaRSs identify their cognate tRNA and catalyze the charging reaction of a specific amino acid at the tRNA 3′ end (22). The interaction of a specific tRNA with its cognate aaRS depends on unique tRNA identity elements. These elements are located mainly at the acceptor stem and the anticodon stem–loop (23–25). tRNA identity elements are also a hotspot of RNA modifications. These post-transcriptional RNA modifications can change the tRNA structure and may play a central role in tRNA–aaRS interactions (26). However, the in vivo impact of tRNA pseudouridylation on aaRS binding is not known. The association of aaRSs with RNA is not restricted to tRNAs (27,28). Recent interactome studies revealed that aaRS interaction regulates its cognate aaRS depends on unique tRNA identity elements (31). Various anticodon-like elements within mRNA appeared key for aaRSmRNA interaction and regulation (32–34); however, a role for mRNA modification is unknown.

Herein, we performed an in vivo binding-site mapping for Saccharomyces cerevisiae MetRS. Comparing these data to previously published pseudouridine-seq data revealed that MetRS preferentially binds Ψ-enriched regions. We revealed that Ψ at position 1074 of YEF3 mRNA and Ψ 31 of elongator tRNA Met are important for MetRS binding. The former interaction regulates YEF3 protein synthesis while the latter has a global effect on translation. Furthermore, our data suggest that both modifications are made by Pus6, suggesting a pivotal role for this enzyme in regulating global and gene-specific translation.

**MATERIALS AND METHODS**

**Yeast strains and growth conditions**

The following yeast strains were used: endogenously TAP-tagged (TAP-His3MX) Mesi (MetRS) in the following genetic background (ATCC 20188: MATα his3Δ1 leu2Δ0 met15Δ0 ura3Δ0) (35); endogenously GFP-tagged (GFP-S65T)-His3MX-Mesi (MetRS) in the following genetic background (ATCC 20188: MATα his3Δ1 leu2Δ0 met15Δ0 ura3Δ0) (36); and endogenously N terminally GFP-tagged Pus6 (YGR169C) in the background of BY4741 as described in (37). Proper fusion and expression of the tags were verified by PCR and western analysis. The knockout of PUS6 was done in the MetRS-GFP background by homologous recombination using a LEU2 cassette amplified from pRS405 as described in (38). Cells were usually grown in YPD or SCD media (39) at 30°C to mid-logarithmic phase.

**Fragmentation of RNA and ImmunoPrecipitation (fRIP)**

MetRS-TAP strain was grown in YPD to mid-logarithmic phase and subjected to cross-linking by the addition of formaldehyde (0.1% final concentration) for 10 min at room temperature. Cross-linking was terminated with 0.125 M glycine for 3 min, and cells were lysed in Buffer B (20 mM Tris–HCl [pH 7.5], 140 mM NaCl, 0.1% NP40, 0.1% SDS, 0.5 mM EDTA, 1 mM DTT, 2 mM PMSF, 10 μg/ml Leupeptin, 14 μg/ml Pepstatin, 0.02 U/ml RQ1 RNase-free DNase [Promega], 0.24 U/ml Ribolock RNase Inhibitor [Thermo Scientific]). Lysate was cleared by centrifugation for 10 min at 10,600 g at 4°C and fragmented by 0.2 U/ml RNase I (LifeTech, AM2295) for 4 min at 37°C with gentle shaking. For background cleavage pattern, 10% of the cleaved lysate was set aside and RNA was subjected to phenol:chloroform extraction (‘Input’ sample). The remaining 90% of the cleaved lysate was loaded on IgG Sepharose beads for 2 h at 4°C and washed four times in Buffer C (20 mM Tris–HCl [pH 7.5], 1 M NaCl, 0.5% NP40, 0.1% SDS, 0.5 mM EDTA, 0.5 mM DTT, 0.01 U/ml Ribolock RNase Inhibitor [Thermo Scientific]). MetRS was eluted by cleavage with 80 U of TEV (Invitrogen 12575-015) for 2 hr at 16°C in TEV buffer. Crosslinking was reversed by heating at 65°C for 16 h in reverse cross-linking buffer (50 mM Tris–HCl [pH 7.5], 5 mM EDTA, 10 mM DTT, 1% SDS, 0.1 U/ml Ribolock RNase Inhibitor [Thermo Scientific]) and RNA was precipitated following phenol:chloroform extraction (‘IP’ sample).

