



Metabolic turnover and dynamics of modified ribonucleosides by ^{13}C labeling

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Tandem mass spectrometry (MS/MS) is an accurate tool to assess modified ribonucleosides and their dynamics in mammalian cells. However, MS/MS quantification of lowly abundant modifications in non-ribosomal RNAs is unreliable, and the dynamic features of various modifications are poorly understood. Here, we developed a ^{13}C labeling approach, called ^{13}C -dynamods, to quantify the turnover of base modifications in newly transcribed RNA. This turnover-based approach helped to resolve mRNA from ncRNA modifications in purified RNA or free ribonucleoside samples and showed the distinct kinetics of the N6-methyladenosine (m^6A) versus 7-methylguanosine (m^7G) modification in polyA+-purified RNA. We uncovered that N6,N6-dimethyladenosine (m^6_2A) exhibits distinct turnover in small RNAs and free ribonucleosides when compared to known m^6_2A -modified large rRNAs. Finally, combined measurements of turnover and abundance of these modifications informed on the transcriptional versus posttranscriptional sensitivity of modified ncRNAs and mRNAs, respectively, to stress conditions. Thus, ^{13}C -dynamods enables studies of the origin of modified RNAs at steady-state and subsequent dynamics under nonstationary conditions. These results open new directions to probe the presence and biological regulation of modifications in particular RNAs.

RNA methylation modulates crucial RNA–protein interactions at various stages of RNA metabolism. Comprehensive studies of individual RNA modifications have been mostly advanced by applications of next-generation sequencing (NGS), which rely on chemical, enzymatic, and/or antibody-based detection of modified ribonucleosides (1–3). These methods have provided a wealth of information on the sites of modification across the transcriptome (1–3), which have uncovered the dynamic behavior of N6-methyladenosine (m^6A) mRNA methylation in development (4–8). Notably, dysregulation of m^6A levels has been recently linked to cancer, aging, and neurodegeneration (9–15), which can occur *via* metabolic inhibition of m^6A demethylation (9, 10). Furthermore, various tRNA modifications have been reported to be posttranscriptionally sensitive to cellular stress (16–19). A current caveat of NGS-based profiling is the lack of high-specificity reagents for

every modification (1–3). Also, the effects of transcription, which often predominate in differential expression analyses (20), make it challenging to quantify general changes in methylation levels, with limited insight being obtained into biological associations between various modifications. As a consequence, the dynamic behavior of RNA methylation often remains poorly understood, and complementary approaches are needed to quantify multiple modifications across biological contexts and to assess their associations to transcriptional versus posttranscriptional events.

Tandem mass spectrometry (LC-MS/MS) is a highly accurate tool for analysis of modified RNAs, which has so far been applied primarily in two approaches to identify and quantify RNA modifications (21). The first approach employs LC-MS/MS of intact RNA oligonucleotides to detect multiple modifications with positional information in specific RNA sequences. This approach is chromatographically challenging (22) and requires advanced data mining for unambiguous identification of RNA fragments, and it has so far been applied for comprehensive characterization of RNA modifications in abundant or short RNAs (22), such as rRNA (23–26), tRNAs (27, 28), and miRNAs (29). The second approach employs LC-MS/MS of ribonucleosides for sensitive quantification of RNA modifications (21). This approach, often combined with stable isotopes, can simultaneously quantify the abundance of multiple modifications in specific RNA classes of interest, being highly suitable to assess the presence of tRNA and rRNA modifications and their dynamics under various biological scenarios (19, 27, 30–35). These methods have so far been applied mainly to study these abundant RNA species because MS detection of mRNA modifications is unreliable due to the heavily modified ncRNAs, which invariably contaminate mRNA pools purified using either polyA+-enrichment or rRNA depletion (36, 37). Therefore, MS approaches are needed that are capable to account for and quantitatively resolve the origin of multiple RNA modifications in purified RNA. Moreover, the dynamic behavior of RNA modifications is insufficiently explained solely by changes in their abundance (or levels), as these do not inform on the underlying pathways driving these changes. Since the deposition (and removal) of RNA modifications is linked to the life cycle of the RNAs themselves (38–40), the change in methylation levels can result from changes either in RNA (de)methylation rates or in

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the transcription or decay of methylated RNAs, From this lens, methods are needed that can assess transcriptional and post-transcriptional effects on methylated RNAs under nonstationary conditions.

Stable isotope labeling is a well-established method to quantify metabolic activity in cultured cells (41–43). Here, we developed a quantitative approach using [^{13}C -methyl]-methionine labeling and mass spectrometry (MS) to assess the turnover of base modifications (^{13}C -dynamods) in newly transcribed RNA. With ^{13}C -dynamods, we trace the proportion of newly methylated ribonucleosides and their decay through time, *i.e.*, methylation turnover, from digested polyadenylated RNA and ncRNA. We first showed that polyadenylated RNA and ncRNA were distinguished by the different turnover frequencies (in h^{-1}) of modified ribonucleosides, which are inherently linked to the different half-lives of mRNA, rRNA, and tRNA (39, 44, 45). Examining the kinetics of methylation turnover at steady state within and across RNA classes as well as in free ribonucleosides enabled us to resolve the origin of RNA modifications in digested RNA and thereby uncover the presence of modifications in uncharacterized RNA classes, such as *N*6,*N*6-dimethyladenosine (m^6_2A). We then applied ^{13}C -dynamods in conjugation with abundance measurements of modified ribonucleosides in polyA+ and ncRNA, which resolved their transcriptional *versus* posttranscriptional sensitivity in response to actinomycin D and metabolic stresses. Thus, the quantitative nature of ^{13}C -dynamods demonstrates its capacity for sensitive characterization of modified ribonucleosides, their origin and dynamics, in multiple RNA classes of interest.

Results

^{13}C labeling of polyadenylated and ribosomal RNA modifications

S-adenosylmethionine (SAM) is the direct substrate of RNA methylation reactions in eukaryotic cells (46). To trace the incorporation of SAM into RNA, we cultured 786O cells in methionine-free DMEM medium supplemented with either unlabeled methionine (“Unlab”) or [^{13}C -methyl]-methionine and analyzed the isotopologues ($m + 0$, $m + 1$, $m + 2$) of modified and unmodified ribonucleosides by tandem mass spectrometry (LC-MS/MS) (Fig. 1, A and B). The $m + 0$ isotopologue (*e.g.*, 150.1 m/z for m^6A) represents the mass of the analyzed molecule where all atoms are present as the most common isotope, whereas the $m + 1$ isotopologue (*e.g.*, 151.1 m/z for m^6A) indicates the mass shift due to ^{13}C incorporation from the ^{13}C -labeled methionine tracer or from the natural abundance of ^{13}C , ^{15}N , ^{18}O , and ^2H stable isotopes. With ^{13}C -dynamods, we measure the isotopologue fractions [$m + 1/(m + 1 + m + 0)$] of modified and unmodified ribonucleosides, which reflect the amount of ^{13}C enrichment ($m + 1$) in the measured ribonucleoside relative to the total pool of that ribonucleoside ($m + 1 + m + 0$) (Fig. 1A). Thus, the isotopologue fractions of each ribonucleoside during [^{13}C -methyl]-methionine labeling are internally controlled for the amount of preexisting ribonucleosides ($m + 0$ and naturally labeled $m + 1$) prior to ^{13}C labeling. Here,

