Identifying eIF4E-binding protein translationally-controlled transcripts reveals links to mRNAs bound by specific PUF proteins

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ABSTRACT
eIF4E-binding proteins (4E-BPs) regulate translation of mRNAs in eukaryotes. However the extent to which specific mRNA targets are regulated by 4E-BPs remains unknown. We performed translational profiling by microarray analysis of polysome and monosome associated mRNAs in wild-type and mutant cells to identify mRNAs in yeast regulated by the 4E-BPs Caf20p and Eap1p; the first-global comparison of 4E-BP target mRNAs. We find that yeast 4E-BPs modulate the translation of >1000 genes. Most target mRNAs differ between the 4E-BPs revealing mRNA specificity for translational control by each 4E-BP. This is supported by observations that eap1Δ and caf20Δ cells have different nitrogen source utilization defects, implying different mRNA targets. To account for the mRNA specificity shown by each 4E-BP, we found correlations between our data sets and previously determined targets of yeast mRNA-binding proteins. We used affinity chromatography experiments to uncover specific RNA-stabilized complexes formed between Caf20p and Puf4p/Puf5p and between Eap1p and Puf1p/Puf2p. Thus the combined action of each 4E-BP with specific 3′-UTR-binding proteins mediates mRNA-specific translational control in yeast, showing that this form of translational control is more widely employed than previously thought.

INTRODUCTION
It has become increasingly evident that the translation process for a large number of mRNAs is controlled at the level of initiation in a wide variety of cells. The selection of mRNAs for translation is mediated by the binding of eukaryotic initiation factor (eIF) 4F to the 5′-end of each mRNA. The eIF4F complex is composed of eIF4E that binds to the 5′ 7-methyl guanosine mRNA cap, eIF4A an RNA helicase, and eIF4G a scaffolding protein capable of bridging contacts between eIF4E and 40S ribosomal subunits. eIF4G and the eIF4E-binding proteins (4E-BPs) share a motif (YXXXXLφ, where X is any residue and φ is hydrophobic) that is critical for eIF4E interaction. Hence 4E-BP bound mRNAs exclude eIF4G and are translationally silenced (1,2). Mammalian 4E-BP1 phosphorylation status modulates its affinity for eIF4E. mTOR dependent hyperphosphorylation of 4E-BP1 causes its dissociation from eIF4E relieving repression and promoting eIF4F assembly, translation and cell growth (3,4). A second class of 4E-BP has been identified from studies of early development, e.g. in Xenopus and Drosophila where asymmetric expression of proteins is important for determining the body plan of the embryo (2,5). For these 4E-BPs additional protein–protein and protein–mRNA contacts tether the 4E-BP to specific mRNA targets. Thus Xenopus Maskin is a 4E-BP that also binds to the cytoplasmic polyadenylation element-binding (CPEB) protein which itself interacts with sequences within mRNA 3′-UTRs. Together and in combination with other proteins, these factors mediate circularization and repression of transcripts until developmentally
regulated signalling events activate both cytoplasmic polyadenylation, disrupt the Maskin-eIF4E interaction and promote translation (2,5).

*Saccharomyces cerevisiae* possesses two 4E-BPs, Caf20p and Eap1p of 18 and 70 KDa, respectively. Neither protein shares obvious sequence homology with each other or 4E-BPs from non-yeast species, except for the eIF4E interaction consensus motif, however both yeast 4E-BPs interact with eIF4E to inhibit translation of capped reporter mRNAs (6,7). Although deletion of either 4E-BP gene confers no growth defect under standard laboratory conditions, *cafl20A* alleviates growth defects of some translation factor mutations and conversely its overexpression exacerbates these phenotypes (8). *eap1A* causes rapamycin resistance (7) and partial resistance to translation inhibition caused by diame (a thiol oxidant), and cadmium (a heavy metal) (9). In the yeast *Σ1278b* background deletion of either 4E-BP prevents pseudohyphal growth following nitrogen limitation (10). Similarly, *eap1A* and *cafl20A* cells respond differently to membrane stress (11,12). Taken together the data are consistent with the idea that each 4E-BP interacts with eIF4E and competes with eIF4G to modulate the translation of a different subset of specific yeast genes.