**RNA-binding proteins ImmunoPrecipitation and qPCR (RIP-qPCR)**

RIP-qPCR analysis was done as described in (33). Briefly, for mRNA RT-qPCR, RNA was reverse-transcribed using a high-capacity cDNA reverse transcription kit (Maxima according to the manufacturer’s instructions. Gene-specific levels were determined in a 25 μl reaction volume in triplicate with a Power SYBR Green PCR Master Mix® (Applied Biosystems) two-step RT-PCR method following the manufacturer’s instructions using primers for the indicated genes (Supplementary Table S1). All qPCR reactions used the following parameters: 50°C for 2 min, 95°C for 10 min, and then 30 s at 95°C and 1 min at 60°C for 40 cycles. Results were analyzed with Applied Biosystems 7500 Real-Time PCR Software v2.0.6. Fold change was calculated using either $2^{-\Delta \Delta C_{t}}$ or $2^{-\Delta C_{t}}$.

**RNA-seq analyses**

RNA samples were subjected to RNA-seq at the Technion Genome Center. Libraries from the Input and Bound fRIP samples were prepared using a TruSeq RNA Library Prep Kit v2 (Illumina, CA, USA) according to the manufacturer’s instructions. Ribo-Zero RNA Removal Kit was applied according to the manufacturer’s instructions only for the Input samples. All samples were sequenced on Illumina platform, yielding 10 to 30 million reads per sample. Reads were mapped to the S288c S. cerevisiae version R64–2 genome using RNA STAR version 2.6.0–2. Peaks were detected using MACS2 callpeak (40) version 2.1.1.20160309.6 against input sample with essential command attributes.
CRISPR/Cas9 point mutagenesis

Point mutations were introduced using the single-plasmid CRISPR/Cas9 protocol (41,42). Briefly, plasmid (bRA66 backbone (Addgene #100952)) expressing guide RNA recognizing the mutation region was transformed into MetRS-GFP strain together with the double-stranded 80 bp DNA fragment that contains the desired mutation at its center (Supplementary Table S1). Transformants were grown on YP-Gal Hygromycin (200 μg/ml) plate to induce Cas9 expression. Three viable clones from the transformation with the 80 bp fragment were isolated and the modification region was sequenced; all included only a single change, T1074C.

35S methionine pulse labeling

Cells (10 ml) were grown to mid-log phase in synthetic complete (SCD) medium, washed and resuspended in 10 ml SC without methionine.35S methionine (NEG709A005MC, PerkinElmer) was added and growth was continued for another 30 min. The translation was quenched by the addition of cycloheximide (100 μg/ml final concentration). Proteins were extracted and resolved on 10% PAGE. Gels were exposed to a phosphorimager and radioactive signals were quantified by ImageJ.

Polysomal analysis

Polysomal analysis was done as described in (43). Briefly, 100 ml cells grown in YPD to mid-logarithmic stage were harvested and lysed by a bead beater in 0.4 ml of lysis buffer (20 mM Tris–HCl at pH 7.4, 140 mM KCl, 1.5 mM MgCl2, 0.5 mM dithiothreitol, 100 μg/ml cycloheximide, 1 mg/ml heparin, 1% Triton X-100). The lysate was cleared and centrifuged for 15 min at 9000 × g at 4°C and the supernatant was loaded onto a 12 ml 10–50% linear sucrose gradient. Gradients were centrifuged in a SW41 rotor (Beckman) at 35 000 rpm for 160 min and polysomal profiles were determined by monitoring RNA absorbance at 254 nm with ISCO UA-6. Polysomal association was calculated by dividing the polysomal area by the area from the beginning of the 40S to the end of the graph. The entire gradient was fractionated to fractions of similar volume. RNA was extracted from each fraction by the addition of an equal volume of 8 M guanidinium HCl and two volumes of 100% ethanol, overnight incubation at −20°C and centrifugation at 13 000 rpm for 30 min. Pellets were washed with 80% cold ethanol, and usually half of the sample was subjected to northern blot analysis with the indicated probes.