we analyzed the isotopologues of *N*6-methyladenosine (m^6A), 7-methylguanosine (m^7G), 1-methyladenosine (m^1A), *N*6,*N*6-dimethyladenosine (m^6_2A), 2'-*O*-methyladenosine (A_m), 5-methylcytosine (m^5C), and unmodified ribonucleosides from digested polyadenylated (polyA+), large (>200 nt) and small (<200 nt) RNA. We observed increased $m + 1$ and concomitant decreased $m + 0$ ion counts in modified ribonucleosides from polyA+ and large RNA after 4 and 24 h of [^{13}C -methyl]-methionine labeling, while the $m + 1$ fractions (natural abundance) of the unmodified ribonucleosides was unaltered during the labeling period (Figs. 1, C and D and S1A; Supplementary Materials). We determined that the ^{13}C enrichment of intracellular methionine and SAM reaches a 98 to 100% plateau within 30 min and remains constant thereafter (Fig. S1B). Thus, the change in the “heavy” ($m + 1$) isotopologue fraction relative to the unlabeled condition (natural abundance) indicates SAM-dependent RNA methylation of newly synthesized RNA, whose turnover we have examined within and across RNA classes. In contrast to singly methylated ribonucleosides, the m^6_2A modification exhibited an enrichment mostly of the $m + 2$ isotopologue upon ^{13}C labeling (Fig. S1A), and so the $m + 2$ fraction was used to assess m^6_2A dynamics in subsequent analyses.

Quantification of isotopologue fractions showed a faster kinetics for m^6A , m^7G , and m^5C in polyA+-purified RNA when compared with large (>200 nt) and small (<200 nt) RNA (Fig. 1E), in accordance with the faster life cycle of mRNA in mammalian cells (45). In contrast, the kinetics of m^1A , m^6_2A , and A_m methylation was similar between polyA+ and large RNA fractions (Fig. 1E), suggesting that a large portion of the signal for these modifications in the polyA+ fraction might derive from contaminating ribosomal RNA (rRNA). Of note, the isotopologues are analyzed on all ribonucleosides from the same polyA+ pool (Fig. S1C), so the purity of polyA+ enrichment is the same in all cases. This indicates that the relative abundance in mRNAs *versus* contaminating ncRNA is much higher for m^6A , m^7G , and m^5C compared with m^1A , m^6_2A , and A_m . Interestingly, we detected m^6_2A in small RNAs from 786O cells, and its kinetics was different when compared with large RNA (Fig. 1E).

Methylation turnover to resolve the origin of RNA modifications and their presence in uncharacterized RNA classes

To examine the kinetics of methylation turnover, we cultured cells for 24 h with [^{13}C -methyl]-methionine followed by replacement (chase) with unlabeled methionine and analyzed the isotopologues over time. In accordance with the preceding findings (Fig. 1E), we observed an exponential decay of the $m + 1$ isotopologue fraction for $\text{m}^6\text{A}/\text{m}^7\text{G}$ but not for $\text{A}_m/\text{m}^1\text{A}/\text{m}^6_2\text{A}$ in polyA+ RNA (Fig. 2A). Conversely, the modifications of total/large RNAs exhibited uniformly slow turnover (Fig. 2B), consistent with the high stability of rRNA and tRNA in growing cells (44). To test the kinetic behavior underlying methylation turnover, we examined the goodness of fit of a linear *versus* exponential regression of the