Little progress has been made in identifying specific mRNA targets regulated by the yeast 4E-BPs. One possible candidate is the G1 cyclin *CLN3* as it can be controlled by altering eIF4E activity. *CLN3* mRNA contains a short up-stream ORF that plays a role repressing Cln3p expression (13). Temperature sensitive eIF4E mutations (e.g. *cdc33-1*) confer G1 arrest and reduce expression of *CLN3* and enhanced *CLN3* expression is sufficient to restore G1-S phase progression (14). Recently, a second mRNA target, *POM34* mRNA was described, which encodes a membrane protein component of the nuclear pore complex. In cells bearing spindle-pole body duplication defects Eap1p was required for *POM34* translation (15).

Given the paucity of information concerning specific 4E-BP targets, we used a translational profiling approach to identify mRNAs whose translation is altered following loss of each yeast 4E-BP and found that both altered translation of a large fraction of yeast genes. Quantitative real-time reverse transcription-polymerase chain reaction (RT–PCR) and immunoblot analyses have validated the micro-array approach for selected targets and a phenotypic analysis indicates that the observed altered translation of several nitrogen metabolism genes may contribute to the sensitivity of the *eap1A* strain to growth on some nitrogen sources. Despite sharing a common interaction site on eIF4E, we find that most target mRNAs differ between the 4E-BPs suggesting that additional factors are necessary to imparts 4E-BP mRNA specificity. To begin to address this mechanistic issue we compared the identities of mRNAs found by others to interact with specific RNA-interacting proteins, with our translational profiling data. Our analysis suggested that certain yeast PUF proteins may be among those important for mRNA specificity. As PUF proteins have been implicated in translational control in higher organisms, we tested this prediction directly and report that Puf4/5p-bound mRNA complexes associate with a fraction of Caf20p, while Puf1/2p associate with Eap1p. Taken together these studies reveal that ~1000 genes are subject to translational control via the 4E-BPs and provides evidence that 4E-BP association with specific 3'-UTR-binding proteins imparts mRNA target specificity in yeast.

**MATERIALS AND METHODS**

**Yeast strains and growth conditions**

Isogenic yeast strains BY4742 (*MATα his3Δ1 leu2Δ0 lys2Δ0 ura3Δ0*), Y17334 (*MATα his3Δ1 leu2Δ0 lys2Δ0 ura3Δ0 caf20A::kanMX4*) and Y17036 (*MATα his3Δ1 leu2Δ0 lys2Δ0 ura3Δ0 eap1A::kanMX4*) were obtained from Euroscarf (Frankfurt, Germany) and grown in synthetic dextrose (SD) minimal medium containing auxotrophic supplements and grown to *A_{600} = 0.7* for all array experiments.

For nitrogen source growth experiments auxotrophic markers were complemented with plasmid encoded gene copies [pRS313-HIS3; pRS314-TRP1; pRS315-LEU2 (16), and pRSK-URA3 LYS2 (a gift from M. Stark, University of Dundee)]. Wild-type or mutant *EAPI* plasmids: pJF3896 (HA-*EAPI LEU2 CEN*) and pBMK492 (HA-*eap1-Y109A,L114A LEU2 CEN*) replaced pRS315 where indicated (10). Cells were grown on SD medium containing a single indicated nitrogen source at the following concentrations: 0.1% w/v ammonium sulphate, allantoin, 0.02% w/v l-isoleucine, l-leucine, l-tryptophan, l-valine, l-asparagine, l-aspartic acid, l-glutamine, l-glutamic acid, l-serine, 0.01% w/v ammonium chloride, 0.01M urea. All media were adjusted to pH 5.3. Synthetic complete (SC) contained 0.5% w/v ammonium sulphate and 0.2% w/v of a mix of all 20 amino acids, inositol, para-aminobenzoic acid, uracil, adenine as described earlier (17). Agar plates (2% w/v bacto agar) were incubated at 30°C for 2–5 days. Doubling times (*T_d*) in liquid media were determined using 96-well microplate cultures grown in a temperature-controlled absorbance reader with shaking (BMG Labtech). *A_{600}* measurements were recorded every 5 min for 20 h. A previously described MS Excel macro was adapted and used to automate *T_d* determinations (18).