RESULTS

Fragmented RIP-seq (fRIP-seq) protocol for mapping MetRS-bound RNA regions

Our previous RNA-binding protein ImmunoPrecipitation (RIP-seq) study revealed a subset of mRNAs that are highly associated with yeast methionine aminoacyl synthetase (MetRS) (33). To narrow down RNA regions that MetRS is associated with, we established a fragmentation of RNA and RIP-seq (fRIP-seq) protocol. In fRIP-seq,
Figure 1. fRIP-seq protocol to identify RBPs binding regions. (A) Scheme of the fRIP-seq protocol. Cell lysates of TAP-tagged MetRS strain are subjected to mild RNase I treatment. Fragmented lysate was mixed with IgG Sepharose beads to isolate the MetRS-TAP-bound RNA. Input and immunoprecipitated (IP) fragments were subjected to RNA-seq. Peak calling algorithm (40) is used to define regions that are enriched in the IP compared to the Untagged control fragments. (B) fRIP-seq Input RNA samples from two biologically independent MetRS-TAP cultures and from untagged parental strain were subjected to RNA-seq. Scatter plots present the pairwise comparisons of sequencing reads. Pearson correlation coefficient is shown in each box. (C) fRIP-seq for IP samples, collected and analyzed as in B. (D) Examples of fRIP-seq read maps of IP and Input data from MetRS-TAP and untagged strains. Y axis to the left of every map indicates the number of reads (note that scales differ between genes). Schematic depiction of the gene is indicated below each map. (E) Assignment of significant peaks. Scatter plot indicates the fold-enrichment (compared to Untagged control) for each peak defined by MACS2 callpeak algorithm (40) vs. the significance (as log(10) of the q-value). Red-marked dots indicate peaks that are defined as significant (Irreproducible Discovery Rate (IDR) < 0.05). (F) Pie chart indicates the position of significant peaks along detected genes. (G) MetRS-bound peaks were analyzed for an enriched sequence motif by DREME (59). (H) MetRS-bound peaks were analyzed for enriched process terms by SGD GO Term Finder Version 0.86. The most significant terms (corrected P-value < 9.49E–13) are presented.

cellular lysate of a TAP-tagged MetRS strain (or, for that matter, any other RNA-binding protein) is subjected to a mild RNase I treatment prior to RNA-binding protein immunopurification (RIP) with IgG Sepharose beads. Fragmented samples from either before RIP (Input) or those eluted from the beads by TEV cleavage (IP) were analyzed by next-generation sequencing (Figure 1A). The genomic position of Input and IP fragments was compared and enriched positions (‘peaks’) were defined. While we applied this approach herein to MetRS-TAP, we note that we successfully applied it to GFP-tagged proteins using Chromotek GFP-Trap beads (Levi O. et al., in preparation).

The mild fragmentation in fRIP generates RNA fragments of ~200 nts, thereby defining the resolution of the approach (Supplementary Figure S1A). About 25% of tagged MetRS proteins were detected in the elution sample, in-
demonstrates the efficiency of MetRS isolation (Supplementary Figure S1B and C). Western analysis for Hxk1 control protein did not detect any signal at the bound sample, indicating the specificity of isolation. Furthermore, silver staining of all proteins in the bound sample detected only marginal levels of proteins other than MetRS, and all were detected in the control purification of an untagged strain (Supplementary Figure S1C). Overall, under these conditions, MetRS appears to be efficiently and specifically isolated.

RNA samples from two biological repeats of fragmented cell lysate (Input) and the IP sample were sequenced. A similar scheme was applied to the parental untagged strain to control for non-specific binding. A high correlation is apparent between the Input signals of the two MetRS repeats and the untagged strain (Figure 1B). Thus, the TAP-tagging of MetRS does not lead to a general effect on gene expression. Analysis of the IP samples revealed that while both biological repeats of MetRS-TAP were very similar (Pearson correlation of 0.907), both were significantly different from the untagged control IP (Figure 1C). Thus, the rRIP-seq protocol is very reproducible, attested by the high correlation between biological repeats, and highly specific, apparent from the low correlation with the untagged strain. Figure 1D presents read maps of four representative mRNAs. The high efficiency and specificity of binding is apparent upon comparison to the Input sample and/or the Untagged strain data.

Applying a peak calling algorithm (40) to define mRNA regions that are significantly higher compared to the overall background of the mRNA and reproducible in both biological repeats, identified 145 such regions scattered within 131 mRNAs (Figure 1E, Supplementary Table S2). Interestingly, these regions are mostly localized within the coding region of the mRNA (Figure 1F). DREME sequence alignment (44) identified several significant motifs, all containing UA dinucleotide, with GNUANC being the most significant (Figure 1G, Supplementary Figure S1D); we did not find any resemblance to sequences within canonical targets of MetRS (i.e. tRNA\textsuperscript{Met}) (45). GO term analysis revealed that binding is significantly enriched to mRNAs encoding translation factors (GO: 0002181) (Figure 1H, Supplementary Table S3). Altogether, these data suggest that MetRS regulates expression of translation factors through binding to their coding regions.