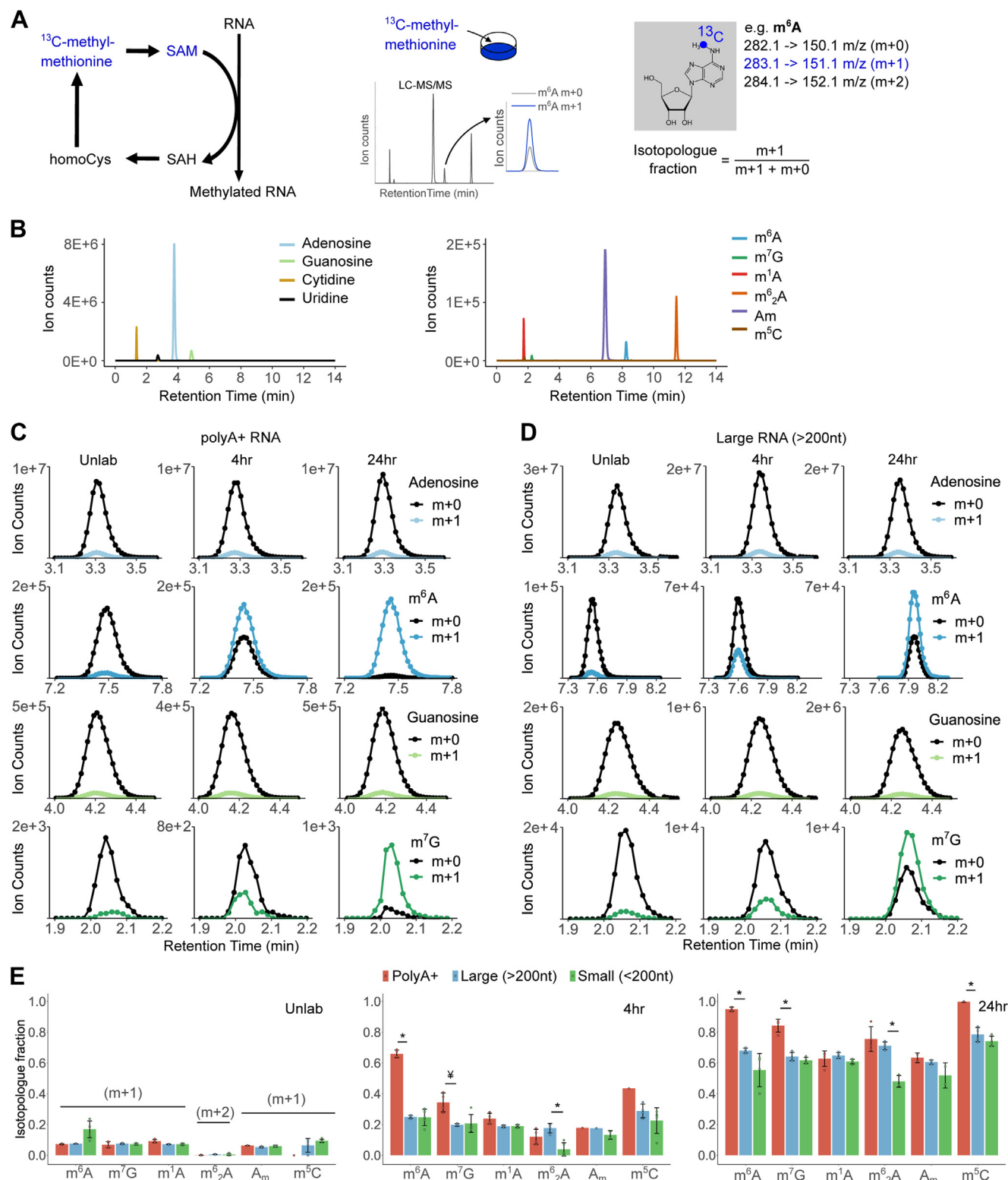


Figure 1. ^{13}C labeling of RNA modifications across RNA classes. A, the ^{13}C -dynamods workflow shows the tracing of RNA methylation: cells are cultured with [^{13}C -methyl]-methionine, the RNA is isolated, digested to ribonucleosides and subjected to LC-MS/MS analysis. The isotopologues detected for m^6A are shown. B, representative chromatogram of unmodified and modified ribonucleosides from total RNA. C and D, the $m+0$ and $m+1$ isotopologues of modified and unmodified ribonucleosides (representative chromatograms) in polyA+ (C) and large RNA (D). E, quantification of the isotopologue fractions of each ribonucleoside in polyA+, large and tRNA under unlabeled ('Unlab') conditions, after 4 and 24 h of culture with [^{13}C -methyl]-methionine. Error bars represent standard deviation of 3 to 4 biological replicates, with the exception of m^5C in polyA+-purified RNA (two replicates at 24 h, one replicate in the Unlab/4 h time points). In all cases, each replicate is the average of two technical replicates. * denotes $p < 0.005$, $\ddagger$ denotes $p < 0.05$, of a two-sided Student's t test comparing samples as indicated. Am, 2'-O-methyladenosine; homoCys, homocysteine; m^1A , 1-methyladenosine; m^1G , 1-methylguanosine; m^2G , 2-methylguanosine; m^5C , 5-methylcytosine; m^6A , N6-methyladenosine; m^6_2A , N6,N6-dimethyladenosine; m^7G , 7-methylguanosine; SAH, sadenosylhomocysteine; SAM, s-adenosylmethionine.