PUF-TAP tagged strains were a kind gift from Gerber and Herschlag (19). *CAF20-Myc* and *EAPI-Myc* tagged versions of each PUF-TAP strain were made using pYM4 and PCR to integrate C-terminal 3Myc-KanMX6 marker as described earlier (20). Similarly, pYM19 (9Myc- *HIS3MX6*) and pYM13 (TAP-KanMX6) was used to create Myc-tagged and TAP-tagged versions, respectively, of *CAF20, EAPI, PUF2, PUF3* and *PUF5* in BY4742, Y17334 and Y17036 (20). PCR was used to verify genomic integrations.

**Polyosome fractionation, RNA preparation and microarray analysis**

Cell extracts, polynosome fractionation, RNA preparation and array bioinformatics analysis were done exactly as described earlier (21). Complete data sets are
available at ArrayExpress (www.ebi.ac.uk/microarray/using accession numbers caf20Δ: E-MEXP-1308; eap1Δ: E-MEXP-1309).

Quantitative RT–PCR analysis

RNA analysis by RT–PCR was carried out using the MyIq single-colour real-time PCR detection system and IQ SYBR Green Supermix (BioRad Laboratories) as described earlier (21,22). Signals were quantified relative to actin mRNA control.

TAP affinity chromatography

PUF-TAP or 4E-BP-TAP and control strains were grown in YPD medium to a concentration of 6 × 10⁶ cells per ml. Cultures were harvested by centrifugation (5500g, 10 min, 4°C), washed in ice cold 1 mM phenylmethylsulphonyl fluoride (PMSF) and resuspended in two cell volumes of Buffer A [20 mM Tris–HCl (pH 8.0), 140 mM KCl, 1.8 mM MgCl₂, 0.1% Nonidet P-40 (NP-40), 0.5 mM dithiothreitol (DTT), 1 mM PMSF, 1× complete EDTA-free protease inhibitor cocktail tablet (Roche), 100 U/ml RNAsin (Promega)]. Cell suspension was frozen and ground under liquid nitrogen and stored at −80°C. Crude cell lysates were cleared by centrifugation (5500g for 20 min, 4°C). Extracts (2 mg) were incubated with 400 µl (50% [v/v]) IgG Sepharose beads (GE Healthcare) for 2 h at 4°C and affinity purified as described earlier (19). PUF proteins were released from the IgG beads by heating at 65°C for 3 min in 2× non-reducing Protein Loading Buffer [6.25 mM Tris–HCl (pH 6.8), 2% SDS, 10% (v/v) glycerol, 0.002% (w/v) bromphenol blue].

Immunoblotting

Whole cell extracts; Cells were harvested and resuspended in three times wet pellet volume of Buffer B [50 mM Tris–HCl pH 7.5, 100 mM NaCl 1 mM EDTA 1 mM PMSF and 1× Protease Inhibitors (Roche)] and ground in liquid Nitrogen. Samples were assayed for protein concentration (Bradford) and diluted to 1 µg protein per ml. TAP proteins were resolved by SDS–PAGE electrophoresis followed by transfer to a nitrocellulose membrane. Immunoblotting detection used horseradish peroxidase (HRP) conjugated primary antibodies to Protein A (Abcam) and c-Myc (9E10, Santa Cruz Biotechnology and 4A6, Millipore), and chemiluminescent detection (Pierce Biotechnology). Other primary antibodies used were caf20p (10), Gcd11p (23), Cic1p (24), Lsm8p (25), Taf3p, Taf7p (26), Sec9p (27), Pub1p (28), Ade2p and Arp2p (Santa Cruz Biotechnology) and were detected using appropriate HRP-conjugated secondary antibodies as indicated earlier.

