Reconsidering Movement of Eukaryotic mRNAs between Polysomes and P Bodies

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Reconsidering Movement of Eukaryotic mRNAs between Polysomes and P Bodies

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SUMMARY

Cell survival in changing environments requires appropriate regulation of gene expression, including posttranscriptional regulatory mechanisms. From reporter gene studies in glucose-starved yeast, it was proposed that translationally silenced eukaryotic mRNAs accumulate in P bodies and can return to active translation. We present evidence contradicting the notion that reversible storage of non-translating mRNAs is a widespread and general phenomenon. First, genome-wide measurements of mRNA abundance, translation, and ribosome occupancy after glucose withdrawal show that most mRNAs are depleted from the cell coincident with their depletion from polysomes. Second, only a limited subpopulation of translationally repressed transcripts, comprising fewer than 400 genes, can be reactivated for translation upon glucose readdition in the absence of new transcription. This highly selective posttranscriptional regulation could be a mechanism for cells to minimize the energetic costs of reversing gene-regulatory decisions in rapidly changing environments by transiently preserving a pool of transcripts whose translation is rate-limiting for growth.

INTRODUCTION

Cells respond to changing environments by regulating gene expression. Regulation can occur at the level of transcription and/or posttranscriptionally during processes including premRNA splicing, messenger RNA (mRNA) export, translation and mRNA decay. In some embryonic cells, gene regulation during early development is entirely posttranscriptional and involves temporally and spatially controlled translation of maternally deposited mRNAs (Johnstone and Lasko, 2001; Richter, 1991). More typically, cells employ a combination of transcriptional and posttranscriptional regulatory strategies. The logical and mechanistic relationships between transcriptional and posttranscriptional regulation are poorly understood, if indeed such relationships exist.

Various hypotheses have been proposed for the role of translational regulation in contexts where transcriptional regulatory mechanisms are also active. For example, translational activation of preexisting mRNAs can produce new protein faster than transcriptional activation of the same genes, and may therefore be important in situations that demand rapid responses. In addition, translational mechanisms can control where proteins are produced within the cell. Furthermore, translational regulation has been suggested to act globally as an amplifier of the effects of transcriptional gene control, increasing the protein output from transcriptionally induced genes and further decreasing the protein output from transcriptionally repressed genes (Melamed et al., 2008; Preiss et al., 2003). On the other hand, translational attenuation has also been proposed to act as a global dampener of transcriptional noise in gene expression (Blake et al., 2003; Ozbudak et al., 2002; Raser and O’Shea, 2005).

We set out to determine the relationship between the programs of transcriptional and translational response to stress. We further sought to determine the biological logic behind selection of specific mRNAs for translational regulation and the molecular differences between genes controlled at transcriptional versus translational levels. The glucose starvation response in yeast is an appropriate model system because glucose withdrawal induces widespread changes in both transcription and translation. Transcriptional changes are mediated by well-characterized signaling pathways and transcription factors (Zaman et al., 2008). Translation activity changes by an incompletely understood mechanism requiring genes that have been variously implicated in deadenylation-dependent mRNA decapping and decay, mRNA subcellular localization, and the formation of translationally repressed mRNP complexes (Ashe et al., 2000; Brengues et al., 2005; Coller and Parker, 2004, 2005; Holmes et al., 2004; Teixeira et al., 2005). In response to glucose starvation, yeast initiate a cellular differentiation program known as haploid invasive growth, which is thought to function as a cellular foraging response (Cullen and Sprague, 2000). Because this cellular adaptive response to glucose starvation requires new protein synthesis, the “global” repression of translation must either be short-lived or in fact affect only a subset of genes.

Here we used DNA microarrays to investigate changes in mRNA abundance, translation activity and ribosome occupancy after glucose withdrawal. We made measurements genome-wide of mRNA abundance, translation, and ribosome occupancy in glucose-starved cells compared to control cultures. Overall, we have found that transcriptional and translational regulation are mechanistically and functionally distinct, and that transcriptional repression in response to glucose withdrawal is a more widespread and general phenomenon than previously proposed.
during a 2 hr time course after glucose withdrawal. We found that the view of “global” translational repression is overgeneralized. While “bulk” translation was greatly reduced, hundreds of newly transcribed mRNAs associated with polysomes within ten minutes of glucose withdrawal. Functionally coherent groups of genes were coregulated at the posttranscriptional as well as transcriptional level. Using computational approaches, we related gene-specific posttranscriptional changes to underlying mRNA properties by exploiting recent genome-wide studies of yeast mRNA characteristics including abundance, half-life, translational efficiency, poly(A) tail length and association with various RNA-binding proteins. Following a lead generated by this analysis, we examined whether all genes or only a specific subpopulation of genes are capable of returning to active translation in the absence of new transcription. In contradiction of the prevailing model in the field, we found that the capacity for translational reactivation is narrowly restricted to a limited subset of mRNAs. Transient preservation of these select mRNAs, whose translation is rate limiting for growth in rich media, could act as a buffer to minimize the fitness costs associated with “false alarms” caused by transient depletion of glucose or by noise in glucose-sensitive signaling pathways.