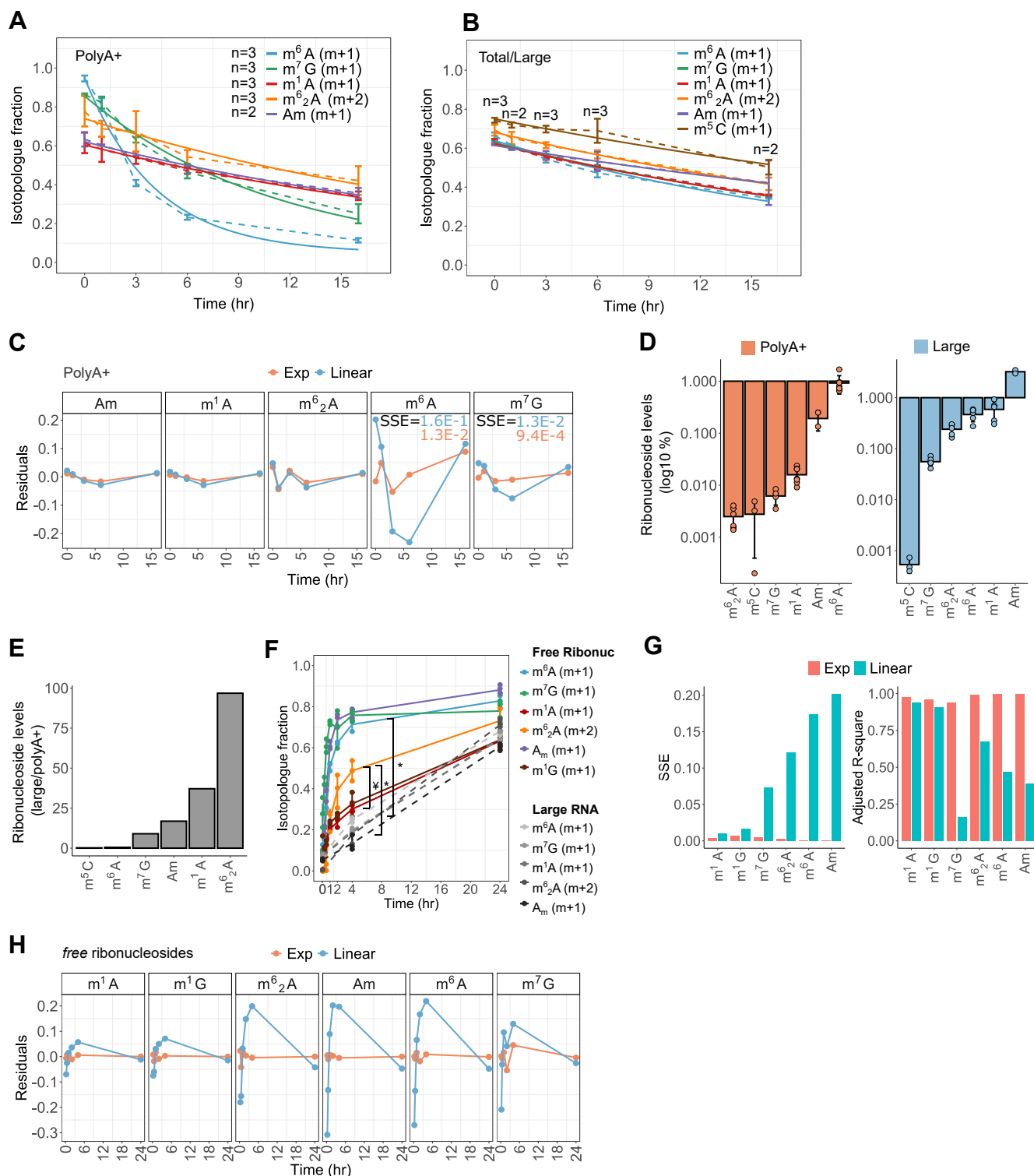


Figure 2. Turnover of modified ribonucleosides in polyA+, ncRNA, and free ribonucleoside pool. A and B, isotopologue fractions during the “chase” of ^{13}C -labeled modifications with naturally labeled methionine for 0, 1, 3, 6, and 16 h, in polyA+ (A) and total/large RNA (B); dashed lines connect data points; solid lines, exponential (A) or linear (B) fit of isotopologue fractions. C, residuals of a linear versus exponential regression of isotopologue fractions in polyA+ RNA in the chase experiment. D, normalized ion counts of modified ribonucleosides relative to the ion counts sum of all ribonucleosides, shown for polyA+ and large RNA. E, ratio of normalized ion counts between polyA+ and large RNA, as determined in (D). F, isotopologue fraction of free ribonucleosides analyzed from metabolic extracts; gray datapoints refer to large RNA modifications (Fig. 1E) for comparison. G and H, goodness of fit of a linear versus exponential fit of the isotopologue fractions in free modified ribonucleosides. Error bars represent 90% confidence intervals in (A and B), or standard deviation in (D and F) of at least three biological replicates with exception of Am (two replicates) and in (B) as indicated. In all cases, each replicate is the average of two technical replicas. * denotes $p < 0.005$, $\ddagger$ denotes $p < 0.05$, of a two-sided Student’s *t* test comparing m^6_2A and m^6A in the free ribonucleoside pool (free ribonuc) to large RNA, or m^6_2A to m^1A in the free ribonuc pool. A, adenosine; Am, 2'-O-methyladenosine; G, guanosine; m^1A , 1-

isotopologue fraction. Analysis of residual errors showed that the linear regression ($m_{(t)} = m_{(0)} + kt$) fits well the turnover of $m^1A/m^6_2A/A_m$ but not m^6A/m^7G in polyA+ RNA, in which the “U-shaped” curve supports a nonlinear model ($m_{(t)} = m_{(0)} e^{(-kt)}$) (Figs. 2C and S2A). Conversely, the linear regression fitted well the turnover of ncRNA modifications (Fig. S2, B and C). The turnover frequency determined for $A_m/m^1A/m^6_2A$ in polyA+ was similar to ncRNA modifications ($k = 0.031 \text{ h}^{-1}$ on average) and significantly slower than m^6A ($k = 0.244 \text{ h}^{-1}$) or m^7G ($k = 0.089 \text{ h}^{-1}$) in polyA+ RNA (Fig. 2, A and B). Thus, the turnover of $m^1A/m^6_2A/A_m$ from polyA+ fractions is incompatible with the exponential turnover of mRNA modifications (44, 45), confirming that they originate from ncRNA contamination.

The normalized abundance (frequency) of m^7G was in between that of m^6_2A and m^1A/A_m in polyA+ RNA (Fig. 2D), and the sensitivity of MS detection was similar across all measured modifications except m^5C , as determined from equimolar injections of pure compounds (Fig. S2D). Thus, the distinct turnover of RNA modifications in the polyA+-purified fraction was not explained by differences in normalized abundances alone or analytical sensitivity of ribonucleoside detection. Instead, the modified ribonucleosides were more abundant in large RNA relative to polyA+ RNA by a factor of 17/37/97/9 for $A_m/m^1A/m^6_2A/m^7G$, respectively, while m^6A level was greater in polyA+ RNA by a factor of 2 (Fig. 2E). From the determined turnover frequencies (h^{-1}) of ncRNA (Fig. 2B), we estimated that a 50% contribution of ncRNA signal to the ribonucleoside pool in polyA+-purified RNA would lower the detected turnover of the *bona fide* m^6A mRNA modification, with a turnover frequency of 0.242 h^{-1} , to that of m^7G , with a turnover frequency of $\sim 0.09 \text{ h}^{-1}$ (Fig. S2E). That is, reliable turnover-based detection of a hypothetical mRNA modification that is more abundant by a factor of 17 to 97 in ncRNA relative to mRNA is attainable with a maximal ncRNA contamination of 0.5% (50%/97) to 2.9% (50%/17), but even an optimal performance of available methods to purify polyA+ RNA commonly contains 2 to 3% of contaminating ncRNA (Fig. S2F) (36, 37). Thus, the ability to resolve *bona fide* mRNA modifications based on methylation turnover is limited by the depletion efficiency of highly abundant rRNA modifications.

As the turnover frequency of m^6A was approximately eight times faster in polyA+ RNA than ncRNA (Fig. 2, A and B), we reasoned that free ribonucleosides derived from metabolic extracts would be derived mainly from endogenous degradation of mRNAs, rather than ncRNAs. Thus, we employed ^{13}C -dynamods to examine the methylation turnover of free ribonucleosides, thereby enriching for mRNA-derived ribonucleosides and decreasing interference from rRNA-derived ribonucleosides. We found a fast kinetics following ^{13}C labeling that was similar for m^6A , m^7G , and A_m , indicating that they are primarily derived from the same RNA type (*i.e.*, mRNAs) (Fig. 2, F–H). Interestingly, the m^6_2A modification

also exhibited a nonlinear, faster kinetics than m^1A or 1-methylguanosine (m^1G) (Fig. 2, F–H), which are known tRNA modifications and were present in the metabolic extracts at high levels (Fig. S2G) (47). We did not include the isotopologue analysis of m^5C in the time-series experiments due to low abundance of its free ribonucleosides and due to low MS sensitivity for m^5C (Figs. 2D and S2, D and G), which compromises the quantification under conditions of partial ^{13}C labeling. Thus, while the turnover analysis of the polyA+ fraction could not be used to validate lowly abundant modifications due to rRNA contamination (Fig. 2E), the turnover of A_m and m^6_2A in the free pool suggests that these modifications indeed partly derive from rapidly decaying RNAs (Fig. 2F), which may include non-ribosomal RNAs. This was supported by the normalized levels of A_m and m^6_2A in the free pool being more similar to those of polyA+ than rRNA (Fig. S2H). These results confirm the presence of A_m in mRNA (48) and suggest that m^6_2A is more common in short-lived RNAs than m^1A , which has already been studied in mRNAs (49, 50).

Sensitivity of RNA modifications to metabolic stresses

The maintenance of methylation reactions requires SAM and serine/glycine, which feed one-carbon metabolism and SAM synthesis (51). Conversely, RNA demethylation is catalyzed by α -ketoglutarate (α -kg)-dependent dioxygenases such as FTO and ALKBH5, which act on m^6A , N6,2-O-dimethyladenosine (m^6A_m), and m^1A (52, 53). We measured both the methylation turnover and abundance of modified ribonucleosides to obtain insights into the sensitivity of mRNA *versus* ncRNA to metabolic stresses linked to RNA (de)methylation. First, we confirmed that actinomycin (ActD), a pan inhibitor of eukaryotic transcription, completely inhibited the methylation turnover of polyA+ and large RNA (Fig. 3A), in accordance with the cotranscriptional deposition of mRNA and most rRNA modifications (54–57). Treatment with ActD decreased the normalized levels of m^6A (m^6A/A) in polyA+ RNA (Fig. 3B), but not of m^7G , which supports the role of m^6A in global mRNA destabilization, as initially reported (40). Deprivation of serine or glutamine inhibited mainly the methylation turnover of large RNA (Fig. 3, C and D). In contrast, the abundance of modified ribonucleosides in large RNA was unaltered under these forms of stress (Fig. 3E), whereas glutamine deprivation led to increased m^6A levels in polyA+ RNA (Fig. 3F), which was in contrast to ActD treatment (Fig. 3B). Thus, the reduced methylation turnover of large RNA simply results from an inhibited transcription of the rRNAs themselves, while the altered abundance of m^6A without significant changes in its turnover indicates its post-transcriptional lability to stress conditions *via* altered (i) decay of m^6A -enriched *versus* m^6A -depleted mRNAs or (ii) dynamics of the m^6A modification itself. These data show that measurements of methylation turnover and ribonucleoside

methyladenosine; m^1G , 1-methylguanosine; m^5C , 5-methylcytidine; m^6A , N6-methyladenosine; m^6_2A , N6,N6-dimethyladenosine; m^7G , 7-methylguanosine; SSE, sum of squared errors.