RESULTS AND DISCUSSION

Microarray approach to determine roles for 4E-BPs

To determine whether translation of specific mRNAs was altered by elimination of Caf20p or Eap1p under steady state growth conditions, we used a microarray-based approach so that the translation state of all mRNAs could be surveyed simultaneously. We used isogenic strains produced by the systematic yeast gene deletion project for this study. Our initial growth characterization revealed that the caf20Δ and wild-type strains behaved identically in both rich and minimal medium, while the eap1Δ strain typically exhibited an initial growth lag when recovering from stationary phase, but then proliferated at a rate indistinguishable from the wild-type strain (data not shown). Polyribosome profile analysis revealed a minor halfmer phenotype for the eap1Δ strain, but no obvious global defect in the caf20Δ strain (Figure 1A). For array analysis the following samples were collected: (i) total RNA samples to control for changes in transcript abundance and (ii) polysome gradient fractionated samples. Fractionated samples were split into monosomal and polysomal fractions as indicated in Figure 1A. Each RNA sample was processed into complementary RNA and hybridized to Affymetrix yeast S98 gene chips arrays using standard protocols (21,22). All analyses were performed in duplicate and the resulting data were analysed and compared using standard bioinformatics procedures (as outlined in the Supplementary Data). MA plots and statistical analysis of variance tests (Supplementary Figure S1A, B, D, E, G and H and data not shown) indicated that there was only modest dispersal between biological replicate samples, confirming that the arrays are reliable reporters of relative transcript abundance in our samples. As anticipated, a greater dispersal of data points was seen between wild-type and mutant samples (Supplementary Figure S1C, F and I).

4E-BPs have minimal impact on mRNA abundance

Under normal cellular conditions, changes in mRNA transcript abundance reflect changes in both mRNA synthesis and decay. In this study, overall transcript abundance changes were relatively minor. caf20Δ alters transcript abundance >2-fold for 100 genes (29 up and 71 down) and eap1Δ 99 genes (56 up and 43 down). A complete listing of significantly altered genes is shown in Supplementary Tables S1–S4. As expected the transcript signals for the deleted gene were down-regulated most in each data set. The small number of genes affected is consistent with 4E-BPs having no widespread direct transcriptional role, instead the changes observed could be an indirect consequence of altered translation of transcription factors (see below). There was also no evidence of ‘potentiation’. This is a term used to describe co-ordinated transcription and translation responses observed during the reprogramming of gene expression to adapt to some cellular stresses (21,29). Fewer than 10 genes were both transcriptionally and translationally up-/down-regulated across both experiments (Figure 1F; numbers in black filled circles).

Overview of translational controls

We measured polysome:monosome [P/M] ratios for each transcript (probe set), termed a ‘translational state’, calculated as log₂[P/M] ratio for each mutant and transcriptionally and translationally up-/down-regulated cellular stresses (21,29). Fewer than 10 genes were both transcriptionally and translationally up-/down-regulated across both experiments (Figure 1F; numbers in black filled circles).
intensities. A ‘change in translational state’ was determined by comparing the ratios from mutant cells (m) to that from wild-type cells (w). These analyses are represented as scatter plots in Figure 1B and C and numerically (Supplementary Tables S5–S8). The plots reveal that the majority of mRNA probe sets fall on or around the diagonal, indicating that the translation of most genes is not affected by deletion of either gene. In line with previous studies using similar techniques, we applied a significance cut off of log2 = ±0.9 (i.e. >1.86-fold) change (21,29). This revealed that caf20Δ altered the translation of 784 mRNAs (471 up- and 313 down-regulated), while eap1Δ affected 329 mRNAs (176 up and 153 down) (Figure 1F). The fact that more mRNAs are Caf20p targets (784/1028 = 76%) than Eap1p agrees with our observations that the relative abundance of Myc tagged Caf20p is 2.8 ± 0.63-fold greater than Eap1 (data not shown). These data are consistent with Caf20p regulating translation of a greater number of mRNAs than Eap1p.

Eap1p acts as a translational repressor for mRNAs that are poorly associated with polysomes under normal growth conditions. This observation is consistent with the idea that Eap1p plays a role repressing translation of a subset of genes during normal growth. This is illustrated graphically in Figure 1D, where the spot location is derived from the caf20Δ experiment, but spots are coloured according to the eap1Δ data. It is evident that most eap1Δ up-regulated genes (red spots) are to the left half of the plot, while the down-regulated ones (blue) are on the right. Therefore the majority of genes which are up-regulated in eap1Δ cells (red spots in Figure 1C and D) are among those transcripts which are poorly associated with polysomes in wild-type cells (81% or 142/176 log2[Pw/Mw]<0). A reciprocal arrangement holds for down-regulated genes (14% or 21/153 log2[Pw/Mw]<0). This is much less evident for Caf20p regulated transcripts. Figure 1E illustrates where the caf20Δ regulated genes are highlighted on the eap1Δ plot. Although >50% of the genes up-regulated by caf20Δ are
on the left in Figure 1E, the bias is less pronounced, suggesting distinct roles for Caf20p and Eap1p in repressing translation of genes under normal growth conditions.