MetRS is a pseudouridine reader

The canonical function of MetRS is charging of initiator and elongator tRNA\textsuperscript{Met} with methionine. These tRNAs are known to be modified extensively, in particular, in uridines that are modified into pseudouridine (Ψ). Therefore, we examined whether MetRS-bound RNAs are enriched with Ψ. Three independent pseudouridine-seq studies were previously published (17–19). While these datasets are likely incomplete, as apparent from the low overlap in their hits (46,47), altogether they report on 750 sites of Ψ within \textit{S. cerevisiae} transcriptome. We compared these sites with the MetRS-bound fragments detected here, and a significant number (49 of 145, P-value 1.8e\textsuperscript{-48}) appeared to overlap (Figure 2A, Supplementary Table S4). Moreover, the peaks that contain a Ψ appear to be better bound by MetRS compared to the 96 not reported to be pseudouridylated (Figure 2B). Thus, pseudouridine enhances MetRS binding to its target regions. GO term analysis for the 49 peaks that contain pseudouridine recaptured the enrichment to mRNAs encoding cytosolic translation factors (Figure 2C, Supplementary Table S3), suggesting that regulation of this family occurs through Ψ.

Ψ1074 of YEF3 is important for MetRS binding

MetRS appeared to significantly bind a region around nt 1000 of the mRNA-encoding eEF3 elongation factor (YEF3) (Figure 3A, red bar). This region was reported by two independent studies to include a Ψ site at position 1074 downstream to the start codon (chrXII:637854) (18,19). Nucleotide 1074 is a wobble position of a UUU codon (Phe amino acid). Position 1075 is occupied by an A, thus generating a UA dinucleotide, in line with the core apparent by computational predictions (Supplementary Figure S1D). To examine the possibility that MetRS binds this site, we expressed \textit{S. cerevisiae} MetRS protein in bacteria and purified it through MBP column (Figure 3B). Purified protein was mixed and crosslinked with labeled 9-mer oligos identical to the 1074 region with either Ψ or U at its center (Figure 3C). A 2-fold higher association was observed in the Ψ-containing 9-mer compared to the mutant (Figure 3D). Importantly, the addition of a competitor cold tRNA abolished binding to either oligo (Figure 3E). We note that attempts to derive exact disassociation constant (K\textsubscript{D}) using

**Figure 2.** MetRS binds Ψ-enriched regions. (A) Venn diagram was generated from peaks identified as bound by the MetRS rRIP-seq analyses and specific mRNA ribonucleotide that was found to contain pseudouridine by Ψ-mapping studies (17–19). P-value was calculated by the Hypergeometric test. (B) MetRS bound peaks were separated into those that do not contain Ψ (non-Ψ) and those that do. Box plot presents the fold enrichment in binding compared to untagged control. P-value was calculated by the two-tailed t-test. (C) GO terms processes analysis was done to Ψ-containing peaks that are bound by MetRS using SGD GO Term Finder Version 0.86.
Figure 3. MetRS binds YEF3 mRNA at Ψ1074. (A) Genome browser map showing rRIP-seq coverage along YEF3 mRNA (UCSC Genome Browser). Red bar under the map indicates the MetRS binding region, as identified by the MACS2 callpeak (40). Arrows point to Ψ positions revealed by (18,19) along YEF3 ORF. HS indicates sites apparent upon heat shock. (B) Coomassie blue staining of MetRS purification steps. Noninduced lane (NI) contains a protein sample from bacteria transformed with MBP-MetRS plasmid before induction. Input (I) contains a protein sample from bacteria after expression induction. Elution includes the protein sample eluted from the MBP column. (C) Labeled 9-mer oligonucleotide (identical to the sequence surrounding Ψ1074), either with or without Ψ at its center, was UV-crosslinked to the indicated extent with purified MetRS and resolved on PAGE. Gel was dried and exposed to a phosphorimager. (D) Quantification of the MetRS-bound oligo to free oligo ratio from two independent crosslinking experiments. ** indicates P-value ≤ 0.01 for the difference between Ψ and U containing oligos at each UV exposure. Note that error bars for the U oligo are smaller than the data points. (E) UV crosslinking was done at the indicated amounts. Competitor cold yeast tRNA was added prior to crosslinking where indicated. (F) Point mutation was introduced by CRISPR/Cas9 into the genomic site of YEF3, replacing the encoded U1074 with C. Steady-state levels of wildtype (WT) and mutant (U1074C) YEF3 mRNA were quantified by RT-qPCR analysis from three independent biological repeats, each with three technical repeats, normalized to ACT1 mRNA levels. (G) MetRS-GFP expressing yeast strains, co-expressing endogenously either WT or U1074C YEF3 mRNA, were subjected to RIP. Amounts of bound YEF3 mRNAs were quantified by RT-qPCR analysis and normalized to their expression levels. The histogram presents the quantification of two independent biological repeats, each with three technical repeats. P-value was calculated by the dependent samples one-tailed t-test.