RESULTS

We studied wild-type invasive-growth competent yeast subjected to acute glucose starvation. Cells were starved for 0, 10, 20, 30, 60, or 120 min before processing. For each time point, polysome profiles were generated to monitor global translation. In agreement with previous reports (Ashe et al., 2000; Kuhn et al., 2001), a collapse of the polysome region and concomitant increase in the 80S monosome peak occurred after 10 min of glucose withdrawal, indicating a bulk reduction in translation initiation. Previous investigations of the translational response to acute glucose withdrawal with either bulk measurements (Ashe et al., 2000; Holmes et al., 2004) or examination of a few genes (Brengues et al., 2005) suggested that little translation initiation occurs 10–20 min after glucose removal. In light of prior studies of glucose-stimulated transcriptional regulation, as well as genetic evidence that many of the genes induced upon glucose depletion are required for growth in the absence of glucose, we reasoned that such transcripts must at some time associate with polysomes in glucose-starved cells (Zaman et al., 2008). We extended the time course of the experiment to learn when bulk translation recovers as cells adapt to growth in low glucose concentrations. Polysomes remained low between 10 and 30 min after glucose withdrawal and showed signs of partial recovery after 60 min, with further increases by 120 min (Figure 1A).

Changes in Polysomal mRNA Levels Closely Parallel Changes in Transcript Abundance

To examine the kinetics of gene-specific translation activity after glucose withdrawal, we isolated RNA from polysome fractions and total cell lysates and prepared fluorescently labeled complementary DNA (cDNA) for competitive hybridization on custom microarrays. In order to determine relative changes in translation activity, mRNA abundance, and the fraction of total mRNA associated with polysomes (ribosome occupancy), we performed the following comparisons for each time point after glucose withdrawal: polysomal RNA starved with polysomal RNA mock-starved (P/Px), total RNA starved with total RNA mock-starved (Tx/Tx), and polysome starved x min with total starved x min (Px/Tx), respectively (Figure 1B). All experiments were performed twice (biological replicates), with each RNA sample processed in duplicate (technical replicates). “Noisy” genes for which mRNA abundance or polysome association varied by >1.5-fold between biological replicate experiments in unstarved cells were omitted from further analysis (495 genes). Reproducible results were obtained for 5,590 genes.

Glucose withdrawal led to changes in the relative mRNA abundance of hundreds of genes within 10 min, consistent with prior studies of carbon source-mediated regulation of yeast transcription (Zaman et al., 2008). These changes persisted and were amplified over the course of 120 min of starvation. Our data do not distinguish between transcriptional induction/repression and mRNA stabilization/destabilization as the mechanism responsible for changing total mRNA abundance. Both mechanisms likely contribute. Many of the genes that showed relatively increased mRNA levels following glucose withdrawal are known targets of glucose regulated transcription factors (Figure S1 available online). For most genes, changes in their relative mRNA abundance in polysomes (P/P) mirrored changes in overall mRNA levels (T/T) (Figure 1C). Notably, for the more than 1,000 genes whose relative mRNA abundance increased by 2-fold or more after glucose withdrawal, relative polysomal abundance similarly increased. Furthermore, most induced genes did not show a noticeable lag between the increase of mRNA in the total RNA pool and appearance in the polysomal fraction. Thus, despite the reduction in the rate of “bulk” protein synthesis, translation initiation occurred on transcriptionally upregulated mRNAs as early as 10 min after glucose withdrawal. Reduction in polysomal mRNA closely paralleled reduction in total mRNA levels for almost all downregulated genes. These data contradict the model that glucose withdrawal leads to widespread sequestration of mRNAs in stable translationally repressed mRNP (Brengues and Parker, 2007; Brengues et al., 2005; Hoyle et al., 2007; Teixeira et al., 2005).

Unsupervised hierarchical clustering revealed small groups of genes that appear to be regulated primarily at the posttranscriptional level: cluster I genes showed relatively reduced polysomal mRNA levels despite increased total mRNA abundance due to low ribosome occupancy; conversely, cluster II genes had relatively unaffected polysomal mRNA levels despite reduced total mRNA abundance due to high ribosome occupancy (Figure 1D). Nevertheless, there was strong overall agreement between the relative increase or decrease of an mRNA in the cell at any time point after glucose withdrawal (T/T) and the change in polysome association at that time point (P/P). In contrast, ribosome occupancy (P/T) at any time point was more highly correlated with ribosome occupancy at other times than with either P/P or T/T at the same time point after glucose withdrawal (Figure 1E), suggesting that it is largely an intrinsic property of each mRNA, although some starvation-induced changes in P/T were observed. At the global scale,
fold changes in polysomal mRNA levels were somewhat compressed compared to changes in total mRNA levels (Figure 1F and Figure S2). Thus, our data do not indicate widespread “potentiation,” whereby changes in total mRNA levels are amplified by homodirectional changes in translation efficiency, as was suggested in studies of yeast subject to rapamycin treatment or heat shock (Preiss et al., 2003). Stress specificity of “potentiation” was previously noticed in comparison of the translational responses to amino acid starvation and butanol stress (Smirnova et al., 2005).

See also Figures S1 and S2 and Table S2.