As an independent confirmation that the arrays accurately reported translation state changes, quantitative RT–PCR analysis was performed on polysome fractionated RNA samples with oligonucleotide primers specific for a subset of genes (Figure 2A). Genes were selected to cover a range of up- and down-regulated mRNAs and also some unregulated mRNAs. Despite differences between these techniques, a good correlation was observed between the array and RT–PCR approaches.

The 4E-BPs interact with eIF4E via a common motif, and eIF4E itself binds the cap structure common to all mRNAs. Therefore it seems unlikely that these features alone impart the observed mRNA specific translational controls. Therefore our data suggest that there must be other factors, which work in concert with the yeast 4E-BPs to mediate control of translation (see below). It seemed likely to us that these other factors may bind to specific sequences within each mRNA and contribute to the observed control. As such, it was important to verify whether protein levels vary as anticipated by the translational profiling data. Almost all efficient strategies for epitope-tagging yeast genes involve disrupting either the 3′-UTR or replacing elements 5′ of the ORF. The 5′- and 3′-UTRs are common sites used by RNA-binding factors to mediate mRNA specific controls in many eukaryotes, so we reasoned that using a tagging approach was not appropriate here. Instead we searched the literature to identify available protein specific antibodies. We screened the antibodies available and identified several that functioned with our whole-cell extracts. The antibodies were used and compared with an actin loading control that did not vary between strains

A significant overlap between caf20 and eap1 data sets

Although the majority of mRNA targets differ between the array experiments, we observed a statistically significant overlap between the mRNAs similarly up-regulated by both eap1Δ and caf20Δ (52 genes $P = 9.8 \times 10^{-23}$; Figure 1D–F, Supplementary Table S9). In contrast, there are very few reciprocally regulated genes (e.g. only four genes up-regulated by caf20Δ are down-regulated by eap1Δ; Figure 1F).

During the course of our analyses, immunoblotting with anti-Caf20p anti-serum revealed that the steady state expression levels of Caf20p are reduced in eap1Δ cells (data not shown). As this observation may explain the observed overlap in translational control between these factors, we examined carefully the expression levels of both 4E-BPs. Myc-epitope tags were added by homologous recombination to the C-termini of CAF20 and EAP1 in separate strains. Immunoblotting confirmed that Caf20p expression was reduced ~50% in eap1Δ cells, while Eap1p levels were maintained in caf20Δ cells (Figure 1G). Therefore the observed reduced Caf20p levels may cause lower affinity Caf20p target mRNAs to lose normal translational control in eap1Δ cells and contribute to the overlap in array data sets.

4E-BPs repress genes controlling gene expression

To gain insight into the functions of the translationally regulated transcripts, each regulated gene was placed into a functional class based upon classifications included within the GOslim mapper (Saccharomyces Genome Database) and annotations at the Yeast Proteome Database (Figure 3 and Supplementary Tables S1–S8 and S10). This analysis revealed that many transcripts with enhanced [P/M] ratios are themselves involved in gene expression. Deletion of either 4E-BP enhances translation of many transcription factors and chromatin remodelling enzymes. These changes did not cause large changes in transcript abundance as arrays of total mRNA showed that few genes are altered by >2-fold (Supplementary Tables S1–S4). We assume that these cell have adapted to the loss of each factor and that the changes to the translation profile observed have normalized overall transcription as far as possible under the standard growth conditions examined. RNA metabolism is another abundant class of up-regulated genes for both 4E-BPs. In addition, Eap1p represses translation of many ribosome biogenesis factors involved in synthesis/assembly of both ribosomal subunits. This may account for the apparent modest halfmer phenotype observed in eap1Δ cells (arrowed in Figure 1A). Overall these data suggest that regulation of genes involved in RNA metabolism is an important function of these 4E-BPs and that each protein regulates specific targets involved in the same broad pathways/processes.