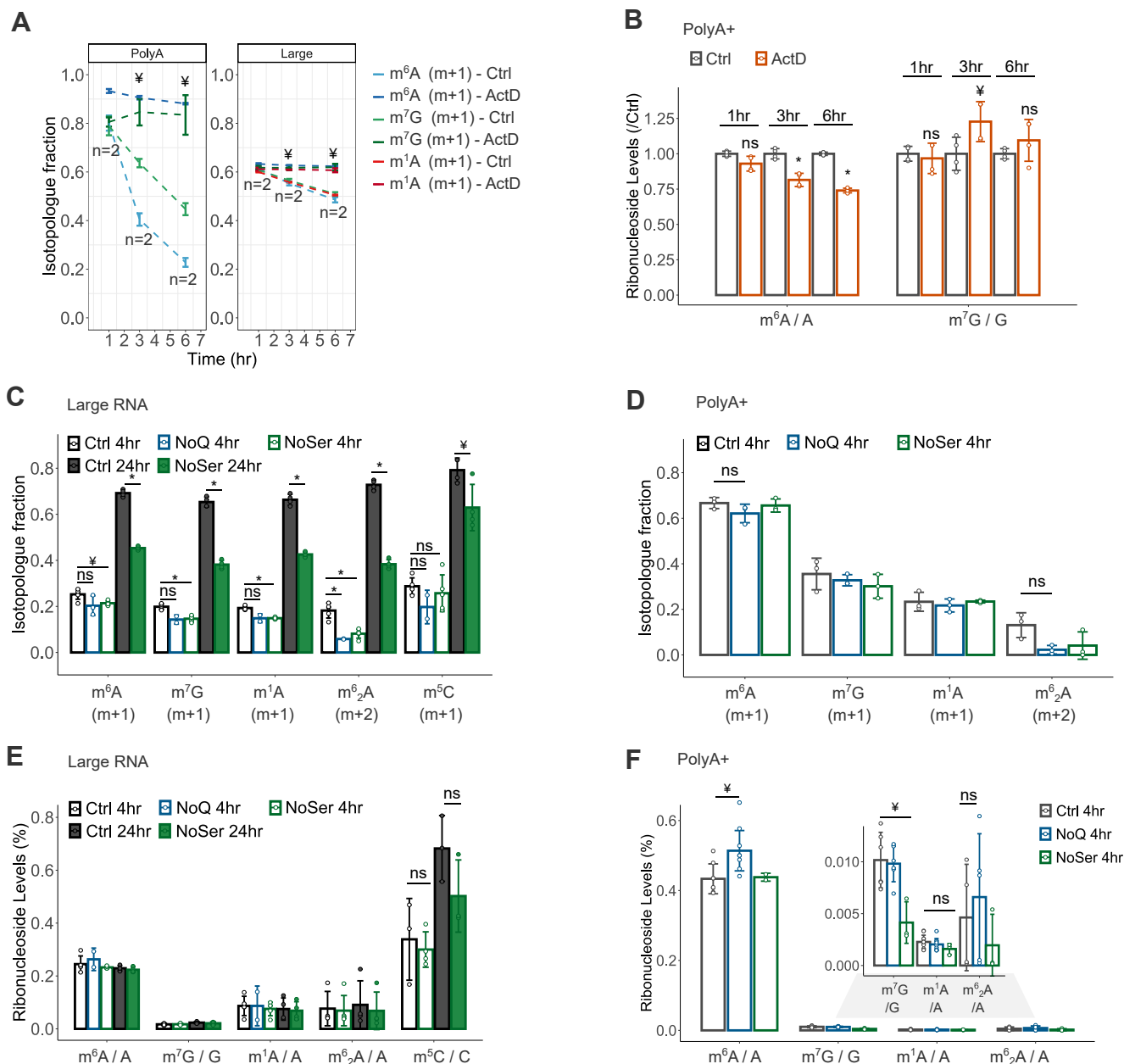


Figure 3. Dynamics of RNA modifications under stress conditions. A, isotopologue fractions of modified ribonucleosides in polyA+ and large RNA during a ^{13}C -dynamods chase experiment in 786O cells cultured with complete DMEM medium (ctrl) or DMEM supplemented with actinomycin D (ActD, 10 μM); dashed lines connect data points. B, normalized levels of m^6A (m^6A/A) and m^7G (m^7G/G) in polyA+ RNA derived from ctrl and ActD-treated 786O cells, normalized relative to the ctrl data points in the chase experiment as indicated. C and D, isotopologue fractions of modified ribonucleosides in large RNA (C) and polyA+ RNA (D) following [^{13}C -methyl]-methionine labeling under control (ctrl), glutamine (NoQ), or serine (NoSer) deprived conditions. E and F, normalized levels of modified ribonucleosides in large RNA (E) and polyA+ RNA (F) in 786O cells cultured with ctrl, NoQ or NoSer DMEM medium for the times indicated. Error bars represent standard deviation of three or more biological replicates, with exception of (A), the NoQ condition in (C and E) and m^6A under NoSer (D and F) (two biological replicates). In all cases, each replicate is the average of two technical replicates. * denotes $p < 0.005$, $\dagger$ denotes $p < 0.05$, ns denotes not significant, of a two-sided Student's t test comparing stress to ctrl samples, as indicated.

abundances resolve transcriptional *versus* posttranscriptional effects on RNA modifications in nonstationary (stimulus-dependent) experiments.