Figure 1. Regulation of Transcription and Translation in Glucose-Starved Cells

(A) Polysome profiles of yeast starved for glucose by transfer from YPD to YP media.

(B) Schematic of the microarray comparisons performed.

(C) Time-resolved gene expression profiles resulting from the comparisons shown in (B). Ratio values are indicated by color scale and represent the average of eight measurements (two biological replicates x two technical replicates x two probes per gene). Rows (genes) are ordered according to the results of hierarchical clustering based on Euclidean distance with P/T values assigned twice the weight of P/P or T/T.

(D) Unusual gene clusters for which polysomal mRNA abundance diverged from total mRNA abundance.

(E) P/T ratios are more similar to each other than to T/T or P/T for any time point. Branch lengths represent Spearman rank correlation coefficients between data columns from (C).

(F) Total mRNA abundance changes parallel and exceed polysomal mRNA abundance changes after 60 min minus glucose. Dotted line indicates best fit by linear regression.

See also Figures S1 and S2 and Table S2.
Relationships between Changes in Transcript Levels and Ribosome Occupancy

The complex groupings of genes produced by combining the analysis of transcriptional and posttranscriptional regulatory behavior using unsupervised hierarchical clustering did not readily reveal the logic underlying the relationship between the two modes of regulation. To investigate this relationship more directly, changes in total mRNA levels (T/T) and changes in ribosome occupancy (P/T) were analyzed separately using k-means clustering to identify groups of genes displaying similar behavior for each mode of regulation. Experimenting with various group numbers (k = 2–20) revealed that k = 7 for the T/T comparisons and k = 3 for the P/T comparisons gave robust solutions reflecting a reasonable compromise between preserving the complexity of the data and simplifying the subsequent analysis (Figure S3). Clustering genes by their ribosome occupancy (P/T) produced simple divisions into groups with high, low, and “neutral” P/T ratios (Figure 2A).

Although we anticipated that kinetic analysis of the total mRNA changes in response to glucose starvation would reveal temporal distinctions between genes, the groups of coregulated genes identified by k means clustering at k = 7 differed primarily by the magnitude of relative increase/decrease rather than by the timing of the maximal changes in gene expression (Figure 2A). Clustering with values of k > 12 did reveal kinetically distinct patterns of mRNA accumulation that are consistent with current understanding of transcriptional responses to glucose withdrawal (Figure S3). For example, genes subject to repression by Mig1 in the presence of glucose (e.g., SUC2 and CAT8) were derepressed within 10 min after glucose withdrawal, whereas known targets of the Cat8 transcription factor (e.g., PCK1, ICL1, and FBP1) accumulated mRNA strongly only after 120 min (Figure S1). Groups of genes clustered based on starvation-induced changes in total relative mRNA levels ranged from strongly induced (119 genes, median induction at t = 60 min of 19.7-fold) to strongly repressed (311 genes, median repression...
at t = 60 min of 11.9-fold). The use of multiple time points as well as both biological and technical replicates allowed the confident identification of genes displaying modest yet consistent changes in mRNA levels. The most weakly induced category includes more than 100 genes encoding mitochondrial proteins known to be important for growth in the absence of glucose, highlighting the potential biological significance of coordinated small changes in gene expression.

Changes in mRNA relative abundance and ribosome occupancy were not independent of one another ($X^2 = 179, p < 0.0001$ for T/T versus P/T). Starvation-induced genes were more likely to show low ribosome occupancy after glucose withdrawal, and repressed genes were more likely to show high ribosome occupancy (Figure 2B). Despite the statistical interdependence of changes in total mRNA levels and ribosome occupancy, for each group of genes having similarly induced/repressed total mRNA levels there were many genes displaying each of the possible posttranscriptional regulatory behaviors.

Functionally Distinct Groups of Genes Are Coregulated at the Posttranscriptional Level

To investigate the possibility that differences in posttranscriptional regulatory behavior are biologically significant, the function of genes in each category was examined by gene ontology (GO) analysis. Notably, the GO terms that were significantly enriched ($p < 0.01$ with Bonferroni correction for multiple hypothesis testing) for the seven mRNA abundance-based groups (highly induced, moderately induced, weakly induced, strongly repressed, moderately repressed, weakly repressed, and unchanged) segregated within these groups along posttranscriptional regulatory divisions. A complete list of significant GO terms for each of the 21 regulatory groups (seven T/T × three P/T) is provided in Table S1. Rarely were GO categories split between multiple post-transcriptional (P/T) regulatory groups, and where such a split occurred, the GO category spanned two out of three most similar groups—“high” and “neutral” or “low” and “neutral;” not “high” and “low.” This suggests that the posttranscriptional behavior of a gene is related to its biological function in the starved cell.