Caf20p represses translation of CLN genes and those important for polarised growth

The cell cycle, budding and site of polarised growth GO Slim categories are particularly over-represented within the translationally up-regulated caf20Δ mRNAs. Cyclins CLN1, CLN2, CLN3, CLB1, and CLB4 were among many cell cycle associated mRNAs. As indicated in the introduction, one of the best-characterized examples of a gene regulated by elf4E availability in yeast is the G1 cyclin CLN3. A temperature sensitive mutation in yeast elf4E (cde33-1) has a G1-S phase cell cycle defect that is overcome by elevated expression of CLN3 (14). We interpret our array data as confirming the importance of
translational controls for regulating cyclin expression and further suggest that an eIF4E-Caf20p interaction is important for translational repression of CLN3.

In metazoa correct intracellular localization of specific transcripts is important for both early developmental decisions and memory functions in neurons. 4E-BPs provide a mechanism to repress translation of delocalized mRNAs e.g. Drosophila 4E-BP Cup acts with Bruno to repress translation of oskar mRNA (30). Polarised growth in budding yeast requires asymmetric localization of components to the growing bud. Several mRNAs have also been found to be bud-localized. A survey of yeast genes uncovered 22-bud localized transcripts, of which 10 were asymmetrically localized to the bud (31). Intriguingly, six of these 10 genes (EGT2, IST2, MMRI, SRL1, TCB3 and TPO1) are translationally up-regulated in caf20Δ cells. GO Slim mapping analysis suggests that bud, cell wall, and site of polarized cell growth-localized proteins are among those enriched in the caf20Δ data set. Together, these observations suggest that Caf20p may

Figure 2. qPCR and immunoblotting confirms translational profiling. (A) Quantitative reverse-transcription PCR analysis of translation state for qPCR (light grey bars ± SD, n = 3) and array data (black bars) for indicated transcripts (plotted on a log2 scale so that reduced polysome association in mutant cells is shown as a negative number). (B) Top: total protein levels of Sec9p, Taf7p, Taf3p, Cic1p, Lsm8p, Pub1p and Ade2p were analysed by immunoblot analysis in indicated yeast strains. Experiments were done in triplicate. Loading control Arp2p (Actin Related Protein 2) is shown beneath each. Middle: densitometry analysis for each protein relative to Arp2p is beneath each panel (dark grey bars). Bottom: polysome/monosome [P/M] ratios from translational profiling (light grey bars).
play a role repressing translation of these mRNAs in the mother cell, but that localized transcripts could become translatable. Bud-localized activation of ASH1 mRNA translation has been studied directly and two alternative hypotheses have been proposed to account for its translational repression involving Khd1p, eIF4G and the kinase Yck1p (32) or involving eIF5B and Puf6p (33). Our data is consistent with and suggestive that the yeast 4E-BPs could play a role in spatial regulation of translation for specific transcripts in yeast but elucidation of molecular mechanisms is beyond the scope of this article.

Nitrogen and amino acid metabolism defects

Filamentous or pseudohyphal growth on poor nitrogen sources in Σ1278b yeast strains requires reprogramming of cellular growth. We recently showed that each 4E-BP was required for this response (10) highlighting a role for translational control in the response to nitrogen limitation. Others found that both CLN1 and STE12 translation is induced following induction of filamentous growth on poor nitrogen sources in Σ1278b yeast (34). Our arrays is consistent with and suggestive that the yeast 4E-BPs could play a role in spatial regulation of translation for specific transcripts in yeast but elucidation of molecular mechanisms is beyond the scope of this article.


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growth on alternate nitrogen sources, while a plasmid bearing missense mutations that disrupt the eIF4E interaction motif in Eap1p (EAP1-mt in Figure 4) shown to eliminate eIF4E binding (10) was significantly impaired in this response, suggesting that eIF4E-binding contributes to this phenotype. These data show that 4E-BP-mediated translational controls are important for normal amino acid metabolism in yeast in addition to their role in pseudohyphal development in \( \Sigma \)1278b yeast cells (10).