EMSA were futile, probably because affinity to the 9-mer was too low (not shown). Thus, the optimal MetRS binding site is probably longer.

To examine the significance of Ψ at position 1074 (Ψ1074 in vivo), we replaced the thymidine (T) in this genomic position with cytosine (C), thereby eliminating pseudouridylation, yet maintaining the same amino acid (both UUU and UUC encode Phe). The mutation was introduced in a seamless manner by CRISPR/Cas9, without the insertion of any selection marker or any other change in the YEF3 gene (42). Analysis of steady-state YEF3 mRNA levels revealed a small and insignificant change in the U1074C mRNA levels (Figure 3F). Significantly, however, RIP followed by RT-qPCR to cells expressing MetRS-GFP with
either normal or U1074C *YEF3* mRNA revealed a strong reduction in MetRS association with U1074C mutant compared to the WT transcript (Figure 3G). This demonstrates that Ψ1074 is important for MetRS association with *YEF3* mRNA *in vivo*.

### Ψ1074 confers a translation regulatory role

Protein binding to an mRNA usually serves expression regulation purposes, which can be exerted through changes in mRNA stability or translation. Since we did not detect significant changes in mRNA steady-state levels upon U1074C change (Figure 3F), we tested possible impact on protein synthesis. Western analysis revealed a clear increase in YeF3 protein levels upon U1074C point mutation (Figure 4A and B). We note that U1074C mutation leads to a silent codon change from UUU to UUC (both code for Phe), with UUC having a lower codon usage than UUU (18.4 versus 26.1) (48). Since lower usage is associated with lower translation (49), it is unlikely to explain the increased protein amounts we observe. Furthermore, both codons are recognized by the same tRNA (tRNA^GAA^) as no tRNA^AAA^ exist *S. cerevisiae* (45); hence, differences in cognate tRNA abundance also cannot explain the increased protein levels.

To substantiate the link to translation, polysome profiling through sucrose gradients was done to both strains. While ribosomal association of unmutated transcripts (i.e. *HXK1* and *CDC33*) appeared not to change, and possibly even decreased in the U1074C strain, the ribosomal association of the U1074C transcript appeared to increase (i.e. shifted to the right) (Figure 4C, Supplementary Figure S3B). Thus, Ψ1074 mediates *YEF3* translation.

### Pus6 affects MetRS binding to tRNA

Pseudouridine synthetases (PUSs) isomerize U to Ψ among many tRNAs. The Pus6 member of this family has only one cytosolic tRNA target, elongator tRNA^{Met}. Furthermore, U31 within elongator tRNA^{Met} is the only known cytosolic target of Pus6 (14). We performed RIP analysis for Pus6 tagged by GFP to examine its tRNA association *in vivo*. Purification efficiency of this protein appears similar to the other GFP-tagged MetRS strains (Supplementary Figure S2). RT-qPCR designed for tRNA detection (35) was used.
to explore its targets. A strong association with tRNA\textsubscript{Met} elongator is apparent, more than 10-fold higher than the control, non-target tRNA\textsubscript{Pro} (Figure 5A). A similar preference was observed for MetRS binding to tRNA\textsubscript{Met} (Figure 5B). These data provide an \textit{in vivo} support to tRNA binding specificity of Pus6 and MetRS.

We next wished to determine the impact of pseudouridylation by Pus6. The deletion of Pus6 did not have a significant effect on steady-state levels of either tRNA\textsubscript{Met} or tRNA\textsubscript{Pro}, suggesting that the modification is not important for tRNA stability (Figure 5C). Importantly, however, RIP analysis to MetRS-GFP detected a strong reduction in MetRS binding to elongator tRNA\textsubscript{Met} in pus6\textsubscript{Δ}, and a smaller and insignificant impact on binding to tRNA\textsubscript{Pro} (Figure 5D). This is not due to a decrease in MetRS purification upon Pus6 deletion (Supplementary Figure S2).