The posttranscriptional partitioning of functionally related genes may derive from mechanistic similarities in their gene expression pathways. For example, the nuclear-encoded mitochondrial protein genes showed consistently low ribosome occupancy despite the fact that these genes are transcriptionally upregulated in response to glucose withdrawal and encode proteins required for the cellular adaptation to low glucose conditions. mRNAs of some nuclear-encoded mitochondrial proteins are translated on cytosolic ribosomes that associate with mitochondria (Marc et al., 2002; Sylvestre et al., 2003). Subcellular localization of these mRNAs is driven by cis-acting elements in their 3′ untranslated regions (UTRs), and requires both the trans-acting RNA-binding protein Puf3 and the translocon of the mitochondrial outer membrane complex for full localization (Eliyahu et al., 2010; Saint-Georges et al., 2008). The apparent lag between the appearance of these transcriptionally induced mRNAs in the cell and their association with polysomes in our experiments could be explained by translational silencing of messages in transit from the nucleus to the periphery of mitochondria. In light of this explanation for the low P/T ratios of mRNAs encoding mitochondrial proteins, it is interesting to consider whether other transcriptionally induced yet poorly translated mRNAs identified in our analysis might also be subject to localization-dependent translational control.

The most striking example of gene function partitioning according to posttranscriptional regulatory behavior was the separation of cytoplasmic ribosomal protein genes (RPGs) from ribosome biogenesis factors (RBGs) (Figures 2A and 3 and Figure S4). Genes from both functional groups were moderately to strongly reduced in both the total and polysomal mRNA pools after glucose withdrawal. This downregulation is likely due to the greatly reduced demands for new ribosome synthesis as the cells transition from rapid growth and division, requiring the assembly of ~200,000 new ribosomes every 90 min (Warner, 1999), to cellular differentiation and slower growth in the invasive filamentous form (Cullen and Sprague, 2000). The two groups diverged in their posttranscriptional responses to glucose starvation. The P/T ratios of RBG mRNAs increased after glucose withdrawal and remained relatively high throughout the 2 hr experiment. In contrast, the mRNAs encoding RPGs showed very low P/T ratios between 10 and 30 min after glucose withdrawal, and these ratios increased between 30 and 120 min (Figure 2A). Low P/T ratios indicate that a population of nontranslating mRNA exists in the cell. To test this interpretation directly, we performed qRT-PCR on polysome gradient fractions after 10 min of glucose starvation. RPG mRNAs (low P/T genes) accumulated in ribosome-free fractions at the top of the gradient, whereas mRNAs from genes with high P/T ratios did not (Figure S4). In principle, the apparent increase in ribosome occupancy for RPGs after 60–120 min could result from improved translation (increased P) or from degradation of the nonpolysomal pool of mRNA. Given that P/P and T/T ratios for RPGs were divergent at early times minus glucose and converged after 60 min of starvation (Figure 3), the second interpretation is more plausible, suggesting that this subpopulation of mRNA is only transiently stable as a non-translating pool. Notably, the messages that were most dramatically downregulated upon glucose withdrawal and had low P/T values in acutely starved cells are the most abundant (Figure 4). Transient preservation of these mRNAs in a non-translating pool (Figure S4) may reflect a bet-hedging strategy (see the Discussion).

Only a Subset of mRNAs Can Return to Polysomes after Starvation and Refeeding

Previous reports showed that the global reduction in protein synthesis caused by 10 min of glucose withdrawal can be reversed within 5 min of glucose readdition (Ashe et al., 2000; Brengues et al., 2005). It was proposed that this rapid recovery of translation is due to mobilization of stored, translationally silenced mRNPs from P bodies (cytoplasmic RNP granules) to polysomes. Only two reporter genes were tested for the capacity to return to polysomes in the absence of new transcription (Brengues et al., 2005). Our data suggest that most mRNAs are depleted from the total mRNA pool coincident with their loss from polysomes (Figure 1C and Figure S4D), and that the capacity for translational resurrection of preexisting messages is thus narrowly restricted to a select subpopulation that
includes the RPG mRNAs. To test this hypothesis directly, we examined global as well as gene-specific recovery of translation after glucose readdition to starved cells. New transcription was inhibited with the drug thiolutin (Grigull et al., 2004), to assess the capacity of preexisting mRNAs to return to polysomes. Treatment with thiolutin slightly blunted translational recovery upon glucose readdition, resulting in fewer heavy polysomes, more disomes, and persistence of a larger 80S monosome peak (Figure 5A). These results indicate that initiation of translation on newly synthesized mRNAs accounts for some of the “recovery” of translation after glucose readdition, even after only 10 min of glucose starvation.

Nevertheless, substantial polysome recovery occurred even when new transcription was inhibited (Figure 5A). To determine which genes participate in this recovery, the polysomal and nonpolysomal mRNA pools were examined with microarrays after 10 min of glucose starvation and again after 5 min of glucose repletion, with thiolutin present throughout. A gene’s P/T ratio during glucose starvation predicted its ability to return to polysomes upon glucose readdition. Genes that showed high or neutral P/T ratios after glucose removal (Figure 2A) were depleted from the nonpolysomal as well as the polysomal mRNA pools after 10 min of glucose starvation and showed very little mobilization into polysomes 5 min after glucose readddition (Figures 5B and 5C). This argues that genes like the RBGs, which have high P/T ratios under conditions of decreasing P, do so because of rapid depletion of the nonpolysomal mRNA. Consistent with this interpretation, we found by qRT-PCR that 80%–95% of the total mRNA from these high P/T genes is gone from the total mRNA pool by 10 min after glucose starvation. In contrast, the genes that showed low P/T after glucose withdrawal were preferentially depleted from the polysomal compared to the nonpolysomal pool (Figure S4D).
This group of genes also showed significantly greater mobilization into polysomes upon glucose readdition by microarray (p < 0.0001), and by qRT-PCR analysis of select genes' mRNA abundance in polysome gradient fractions (Figure 6). Similar results were observed for more moderately downregulated genes, with the low P/T subpopulation showing significantly greater recovery than the high or neutral P/T genes, although the extent of recovery was somewhat less. The RPGs as a class showed ~6-fold more recovery than all other starvation-repressed genes (T/T clusters 5, 6, and 7 from Figure 2A) combined (p < 6.0 x 10^{-54}), whereas RBGs did not recover (Figure 5D).