Discussion

Isotopic labeling of cultured cells is a well-established method to quantify metabolic activity, but its application to

RNA modifications has been limited to ncRNAs (33, 58). Here, we demonstrate that quantification of methylation turnover with dynamic ^{13}C labeling informs on the distinct dynamics of polyA+ and ncRNA modifications and their sensitivity to metabolic perturbations of mammalian cells. ^{13}C -dynamods presents several advances in the application of MS approaches to study RNA turnover and dynamics. First, in contrast to approaches that rely on prior/in-parallel labeling with

$^{15}\text{N}/^{13}\text{C}$ -enriched nucleosides to distinguish between preexisting and newly modified RNA (33, 58), the “heavy” isotopologue fraction derived from [^{13}C -methyl]-methionine is internally controlled for the amount of unlabeled, preexisting ribonucleosides and is thus specific to modified ribonucleosides. Second, in contrast to approaches using multiplexing or spiking from isotopically labeled cultures (24, 25, 32, 58), the dynamic ^{13}C labeling over time directly informs on the turnover frequency (in h^{-1}) (41, 43) of various RNA modifications, as the direct substrate (SAM) of the targeted reaction (RNA methylation) is close to 100% labeled in less than 30 min in cell culture (Fig. S1B) (43). Moreover, SAM need not be fully labeled for comparisons of dynamic ^{13}C labeling between modifications at early, single time points as its ^{13}C enrichment is expected to be equal across SAM-dependent RNA modifications. Third, conventional RNA labeling approaches using ribonucleoside analogues or precursors rely on their incorporation into newly synthesized RNA through the salvage pathway (e.g., uridine and its analogs) (45, 59) or the *de novo* pathway (glucose, serine/glycine, or water tracers) of ribonucleotide (NTP) synthesis (60–64). In contrast, the methylation of newly transcribed RNA occurs downstream (cotranscriptionally) of NTP synthesis, making it a reliable readout of RNA turnover, particularly under nonstationary conditions that may alter the relative activity of NTP synthesis pathways (65, 66). Also, the nucleotide recycling through the salvage pathway may lead to ineffective “washout” with unlabeled nucleotides in chase experiments (59, 66), whereas a new SAM molecule is used in each (cotranscriptional) methylation cycle in ^{13}C -dynamods, adding to the specificity and versatility of experimental designs (59). Finally, combined measurements of methylation turnover and ribonucleoside levels allow ^{13}C -dynamods to also inform on transcriptional *versus* post-transcriptional sensitivity of RNA modifications in response to a stimulus.

Reliable study of mRNA modifications by MS analyses of ribonucleosides is challenging due to the high abundance of heavily modified ncRNAs that invariably contaminate the polyA+ fraction (36, 37). To address this challenge, we exploited the fact that the life cycle of methylated RNA follows the life cycle of the RNAs at metabolic steady state (38–40). The decay of mRNA is fast with a median half-life from 40 min to 9 h in mammalian cells (44, 67, 68), while rRNA and tRNA exhibit half-lives of 60 to 70 h in growing fibroblasts (44). By definition, the methylation turnover as determined by ^{13}C incorporation into newly methylated RNA does not assess nonmodified RNAs and hence does not capture the removal of RNA modifications. That is, any RNA demethylation activity can only affect the abundances of modified ribonucleosides as a whole (sum of $m + 1$ and $m + 1$), not the $m + 1$ or $m + 0$ isotopologues individually, which would be required to cause a differential $m + 1$ fraction. Since RNA transcription and decay rates are constant at metabolic steady state, we propose that the methylation turnover is an accurate readout of RNA turnover, which allowed us to assess the contributions of short-lived (mRNAs) and long-lived (noncoding) RNAs to a pool of modified RNAs and free ribonucleosides. Indeed, this

information could not be drawn if labeled ribonucleoside themselves or its precursors were used to trace ribonucleotide synthesis (*via* the salvage and *de novo* pathways) or transcription. Likewise, the source RNA cannot be pinpointed if the abundance (levels) of modified ribonucleosides is measured in isolation.

Using this approach, we could not reliably determine highly abundant rRNA modifications ($\text{Am}/m^1\text{A}/m^6_2\text{A}$) in polyA+-purified RNA as being derived from mRNAs. Based on methylation turnover, we estimated that these modifications in the polyA+ fraction could be detected with a maximal ncRNA contamination of 0.5 to 2.9%, since $\text{Am}/m^1\text{A}/m^6_2\text{A}$ levels were found higher in ncRNAs by a factor of 17–37–97, respectively. In contrast, a ~6% of ncRNA contamination would be required to fully account for the slower $m^7\text{G}$ turnover detected in polyA+ RNA, since $m^7\text{G}$ levels were found only approximately nine times higher in ncRNAs. Since the $m^7\text{G}$ turnover reflected that of a *bona fide* mRNA modification, its slower kinetics in polyA+-purified RNA suggests a temporal delay relative to $m^6\text{A}$ turnover. A further evidence for the distinct turnover of $m^7\text{G}$ relative to $m^6\text{A}$ in mRNAs is the response to metabolic stresses, where the $m^7\text{G}$ turnover changed in large RNA but not polyA+ RNA following 4 h of ^{13}C labeling (Fig. 3, C and D). In this respect, while $m^6\text{A}$ is cotranscriptionally deposited in mRNA (39, 54, 56), $m^7\text{G}$ is an essential modification at the 5' cap of mRNA that can be placed in both the nucleus and cytoplasm (69, 70). Moreover, $m^7\text{G}$ sites have been reported internally within mRNA, catalyzed by the cytosolic METTL1 methyltransferase (71). Taken together, the distinct kinetics of $m^7\text{G}$ and $m^6\text{A}$ in polyadenylated RNA suggest compartmentalization differences affecting their temporal deposition into newly synthesized mRNA.

Previous reports identified $m^5\text{C}$ and Am as mRNA modifications (48, 72) and more recently also $m^1\text{A}$ (49, 50). Our analysis of free ribonucleosides showed a similarly fast kinetics for the turnover of Am, $m^6\text{A}$ and $m^7\text{G}$, consistent with these modifications being derived from the rapidly decaying mRNAs. This was supported by the normalized levels of free modified ribonucleosides being more similar to those of polyA+-purified than of ncRNAs. In contrast, the turnover of free $m^1\text{A}$ and $m^1\text{G}$ ribonucleosides was significantly slower, indicating that these modifications are predominantly derived from an ncRNA class, likely tRNAs, where they are present at high levels (47). This aligns with the recent conclusions that $m^1\text{A}$ might be restricted to a handful of mRNAs (3, 49). Finally, despite the low sensitivity of $m^5\text{C}$ detection in chase experiments, the quantification of $m^5\text{C}$ isotopologues at 24 h is consistent with it being derived from mRNA (72).