To the best of our knowledge, this is the first time that pseudouridylation of a tRNA is linked to tRNA synthetase binding \textit{in vivo}.

**Pus6 mediates global protein synthesis**

The impact of pseudouridylation on tRNA\textsubscript{Met} binding to MetRS may lead to a broad impact on protein synthesis. To examine this, we performed pulse labeling for all cellular proteins by \textsuperscript{35}S-met labeling of pus6\textsubscript{Δ} and its parental strain (WT) (Figure 6). A clear reduction in their synthesis rates upon Pus6 deletion is observed (Figure 6A and B). Intriguingly, polysomal RNA analysis revealed an increase in OD \textsubscript{254} signal of polysomal fractions in pus6\textsubscript{Δ} cells (Figure 6C and D). This suggests that the lower protein levels are due to slowed translation elongation, leading to accumulation of ribosomes on mRNAs upon Pus6 deletion.

**Pus6 binds and affects YEF3 translation**

Pus6 was previously reported to modify mRNAs (18). Therefore, we tested its possible role in mRNA regulation. \textit{In vivo} binding of Pus6 to mRNAs was assayed by RIP followed by RT-qPCR (Figure 7A). A significantly
higher association with YEF3 than with an mRNA that was not found to be pseudouridylated (ACT1) was observed. A strong association was also observed with CDC33 mRNA (encoding the translation initiation factor eIF4E), supplementing previous in vitro pseudouridylation analysis by purified Pus6 on CDC33 mRNA (18). Furthermore, Pus6 deletion did not affect steady-state YEF3 mRNA levels (Figure 7B), yet led to a significant reduction in MetRS association with YEF3 (Figure 7C). Thus, Pus6 is a ‘writer’ of pseudouridine on YEF3 that is ‘read’ by MetRS.

Concomitant with the decreased MetRS association, analysis of YEF3 protein levels revealed a clear increase upon Pus6 deletion (Figure 7D and E). A similar increase is observed for CDC33, another target of Pus6 (Figure 7D and E). Polysomal analysis to pinpoint the impact to translation revealed no change in the sedimentation of the control HXK1 mRNA, while both modified mRNAs, CDC33 and YEF3, have a higher polysomal association in pus6Δ cells (Figure 7F). Together, these data reveal that Pus6 deletion imposes an activating effect on translation of its target mRNAs, presumably through reduced MetRS binding to Ψ sites.

DISCUSSION

Functions of aaRS other than tRNA charging are well known and include diverse roles in mRNA regulation (50,51). Interestingly, some of these functions are mediated through recognition of mRNA elements that resemble tRNA elements (24). For example, elements that are similar to anticodons of tRNAHis or tRNAThr were found within mRNA targets of HisRS and ThrRS, respectively, (33,34) and both exert a translation regulatory role. Here, we show for the first time, the involvement of another established tRNA feature, Ψ modification, in mRNA recognition by aaRS followed by translation control. Specifically, we show that MetRS interacts with both tRNA and mRNA through a pseudouridine that is generated by Pus6. Interaction with these two types of RNA offers coordination between tRNA charging and mRNA translation. Our working model (Figure 8) poses that Pus6 modifies elongator tRNA Met, and this modification is important for MetRS binding (Figure 5D). The decreased association with MetRS likely leads to a decrease in tRNA Met charging. We speculate that lower levels of charged tRNA Met reduce rates of translation elongation, hence, the increase in polysomal complexes observed in pus6Δ cells (Figure 6D). In parallel, Pus6 is a pseudouridine writer on specific mRNA targets (Figure 7A) (17,18).

These sites are probably important for MetRS binding to mRNA, therefore upon Pus6 deletion the association of MetRS decreases (Figure 7C). Importantly, we show that Pus6 deletion leads to increased translation of mRNAs that are known to have a pseudouridine (Figure 7D–F). Altogether, we propose that pseudouridines introduced by Pus6 are read by MetRS to exert a global and a gene-specific translation regulation.