We verified that P bodies formed under these conditions based on the localization of previously characterized protein and RNA reporters (Figure S5). In addition, we examined the localization of RPS26A-U1A, a low P/T gene capable of returning to polysomes upon refeeding, and of RRP4-U1A and LSG1-U1A, high P/T genes that are largely depleted within 10 min of glucose withdrawal and are not capable of returning to polysomes in the absence of new transcription. We detected P body localization for RPS26A-U1A and RRP4-U1A, whereas RBGs did not localize in the absence of new transcription (Figure 6B), or alternatively, if the P body localized portion of RPS26A mRNA is the decay intermediates of the fraction (data not shown). These reporter experiments don’t distinguish whether the P body-localized portion of RPS26A mRNA is the decay intermediates of the fraction (data not shown). These reporter experiments don’t distinguish whether the P body-localized portion of RPS26A mRNA is the decay intermediates of the fraction (data not shown). These reporter experiments don’t distinguish whether the P body-localized portion of RPS26A mRNA is the decay intermediates of the fraction (data not shown). These reporter experiments don’t distinguish whether the P body-localized portion of RPS26A mRNA is the fraction of intact mRNA that returns to polysomes upon refeeding. Note the relative heights of the disome (open arrowhead) and polysome (filled arrowhead) peaks and the widths of the monosome peaks ± thio. Analytical polysome assays (A, E, and F) were repeated two to four times. Representative traces are shown for each.

(B) Ratio values shown by color scale from microarray analysis of nonpolysomal (NP) and polysomal (P) mRNA from cells starved for 10 min (YP 10' +thio) or starved and refed for 5 min (+D 5' +thio). Polysomal mRNA from YPD cultures served as the reference sample for each array. Genes are organized according to T/T and P/T groups from Figure 2. Group 5 is shown at 50% vertical scale. Genes that showed low P/T ratios in starved cells were less depleted from the nonpolysomal fraction than genes with high or neutral P/T ratios and showed greater mobilization into polysomes upon refeeding. "Δpoly" values were derived from the ratio of ratios (YPD5_P versus YP10_P).

(C) Graphs show quantification of the Δpoly values from (B).

(D) RPGs as a class preferentially recovered in polysomes upon glucose readdition. All, all genes from T/T 5–7. Error bars in (C) and (D) indicate SEM. Asterisks (*) indicate Student’s t test p value < 0.0001.

(E) Prolonged glucose starvation in the absence of new transcription leads to reduced polysome recovery upon refeeding. Cells were starved for 45 (top row) or 60 (bottom row) minutes in the presence of thiolutin before refeeding for 5 min. Polysome profiles of recovery (+D 5') after only 10 min –glucose +thio are overlaid for comparison (red lines).

(F) New transcription contributes substantially to polysome recovery after prolonged starvation. Polysome profiles of recovery (+D 5') after 45' or 60' of starvation are shown in black. Polysome recovery after starvation for 45 or 60 min +thiolutin is shown (blue lines) for comparison. See also Figure S5.
Figure 6. Quantitative RT-PCR Validation of Select Genes’ mRNA Abundance in Polysome Fractions after Glucose Starvation and Refeeding

(A) Polysome gradient fractions from plus glucose (left), starved (10 min minus glucose, with thiolutin, center), and re-fed (10 min minus glucose and 5 min plus glucose, with thiolutin, right).

(B and C) mRNA abundance per fraction, relative to one-twelfth input and normalized to Fluc dope-in control RNA (left). Adjusted mRNA abundance per fraction was determined by qRT-PCR comparison with plus glucose fractions, shown on the same scale as plus glucose samples (center, right). Error bars indicate standard deviation of the mean of three replicates.
Thus, P bodies likely contain endogenous mRNAs undergoing decay. Whether or not they also contain stored translationally repressed RPG mRNAs remains an open question. Wherever translationally repressed mRNAs reside in the cell, only select mRNAs are able to persist in a nontranslating pool and resume active translation after a short period of glucose starvation.