Overlap between 4E-BP regulated and PUF-bound mRNAs

One paradigm for the action of 4E-BPs is that translation-repression is mediated in part via protein–protein interactions between proteins bound to the 5′-cap and those attached to the 3′-UTR of specific mRNAs (2). As our array data reveal that Eap1p and Caf20p largely regulate translation of distinct genes (Figure 1), it seemed likely that additional mRNA-binding factors may contribute to the control mechanism. Several studies have used microarrays to identify RNAs associated with specific RNA-binding proteins following immune-precipitation of epitope tagged proteins (RIP-chip). We used statistical comparisons to assess if there were any significant overlaps between the identities of our 4E-BP regulated transcripts and transcripts identified by others as specifically associated with a range of mRNA-binding proteins in different RIP-chip experiments (19,37–39). The question addressed was: are genes identified as enriched with specific mRNA binding proteins also those identified as translationally controlled by the 4E-BPs or are the associations random? This analysis considers each transcript equally and does not account for mRNA abundance or any biases in the data sets. We found that there was a potentially significant overlap between genes up-regulated by caf20\( \Delta \) (i.e. translation state change >0.9) and mRNAs associated with Puf1p, Puf2p, and Puf5p (\( P \leq 0.0001 \)) (Supplementary Table S9). The full distribution is shown graphically in Figure 5, where the translational state change for each PUF-bound mRNA is shown for the caf20\( \Delta \) data. This additionally shows that the mean translational states for Puf1p, Puf2p and Puf5p are each shifted positively away from zero, while the means for Puf3p and Puf4p are not. As Gerber and colleagues found fewer mRNAs associated with Puf1p and Puf2p than Puf3-5p, the apparent enrichment in our caf20\( \Delta \) up-regulated genes associated with Puf5p appeared most significant. Overlaps between genes translationally up-regulated by eap1\( \Delta \) and bound by PUF

Table 1. Maximum doubling times of strains in liquid medium with indicated nitrogen sources

<table>
<thead>
<tr>
<th>Strain</th>
<th>Complete</th>
<th>Serine</th>
<th>Glutamate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild-type</td>
<td>118 ± 2( ^{a} )</td>
<td>148 ± 20</td>
<td>153 ± 10</td>
</tr>
<tr>
<td>Caf20( \Delta )</td>
<td>124 ± 5</td>
<td>182 ± 20</td>
<td>189 ± 21</td>
</tr>
<tr>
<td>eap1( \Delta )</td>
<td>140 ± 15</td>
<td>212 ± 19</td>
<td>233 ± 35</td>
</tr>
</tbody>
</table>

\( ^{a} \)Mean maximum doubling times in min ± SD (\( n = 9 \) cultures) for cultures grown in liquid media equivalent to solid media shown in Figure 4.

Overgrowth on alternate nitrogen sources, while a plasmid bearing missense mutations that disrupt the eIF4E interaction motif in Eap1p (EAP1-mt in Figure 4) shown to eliminate eIF4E binding (10) was significantly impaired in this response, suggesting that eIF4E-binding contributes to this phenotype. These data show that 4E-BP-mediated translational controls are important for normal amino acid metabolism in yeast in addition to their role in pseudohyphal development in \( \Sigma \)1278b yeast cells (10).

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whether there were any interactions between the 4E-BPs and the PUF proteins that could help to explain the polysome microarray data.

4E-BPs physically associate with specific PUFs

PUF proteins are found in diverse eukaryotes (as are 4E-BPs) and have been implicated in post-transcriptional control of gene expression in early development, germ cells and neuronal cells (40,41). PUFs have been implicated both in the control of mRNA stability and translational repression (40). Unlike the 4E-BPs, PUFs are sequence specific RNA-binding proteins that contain eight repeats of an ~40 residue motif, the PUF motif (42). PUF proteins often associate with 3′-UTRs (19) where they can form complexes with other proteins (43). In yeast, Pufs 1 and 2 preferentially bind mRNAs encoding membrane-associated proteins, Pufrp rRNA-processing factor mRNAs and Puf5p chromatin modifier mRNAs (19). Pufrp was found to bind the ASH1 3′-UTR and repress its translation in mother cells. ASH1 mRNA is normally localized to the distal bud tip and both mRNA localization and translation were affected in puf6Δ cells (44). Most studies to date point to a function in mRNA stability, with each PUF protein having distinct targets, although few mRNA targets have been studied in any detail. Pufrp preferentially binds mitochondrial-localized transcripts e.g. COX17 to stimulate the latter’s degradation (45,46). In another study, the stability of 38 potential PUF-target mRNAs was assessed and only two of these (HXK1 and TIF1) were found to have altered mRNA half-lives in pufA strains under the conditions studied (47).