The $m^6_2\text{A}$ modification is thought to be present primarily in the 18S and 12S rRNA of mammalian transcriptomes (26, 73, 74). While $m^6_2\text{A}$ has been detected in bacterial tRNA (75), care must be taken with abundant $m^6_2\text{A}$ -modified rRNA fragments that copurify with tRNA (76), and are not resolved by standard MS quantifications. Here, we detected slower turnover of $m^6_2\text{A}$ in mammalian small RNAs relative to large RNA, raising the possibility that this modification is present in non-18S/12S RNA species, e.g., in small rRNA and/or tRNAs. Nevertheless,

an unlikely possibility from ^{13}C -dynamods measurements remains that m^6_2A -containing 18S/12S rRNA fragments are turned over at a slower rate than m^6_2A -containing intact rRNAs. Interestingly, the turnover of free m^6_2A ribonucleosides exhibited a faster kinetics than free $\text{m}^1\text{A}/\text{m}^1\text{G}$ (canonical ncRNA modifications) (47) and large RNA modifications, suggesting that m^6_2A is also present in high turnover RNAs, which likely include short-lived ncRNAs and mRNAs. The presence and role of m^6_2A beyond intact 18S/12S rRNAs thus merit further investigation. Of note, single methylation intermediates of m^6_2A have been detected *in vitro* (77), but we did not detect $\text{m} + 1$ isotopologues for m^6_2A above its natural abundance. As SAM was 98 to 100% ^{13}C -labeled within 30 min, the time resolution of our experiments does not capture sequential methylation of adenosine into m^6_2A . These various findings highlight the value examining the turnover of methylated RNA to uncover the presence of modifications in uncharacterized in RNA subclasses, which warrants investigation.

A nonstationary condition, *e.g.*, a stress condition, may affect the afferent (*i.e.*, methylation of newly transcribed RNA) or efferent (*i.e.*, decay of methylated RNAs) turnover parts of methylated RNAs. Combined with abundance measurements, this allowed us to assess transcriptional *versus* nontranscriptional sensitivity of modified RNAs to stress conditions. This was evidenced by the reduced methylation turnover of ncRNAs following serine deprivation and ActD treatment, wherein the abundance of modifications was unaltered in the former (transcriptional effect) but m^6A levels decreased in the latter (posttranscriptional effect). The observed inhibited turnover of methylated ncRNAs under serine deprivation is in line with the expected inhibition of mTOR activity and thereby of rRNA biogenesis (78). In contrast to ncRNAs, glutamine deprivation increased m^6A levels in polyA⁺-purified RNA without significantly changing (not increasing) its turnover. These data suggest that m^6A is posttranscriptionally sensitive to glutamine levels through either (i) an altered decay of m^6A -enriched *versus* m^6A -depleted RNAs or (ii) inhibition of m^6A demethylation itself. The demethylation of m^6A is a stepwise conversion into adenosine through the formation of N^6 -hydroxymethyladenosine (hm^6A) and N^6 -formyladenosine (f^6A) intermediates (79), whose presence or turnover was not assessed here. Thus, related developments are needed to address if posttranscriptional dynamics of m^6A by metabolic stress conditions are due to RNA demethylation or differential decay of m^6A -modified *versus* m^6A -depleted RNAs. As glutamine is the main carbon source of α -ketoglutarate, a cosubstrate of RNA demethylases (52, 53), it is plausible that its depletion could inhibit RNA demethylation. This supports the notion that the reversibility of m^6A in mRNA is likely context-dependent (80). With exception of ActD treatment, the stimuli examined did not lead to changes in both the turnover and abundances of modified ribonucleosides. In this respect, while altered methylation turnover (and unaltered abundance) indicates a transcriptional effect, it is plausible that a stimulus may alter methylation turnover at the posttranscriptional level, *e.g.*, if the newly methylated ($\text{m} + 1$

fraction) and preexisting methylated ($\text{m} + 0$ fraction) RNAs differently are exposed differently to RNA degradation, *e.g.*, through nuclear/cytosolic compartmentalization.

Our study demonstrates how quantification of methylation turnover and abundance can be used to examine the presence of RNA modifications in RNA classes, their temporal dynamics, and sensitivity to stress conditions. These insights open new directions to be further explored by MS and orthogonal approaches to obtain information on particular RNAs.

Experimental procedures

Cell culture and metabolic labeling

786O cells were obtained from the Crick Cell Services and cultured at 37 °C with 5% CO_2 in high-glucose DMEM medium (Thermo Fisher Scientific, #61965026) supplemented with 10% fetal bovine serum (FBS) (Thermo Fisher Scientific, #21875034). For ^{13}C labeling experiments, 786O cells were grown in high-glucose DMEM medium without glutamine, methionine, and cystine (Thermo Fisher Scientific, #21013024) supplemented with 10% dialyzed FBS (Thermo Fisher Scientific, #26400044) to minimize contributions of unlabeled amino acids typically present in regular FBS, 2 mM glutamine, 0.1 mM cystine, and 0.2 mM [^{13}C -methyl]-methionine (CK Isotopes Limited). Cells were maintained at 50 to 60% confluence (or 30–40% in chase experiments) and washed with PBS before switching to ^{13}C -labeled medium (or regular medium in chase) for the indicated time periods.

RNA purification

At the conclusion of metabolic labeling, the medium was aspirated and extraction of total RNA was performed with the mirVana isolation kit according to manufacturer's instructions (Thermo Fisher Scientific, #AM1560). Large (>200 nt) RNAs were purified by adding 1/3 volume of 100% ethanol to the aqueous phase recovered from the organic extraction before loading into the filter cartridge of the mirVana Kit. Small (<200 nt) RNAs were purified by collecting the total filtrate, addition of 2/3 volume of 100% ethanol, and loading into the filter cartridge. Polyadenylated RNA was purified from total/large RNA *via* two rounds of polyA tail hybridization with Oligo-dT magnetic Dynabeads (Thermo Fisher Scientific, #61002).

LC-MS/MS analysis of ribonucleosides

Purified RNA (100–250 ng) was digested into ribonucleosides using one unit of nuclease P1 (Sigma-Aldrich, #N8630-1VL) in 25 μl of buffer 25 mM NaCl, 2.5 mM ZnCl_2 , and 10 mM NaCH_3COO pH 5.3 and incubated for 2 h at 37 °C. Subsequently, NH_4HCO_3 (100 mM) and 5 units of alkaline phosphatase (CIP) (NEB, #M0525S) were added and the sample incubated for 2 h (or 20 min, with Quick CIP) at 37 °C. Formic acid was added at 0.1% v/v in a final volume of 50 μl , samples were filtered (0.22 μm , Millipore) and 15 to 20 μl analyzed in duplicate by LC-MS. Ribonucleosides were resolved with a C18 reverse-phase column (100 $\times$ 2.1 mm, 3 μm particle size, Chromex Scientific, #F18-020503) and eluted with a gradient of 0.1% v/v formic acid (solvent A) and