The glucose starvation microarray time course data suggest that the nontranslating subpopulation of mRNAs that accumulates at early times is depleted after more prolonged glucose starvation (Figures 2A and 3). This interpretation of the data predicts a turning point after which any recovery of polysomes upon glucose readdition would require new transcription. To test this prediction, we subjected cells to varying periods of glucose starvation, with and without inhibition of new transcription by the drug thiolutin, and examined global translation activity after five minutes of glucose readdition. After 45 or 60 min of starvation in the absence of new transcription, polysome recovery was greatly reduced compared to cells starved for only 10 min (Figure 5E). If new transcription was allowed to occur during extended starvation, translational recovery upon glucose readdition increased (Figure 5F). These observations, together with the indication that specific mRNAs (RPGs) are lost between the 10 and 60 min time points (Figure 3), suggests that translational activation of these silenced mRNAs contributes substantially to polysome recovery upon refeeding of briefly starved cells. These data argue that cells’ inability to rapidly restore translation without new transcription after longer periods of starvation is due to the disappearance of a population of nontranslating mRNAs. Between 30 and 60 min appears to be a switch point for P/T ratios in our data, regardless of whether the ratios are increasing or decreasing. This suggests that the timing of the “turning point” for cells to recover translation of stored RPGs may relate to widespread changes in the activity of factors that influence P/T through effects on mRNA stability.

Molecular Insights into Selective Preservation of Nontranslating RPG Transcripts

How are certain mRNAs able to persist in the cell, even transiently, following glucose starvation, when the majority of mRNAs are depleted from the total RNA pool coincident, within the time resolution of our experiments, with their loss from polysomes? Comparison with genome-wide mRNA half-life measurements indicates that stability in glucose-replete conditions is probably not the determining factor for an mRNA’s capacity to persist in a stable nontranslating pool after glucose withdrawal (Figure S6). In particular, the RPGs as a class have
short half-lives in glucose-replete conditions compared to most genes (Grigull et al., 2004; Wang et al., 2002). Alternatively, the “posttranscriptional operon” hypothesis posits a role for RNA-binding proteins (RBPs) in coordinating the posttranscriptional fate of functionally related genes (Keene, 2007; Keene and Tenenbaum, 2002). To investigate the possibility that specific RBPs affect the posttranscriptional behavior of mRNAs after glucose withdrawal, we examined data from a recent genome-wide association study that identified mRNA targets of 40 yeast RBPs (Hogan et al., 2008). Comparing the “transcriptional” (changes in total mRNA abundance) and posttranscriptional (P/T) regulatory groups identified in our study with RBP-mRNA target groups revealed concordance between RBP association patterns and P/T but not T/T (Figure 7A). For simplicity, only the seven RBPs that showed significant enrichment or de-enrichment (p < 0.01, Bonferroni corrected) of target mRNAs within at least one group are displayed. Similar to GO terms, which rarely spanned opposing P/T groups, RBP-association profiles were similar for “high” and “neutral” P/T groups for many RBPs, and for “low” and “neutral” for PuF3, whereas the “high” and “low” groups had almost no enrichments in common and frequently appeared to be mirror images of each other. In contrast, organizing RBP-association patterns by T/T group resulted in a “checkerboard” appearance, despite the fact that T/T groups contain functionally and cyttopically related genes. This suggests that the coherence of the association between P/T behavior and particular RBPs might be the result of direct effects of these RBPs on mRNA translatability and/or mRNA stability outside of polysomes.

Pab1-associated mRNAs were notably enriched with groups having low P/T ratios, and correspondingly depleted in groups with high and neutral P/T ratios (Figure 7A). Comparison of P/T groups with genome-wide measurements of poly(A) tail lengths (Beilharz and Preiss, 2007) revealed a lack of short-tailed and an enrichment of long-tailed genes among mRNAs with low P/T values. Conversely, genes with long poly(A) tails were significantly underrepresented among groups with high P/T ratios (Figure 7B). The association of Pab1 and long poly(A) tails with groups of mRNAs with low P/T ratios was counterintuitive given Pab1’s role in enhancing translation initiation and the positive association with polysomes in glucose-starved cells. Many of these genes are known to be transcriptional targets of well-characterized glucose-regulated transcription factors and encode proteins required for survival and growth in the absence of glucose. Furthermore, the kinetics of accumulation of these mRNAs in polysomes was nearly indistinguishable from their appearance/accumulation in the total mRNA pool. The simplest interpretation of these data is that cells respond to glucose starvation by upregulating expression of genes whose products promote adaptation to low glucose environments. This view agrees with extensive genetic and genome-wide gene expression studies (reviewed in Zaman et al., 2008) and is consistent with an earlier study, which reported (as data not shown) that protein synthesis, measured by [35S]methionine incorporation rates, decreased to 20% after 5 min without glucose, and returned to 40% by 15 min without glucose (Kuhn et al., 2001). We conclude that there is widespread translation at a reduced level in glucose-starved cells.