A more recent RIP-chip study has identified mRNAs associated with a wide range of RNA-binding proteins (39). Equivalent comparative statistical analyses revealed potential overlap between caf20Δ up-regulated genes and mRNA-binding proteins including translational regulators (Khd1p, Mrn1p, Sep160p), factors involved in nuclear export of mRNA (Gbp2 and Nab2p) as well as Pub1p and Ssd1p. Similar comparisons with our eap1Δ data revealed overlaps with Nab2, Nrd1p, and the cytoplasmic polyA binding protein Pab1p (Supplementary Table S9). In addition, we found other relationships between the data sets, including correlations between mRNAs bound by specific proteins and those translationally down-regulated by the 4E-BP deletions. For example Mrn1p and Pub1p-bound mRNAs are both enriched among mRNAs down-regulated by eap1Δ and up-regulated by caf20Δ. We also noted that there were examples where fewer than expected mRNAs in common between specific data sets (Supplementary Table S9). While not conclusive, these comparisons suggest that different proteins binding to specific mRNAs could combine with 4E-BPs to regulate their translation in yeast in a manner similar to that proposed for multicellular eukaryotes. Because there are several yeast PUF proteins, we decided to examine experimentally whether there were any interactions between the 4E-BPs and the PUF proteins that could help to explain the polysome microarray data.
Myc-tagged EAP1 and repeated the TAP-affinity chromatography experiments to identify whether any Eap1p-PUF complexes formed (Figure 6B). Remarkably, these experiments revealed a complementary set of interactions, as a fraction of Eap1p was associated with Puf1p and Puf2p, but not the other PUFs tested. As each set of experiments identified different protein complexes it provides evidence that they are not artifactual. Because the Protein A component of the TAP tag bound our Myc antibodies, we were not able to complete reciprocal precipitation experiments. Instead, we made new strains bearing Myc-tagged PUF proteins (PUF2, PUF3, and PUF5) and TAP tagged 4E-BPs and repeated the TAP-affinity chromatography experiments. In agreement with our original experiment, a small fraction of Puf5p-Myc was associated with Caf20p-TAP and Puf2p-Myc with Eap1p-TAP (Supplementary Figure S2). Importantly, no associations were seen between Puf3p-Myc and either 4E-BP-TAP protein or between Myc-tagged proteins and the protein A resin in control strains lacking a TAP tag, although a tiny fraction of Puf2p-Myc was additionally found associated with Caf20p-TAP.

These experimental data do not entirely agree with the in silico predictions. The predicted Caf20p-Puf5p interaction was found. However, the Caf20p-Puf4p and Eap1p-Puf1/2p interactions we identified were not predicted by our statistical comparisons. There are several possible explanations for this including the fact that our statistical treatment did not take into consideration mRNA abundances or the relative affinities of the interactions, as these are not known. Our experiments represent a ‘snap-shot’ of cells from non-stressed log cultures. Under different growth stresses, phases or synchronous states the associations may differ. However, our data has shown that there is value using this type of approach to devise hypotheses to test experimentally. Importantly, when taken together, these experiments indicate that each yeast 4E-BP can form mRNA–protein complexes with specific PUF-bound mRNAs.

In summary, by using a translational profiling approach, we have identified that translation of a significant fraction of yeast mRNAs is regulated by the yeast eIF4E-BPs Caf20p and Eap1p. Many of the regulated transcripts are poorly translated in wild-type cells during exponential growth and these mRNAs are enriched in function in gene expression, cell cycle or transport. In addition to explore how mRNA specificity is achieved, we provide direct evidence that each yeast 4E-BP can interact with specific PUF proteins. As PUF proteins bind RNAs with sequence specificity, this could present one mechanism by which the yeast 4E-BPs target individual mRNAs. By analogy with higher eukaryotes, it is likely that these complexes contain additional components that remain to be determined and that other 3’-UTR-binding protein/4E-BP combinations also await identification. These studies, therefore, provide the first evidence that the yeast 4E-BPs regulate translation of hundreds of mRNAs and that by further study it will be possible to uncover molecular details in yeast that inform studies in higher eukaryotes.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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