What is the regulatory purpose of this pathway? The opposing effects on general and gene-specific translation echo the well-studied yeast amino acid starvation pathway (52). In response to amino acids depletion, GCN2 phosphorylates eIF2α, which leads to sequestration of eIF2B, followed by inhibition of global translation yet increased translation of GCN4 mRNA (53,54). While the mechanistic details of our observation are different, it suggests another path for coordination between global and gene-specific regulation in response to environmental changes. Thus far, we were unable to identify the environmental conditions upon which Pus6 is activated; the obvious predictions of methionine or general amino acid changes did not appear to change Pus6 levels (data not shown). Nevertheless, pseudouridylation by Pus6 appears as a new means to coordinate between tRNA and mRNA functions and global and gene-specific translation regulation. Since pseudouridine is among the most abundant modifications in RNA, we expect that regulation will be implicated in many transcripts. Furthermore, a significant fraction of mRNAs that include Ψ and bound by MetRS encode components of the translation machinery. These proteins are key regulators of many cellular processes, suggesting an even broader impact on the expression.

The binding of MetRS within the coding region suggests that binding inhibits ribosome transit along the mRNA, thereby affecting the elongation rate. In such a case, one would expect to see an increase in ribosome density upstream to the MetRS binding site. However, a previous Ri-
Figure 7. Pus6 regulates translation of pseudouridylated mRNAs. (A) GFP-tagged Pus6 cells were subjected to RIP followed by RT-qPCR to the indicated mRNAs. The histogram presents the quantification of two independent biological repeats. \( P \)-values are relative to \( ACT1 \) mRNA levels and were calculated by the dependent samples one-tailed \( t \)-test. (B) RT-qPCR analysis of \( YEF3 \) mRNA levels normalized to \( ACT1 \) in WT and pus6 deleted cells (\( pus6\Delta \)). The histogram presents the quantification of two independent biological repeats. (C) MetRS-GFP expressing cells either containing (WT) or deleted of Pus6 (\( pus6\Delta \)) were subjected to RIP followed by RT-qPCR to \( YEF3 \) mRNA. The histogram presents the quantification of two independent biological repeats, normalized to the signal in the Input sample. \( P \)-value was calculated by the dependent samples one-tailed \( t \)-test. (D) Western analysis for the indicated proteins in WT and in \( pus6\Delta \) strains. Samples from two independent protein preparations are presented. Signals are from the same blot, from which irrelevant lanes were cropped. (E) Quantification of Yef3p and Cdc33p signals in WT and \( pus6\Delta \) cells normalized to the Hxk1p signal. Results are from three independent biological repeats. Error bars are SEM, and \( P \)-value was calculated by the dependent samples one-tailed \( t \)-test. (F) WT and \( pus6\Delta \) cells were subjected to polysomal separation followed by northern analysis for \( YEF3 \), \( CDC33 \) and \( HXK1 \) transcripts. Graphs present the quantification of northern signals within all polysomal fractions. Arrowheads point to the fraction with the highest signal (‘peak fraction’) per mRNA. Results are from three independent biological repeats, error bars are SEM.
bosome Density Mapping along YEF3 mRNA from cells grown under similar conditions did not reveal a higher density of ribosomes at that region compared to other regions along the mRNA (55). This suggests that impact is not through an elongation arrest. Due to the lack of apparent impact on elongation, we speculate that under normal conditions, the pseudouridine at position 1074 affects the translation initiation rate. How a modification that is located in the midst of an ORF affects events at the 5' end of the transcript is yet to be revealed.

Yef3 protein is an elongation factor that assists the ribosomes’ movement along the ORF and plays a role in ribosome recycling (56). It may also play a non-translational role (57). Recently, the impact of Yef3p depletion on ribosome density was studied at high resolution (58). Interestingly, the depletion of Yef3p led to a higher representation of methionine codons at the P-site of stalled ribosomes. This implies a preferred role for Yef3p in assisting ribosomes stalled on these codons. Reduction in charged ribosomes’ movement along the ORF and plays a role in ribosome Density Mapping along YEF3 mRNA from cells grown under similar conditions did not reveal a higher density of ribosomes at that region compared to other regions along the mRNA (55). This suggests that impact is not through an elongation arrest. Due to the lack of apparent impact on elongation, we speculate that under normal conditions, the pseudouridine at position 1074 affects the translation initiation rate. How a modification that is located in the midst of an ORF affects events at the 5' end of the transcript is yet to be revealed.

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**SUPPLEMENTARY DATA**

**Supplementary Data** are available at NAR Online.

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