**DISCUSSION**

**Translation Is Reduced, Not “Inhibited,” after Glucose Withdrawal**

Since the initial discovery of a rapid and dramatic signal-mediated reduction in polysomes after glucose withdrawal (Ashe et al., 2000), subsequent studies have investigated the mechanisms responsible for “global inhibition” of translation in glucose-starved cells (for example, Brengues and Parker, 2007; Brengues et al., 2005; Hilgers et al., 2006; Holmes et al., 2004; Hoyle et al., 2007). Thus, we were initially surprised to observe widespread agreement between the relative abundance of mRNAs in the polysomal and total mRNA pools following glucose withdrawal. The data reported here show that mRNAs from more than 1,000 genes rapidly increased their relative association with polysomes in glucose-starved cells. Many of these genes are known to be transcriptional targets of well-characterized glucose-regulated transcription factors and encode proteins required for survival and growth in the absence of glucose. Furthermore, the kinetics of accumulation of these mRNAs in polysomes was nearly indistinguishable from their appearance/accumulation in the total mRNA pool. The simplest interpretation of these data is that cells respond to glucose starvation by upregulating expression of genes whose products promote adaptation to low glucose environments. This view agrees with extensive genetic and genome-wide gene expression studies (reviewed in Zaman et al., 2008) and is consistent with an earlier study, which reported (as data not shown) that protein synthesis, measured by [35S]methionine incorporation rates, decreased to 20% after 5 min without glucose, and returned to 40% by 15 min without glucose (Kuhn et al., 2001). We conclude that there is widespread translation at a reduced level in glucose-starved cells.

**Most mRNAs Are Depleted Coincident with Translational Repression**

Previous studies suggested that widespread degradation of mRNAs is not part of the response to glucose withdrawal based on northern blots of specific genes (Ashe et al., 2000). Examining the behavior of those genes in our genome-wide experiments reveals that their behavior is not typical. Three of the genes examined, PAB1, ACT1, and PGK1, were barely reduced in polysomes after 10 min without glucose (12%, 13%, and 5% reductions, respectively), and the fourth gene, RPL28/CYH2, is unusual in its capacity to survive as a nontranslating mRNA. Among more than 2,000 genes showing consistently reduced mRNA levels in polysomes after glucose withdrawal, only a small minority (<20%), dominated by the RPGs, persisted in a stable nontranslating pool that could return to polysomes if glucose was restored. Most mRNAs were depleted from the total mRNA population as quickly as they were depleted from...
polysomes. This observation is consistent with reports showing increased decapping and decay of specific mRNAs in response to decreased translation initiation (LaGrandeur and Parker, 1999; Schwartz and Parker, 1999).

**Mechanisms of Posttranscriptional Regulation in Glucose-Starved Cells**

As our data are consistent with widespread decay of translationally repressed mRNAs, we reconsidered the evidence for and against a model of widespread mRNA decapping as the cause of translational repression after glucose starvation. Deletion of either component of the decapping enzyme complex (dcp1Δ or dcp2Δ) prevents polysome reduction after glucose withdrawal (Holmes et al., 2004). Moreover, the ability to inhibit protein synthesis after glucose removal correlated with the extent of residual decapping activity retained by various dcp1 mutants. Our genome-wide results are equally consistent with the model that activation of decapping causes widespread translational repression and mRNA decay or, alternatively, that translational repression occurs first and mRNA decay follows soon after. We favor the first model because inhibition of decapping prevents translational repression, whereas no known translational repressive mechanism is required for polysome collapse, despite extensive efforts to identify such a mechanism (Holmes et al., 2004). Nevertheless, decapping and 5′ to 3′ mRNA decay can occur on polysome-associated mRNAs (Hu et al., 2009).

Thus, for the majority of genes whose mRNAs are depleted from the total pool coincident, within the time resolution of our experiments, with their disappearance from polysomes, we cannot conclude whether they are first evicted from polysomes and then degraded, or whether the initiation of decay prevents subsequent loading of ribosomes (and permits runoff of previously loaded ribosomes).

If widespread decapping is in fact responsible for most of the translational repression in glucose-starved cells, three major questions remain. First, what is the trigger that activates decapping within 1 min after glucose withdrawal? The rapidity of the response suggests that it involves posttranslational modification of some factor. Mutations that disrupt either the glucose derepression signaling pathway or the Ras/PKA pathway interfere with translational repression (Ashe et al., 2000), but no direct molecular connections to regulation of the decapping machinery have yet been characterized. Second, how do some messages escape decapping? There are two types of escapes: the “privileged” mRNAs, including the RPG messages, that are present at the moment of glucose withdrawal and shift out of polysomes, and all the mRNAs that are produced afterwards. Third, what mechanism accounts for the translational repression of the RPGs?

Our results implicate the unusually long poly(A) tails of some messages and their association with the poly(A) binding protein as likely determinants for the preservation of nontranslating mRNA. This is an attractive model given the evidence that deadenylation is the rate-limiting step in most mRNA decay (Parker and Song, 2004). For the second class of escapes—mRNAs whose abundance increases in glucose-starved cells—it is conceivable that they are endowed with longer poly(A) tails in the nucleus and/or that they are assembled into RNP with features that render them inherently resistant to decapping.

We do not favor such a model because these mRNAs are depleted from polysomes within 5 min after glucose readdition (see complete array data available in Table S2). An alternative explanation is that the “special” feature of newly transcribed mRNAs is simply that they enter the cytoplasm after the decapping machinery is sequestered in P bodies.