80% acetonitrile in 0.1% formic acid (solvent B) at a flow rate of 0.2 ml/min and 40 °C: 100% solvent A for 3 min, 12% solvent B for 12 min, and 100% solvent B for 2 min after which the column was re-equilibrated with 100% solvent A for 3 min (20 min total run time). The ribonucleoside separation was performed using U3000 HPLC (Thermo Scientific) and the detection by a TSQ Quantiva Triple Quadrupole mass spectrometer (TSQ Quantiva, Thermo Scientific) controlled by the Xcalibur software version 4.0.27 (Thermo Scientific). The HPLC was coupled to the TSQ Quantiva using a HESI (heated electrospray) ion source (Thermo Scientific) operating in positive ionization mode with the following parameters: capillary voltage, 3500 V; sheath gas flow, 7.35 l/min; gas temperature, 325°C. The first and third quadrupoles (Q1 and Q3) were stringently fixed to 0.2 units of resolution and set to detect the mass of the precursor ribonucleoside ion (Q1) and of the base and ribose product ions (Q3). The ribonucleosides were identified by comparison of the retention time and detected mass transitions to commercially available standards. The collision energies were experimentally defined based on the fragmentation pattern of each ribonucleoside standard and chosen based on the maximum intensity of the base product; the ribose ring was used only as a qualifying transition. The retention time, mass transitions (m/z), and collision energies of each ribonucleoside were: adenosine, ~4.1 min, 268.1 $\rightarrow$ 136.1 m/z , 20 V; guanosine, ~5.4 min, 284.1 $\rightarrow$ 135.0 m/z , 35.5 V; cytidine, ~1.4 min, 244.2 $\rightarrow$ 112.05 m/z , 12 V; m^6A , ~8.7 min, 282.1 $\rightarrow$ 150.1 m/z , 20 V; m^1A , ~1.8 min, 282.1 $\rightarrow$ 150.1 m/z , 20 V, m^7G , ~2.3 min, 298.05 $\rightarrow$ 166.1 m/z , 20 V; m^5C , ~1.8 min, 258.2 $\rightarrow$ 126.1 m/z , 13 V; m^6_2A , ~11.9 min, 296.2 $\rightarrow$ 164.1 m/z , 22 V. Each mass transition above corresponded to the $m + 0$ isotopologue and increased by one ($m + 1$), two ($m + 2$), and three ($m + 3$) units for detection of the other isotopologues, e.g., m^6_2A : 297.2 $\rightarrow$ 165.1 ($m + 1$), 298.2 $\rightarrow$ 166.1 ($m + 2$), 299.2 $\rightarrow$ 167.1 ($m + 3$). The dwell time for each transition was 30 ms for a duty cycle of 930 ms (31 transitions), and 8 to 20 data points per chromatographic peak were obtained for “short” and “long” peaks, respectively. A mix of ribonucleoside standards containing 0.5, 1, 5, 10, 50, 100, 500 fmol, 1, 5, 10, 50, or 100 pmol of each ribonucleoside was run in parallel after the biological samples for absolute quantifications, a subset of which is shown in Fig. S2D. Data were recorded using the Xcalibur 3.0.63 software (Thermo Fisher Scientific) and analyzed using Skyline (version 19.1) (81) (Supplementary Materials).

Metabolite extraction and LC-MS analysis of SAM/free ribonucleosides

At the end of cell culture with [^{13}C -methyl]-methionine, metabolic activity quenched by adding ice-cold PBS. Metabolites were extracted by addition of 600 μ l ice-cold 1:1 (vol/vol) methanol/water to the cell pellets, samples were transferred to a chilled microcentrifuge tube containing 300 μ l chloroform and 600 μ l methanol (1500 μ l total, in 3:1:1 vol/vol methanol/water/chloroform). Samples were sonicated in a water bath for 8 min at 4 °C and centrifuged (13,000 rpm) for 10 min at 4 °C. The supernatant containing the extract was transferred to a

new tube for evaporation in a speed-vacuum centrifuge, resuspended in 3:3:1 (vol/vol/vol) methanol/water/chloroform (350 μ l total) to phase separate polar metabolites (upper aqueous phase) from nonpolar metabolites (lower organic phase), and centrifuged. The aqueous phase was transferred to a new tube for evaporation in a speed-vacuum centrifuge and resuspended in 100 μ l water for LC-MS acquisition. LC-MS analysis was performed using a Dionex UltiMate LC system (Thermo Fisher Scientific) with a ZIC-pHILIC column (150 mm $\times$ 4.6 mm, 5 μ m particle, Merck Sequant), as described previously (82). A 15 min elution gradient of 80% Solvent A (20 mM ammonium carbonate in Optima HPLC grade water, Sigma Aldrich) to 20% Solvent B (acetonitrile Optima HPLC grade, Sigma Aldrich) was used, followed by a 5 min wash of 95:5 Solvent A to Solvent B and 5 min re-equilibration. Other parameters were as follows: flow rate, 300 μ l/min; column temperature, 25 °C; injection volume, 10 μ l; autosampler temperature, 4 °C. All metabolites were detected across a mass range of 70 to 1050 m/z using a Q Exactive Orbitrap instrument (Thermo Fisher Scientific) with heated electrospray ionization and polarity switching mode at a resolution of 70,000 (at 200 m/z). MS parameters were as follows: spray voltage 3.5 kV for positive mode and 3.2 kV for negative mode; probe temperature, 320 °C; sheath gas, 30 arbitrary units; auxiliary gas, five arbitrary units. Parallel reaction monitoring (PRM) was used at a resolution of 17,500 to confirm the identification of metabolites; collision energies were set individually in HCD (high-energy collisional dissociation) mode. Data were recorded using the Xcalibur 3.0.63 software and analyzed using Tracefinder 4.1 (ThermoFisher Scientific) according to the manufacturer's workflows.

Quantification of methylation turnover

The isotopologue fractions were defined as the total ion counts of the $m + 1$ isotopologue (except for $m + 2$ in m^6_2A) relative to the total ion counts of the $m + 0$ plus $m + 1$ isotopologues. The kinetics of isotopologue fractions and goodness of fit were determined using the Curve Fitting toolbox of Matlab R2020a (MathWorks) either by a linear regression [$f(x) = p_1 \cdot x + p_2$] or an exponential fit: a one-term function in the chase experiments [$f(x) = a \cdot \exp(b \cdot x)$, Levenberg–Marquardt algorithm], and a two-term function to fit the isotopologue fractions of free ribonucleosides [$f(x) = a \cdot \exp(b \cdot x) + c \cdot \exp(d \cdot x)$, Levenberg–Marquardt algorithm].

Data availability

The mass spectrometry.raw data and Skyline files pertaining Figure 1 and Fig. S1 are made available in the Supplementary Materials. All remaining mass spectrometry data can be obtained from the authors upon request.

Supporting information—This article contains supporting information.

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Abbreviations—The abbreviations used are: f⁶A, N⁶-formyladenosine; FBS, fetal bovine serum; hm⁶A, N⁶-hydroxymethyladenosine; m⁶A, N⁶-methyladenosine; m⁷G, 7-methylguanosine; m⁶₂A, N⁶,N⁶-dimethyladenosine; MS/MS, tandem mass spectrometry; NGS, next-generation sequencing; rRNA, ribosomal RNA; SAM, S-adenosylmethionine.

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