How are the RPG mRNAs translationally repressed? The observation that this group of mRNAs readily returns to active translation upon glucose readdition strongly suggests that they are not decapped. Furthermore, our model that their long poly(A) tails and stable association with Pab1 are important for escaping decapping argues that they retain all of the features thought to be required for ribosome recruitment under normal conditions. The mTOR pathway is implicated in nutrient regulation of RPG translation in mammalian cells (Hamilton et al., 2006), but no such mechanism was thought to exist in yeast (Warner, 1999). Although the molecular details of the yeast mechanism almost certainly differ, our data suggest that a translational control mechanism targeting RPGs does exist. Genome-wide investigations of yeast translational responses to other stresses including high salinity (Melamed et al., 2008), transfer to a nonfermentable carbon source (Kuhn et al., 2001), and rapamycin treatment (Preiss et al., 2003) have also reported translational repression of RPGs, suggesting that this mechanism acts downstream of a common stress response signaling pathway.

**Implications for the Interpretation of P Bodies**

Glucose starvation in yeast has been extensively studied as a model stress to investigate the relationship between translational repression and mRNA localization to cytoplasmic granules or P bodies. A central tenet of the model is that P bodies are not just sites of mRNA decay, but can also be sites for storage of translationally silent intact mRNAs capable of return to the actively translating pool in response to altered cellular conditions (Brengues et al., 2005; Teixeira et al., 2005). Because core P body components and their roles in mRNA metabolism are well conserved, this model has far-reaching implications for the regulation of gene expression in eukaryotic cells (Anderson and Kedersha, 2006; Parker and Sheth, 2007).

Our findings refute the notion that reciprocal movement between a nontranslating mRNA pool and polysomes is a general property of eukaryotic mRNAs (Brengues et al., 2005). This model was based on previous observations of two reporter mRNAs after glucose starvation and repletion. We found that less than 20% of genes have the capacity to persist in a translationally silent state in glucose-starved cells. This raises the possibility that the majority of the proteins accumulated in P bodies under these conditions are engaged with messages undergoing irreversible decay, although we cannot rule out the possibility that a subset of P body associated endogenous mRNAs can escape and resume translation. If so, the RPG messages are good candidates.

**Bet Hedging**

What advantage might transient preservation of RPG messages confer, and why should their behavior differ from the RBGs? Although the ribosome biogenesis factors and ribosomal proteins (RPs) function together in the assembly of new
ribosomes, the factors act catalytically whereas the RPs are required stoichiometrically. Accordingly, the mRNAs encoding the RPs are among the most abundant messages in the cell, comprising up to 30% of cellular mRNA during rapid growth in rich media and requiring a significant investment of energy for their synthesis (Holstege et al., 1998; Nagalakshmi et al., 2008; Warner, 1999). If the cell were to degrade these messages immediately after release from polysomes, reversing this decision would be energetically costly. By retaining these mRNAs in a non-translating state for 10–30 min after the perception of glucose withdrawal, the cell might preserve the capacity for rapid resumption of growth should glucose return or the starvation signal turn out to have been a false alarm. Given that the production of new ribosomes largely determines the growth rate and generation time of yeast in nutrient replete conditions (Warner, 1999; Zaman et al., 2008), it is tempting to speculate that the selection of RPG mRNAs for transient preservation is a bet-hedging strategy by which the cell retains the capacity for a return to rapid growth and division while simultaneously initiating preparations for prolonged starvation. Previous investigations of global relationships between transcriptional and translational changes after other environmental stresses (high-salt shock, transfer to a nongmentable carbon source, or treatment with rapamycin to mimic nutrient deprivation) also described translational repression of RPGs through measurements that required the existence of a stable nontranslating population of mRNA (Kuhn et al., 2001; Melamed et al., 2008; Preiss et al., 2003). The transient preservation of translationally repressed mRNAs may be a general feature of yeast cells’ responses to growth-inhibiting stresses. Although translational activity was not determined, exploration of the common response to diverse environmental stresses revealed reduction in both RBG and RPG mRNAs, with the RPGs showing a delay (Gasch et al., 2000).

An interesting and unexplained phenomenon revealed by our data is the existence of a turning point, beyond which the cell appears to give up on hedging its bets and commits fully to the starvation gene expression program. How is the timing of this turning point determined molecularly? Does the timer run at a constant speed, or can it be adjusted according to features of the environment that might predict how likely it is that the cell will experience a reversal of fortunes in the near future? Do all cells hedge a little, transiently storing 30%–50% of their RPG mRNAs for transient preservation is a bet-hedging strategy by which the cell retains the capacity for a return to rapid growth and division while simultaneously initiating preparations for prolonged starvation. Previous investigations of global relationships between transcriptional and translational changes after other environmental stresses (high-salt shock, transfer to a nongmentable carbon source, or treatment with rapamycin to mimic nutrient deprivation) also described translational repression of RPGs through measurements that required the existence of a stable nontranslating population of mRNA (Kuhn et al., 2001; Melamed et al., 2008; Preiss et al., 2003). The transient preservation of translationally repressed mRNAs may be a general feature of yeast cells’ responses to growth-inhibiting stresses. Although translational activity was not determined, exploration of the common response to diverse environmental stresses revealed reduction in both RBG and RPG mRNAs, with the RPGs showing a delay (Gasch et al., 2000).

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Translational Regulatory Logic in Starved Cells

ACCESSION NUMBERS

The microarray data have been deposited at the Gene Expression Omnibus (GEO) repository under the accession number GSE31393.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, seven figures, and two tables and can be found with this article online at doi:10.1016/j.molcel.2011.09.019.

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