A Transcript-Specific eIF3 Complex Mediates Global Translational Control of Energy Metabolism

Highlights
- eIF3d and eIF3e promote synthesis of the mitochondrial ETC and respiration
- eIF3 deficiency leads to glycolytic switch, oxidative stress, and reduced lifespan
- eIF3e promotes the synthesis of ETC proteins through a 5’ UTR-mediated mechanism
- eIF3 subunits interact with mRNAs encoding ETC proteins

In Brief
eIF3 is frequently dysregulated in cancer. Shah et al. show that lack of eIF3d and eIF3e results in impaired synthesis of mitochondrial OXPHOS proteins, respiratory deficiency, glycolytic switch, oxidative stress, and reduced lifespan. Thus, the eIF3d-eIF3e module mediates mRNA-specific translational control of energy metabolism that may be disrupted in cancer.

Accession Numbers
- GSE80349
- PXD004137
- PXD004138
- PXD004146
- PXD004147

Cell Reports
August 16, 2016 © 2016 The Author(s).
http://dx.doi.org/10.1016/j.celrep.2016.07.006
A Transcript-Specific elf3 Complex Mediates Global Translational Control of Energy Metabolism

Meera Shah,1,8 Dan Su,2,8 Judith S. Scheliga,1,8 Tomás Pluskal,6,9 Susanna Boronat,7 Khaterheh Motamedchaboki,3 Alexandre Rosa Campos,3 Feng Qi,4 Elena Hidalgo,5 Mitsuhito Yanagida,1 and Dieter A. Wolf1,2,5,*

1Tumor Initiation and Maintenance Program, Sanford Burnham Prebys Medical Discovery Institute, 10901 North Torrey Pines Road, La Jolla, CA 92037, USA
2School of Pharmaceutical Sciences and Center for Stress Signaling Networks, Xiamen University, Xiang’An South Road, Xiamen 361102, China
3Proteomics Facility, Sanford Burnham Prebys Medical Discovery Institute, 10901 North Torrey Pines Road, La Jolla, CA 92037, USA
4Applied Bioinformatics Core, Sanford Burnham Prebys Medical Discovery Institute, 10901 North Torrey Pines Road, La Jolla, CA 92037, USA
5San Diego Center for Systems Biology, La Jolla, CA 92093, USA
6GO Cell Unit, Okinawa Institute of Science and Technology Graduate University, 1919-1 Tancha, Onna-son, Okinawa 904-0495, Japan
7Universitat Pompeu Fabra, Calle Doctor Aiguader 88, 08003 Barcelona, Spain
8Co-first author
9Present address: Whitehead Institute for Biomedical Research, Cambridge, MA 02142, USA
*Correspondence: dwolf@sbpdiscovery.org
http://dx.doi.org/10.1016/j.celrep.2016.07.006

SUMMARY

The multi-subunit eukaryotic translation initiation factor elf3 is thought to assist in the recruitment of ribosomes to mRNA. The expression of elf3 subunits is frequently disrupted in human cancers, but the specific roles of individual subunits in mRNA translation and cancer remain elusive. Using global transcriptomic, proteomic, and metabolomic profiling, we found a striking failure of Schizosaccharomyces pombe cells lacking elf3e and elf3d to synthesize components of the mitochondrial electron transport chain, leading to a defect in respiration, endogenous oxidative stress, and premature aging. Energy balance was maintained, however, by a switch to glycolysis with increased glucose uptake, upregulation of glycolytic enzymes, and strict dependence on a fermentable carbon source. This metabolic regulatory function appears to be conserved in human cells where elf3e binds metabolic mRNAs and promotes their translation. Thus, via its elf3d-elf3e module, elf3 orchestrates an mRNA-specific translational mechanism controlling energy metabolism that may be disrupted in cancer.

INTRODUCTION

Protein synthesis through mRNA translation is the dominant determinant of cellular protein levels (Schwanhäusser et al., 2011). Translation initiation is considered a rate-limiting step in protein synthesis that is governed by the availability and activity of eukaryotic translation initiation factors (elfs) (Sonenberg and Hinnebusch, 2009). elf3 is the most complex translation initiation factor (Hinnebusch, 2006), comprising 13 subunits in mammals (Damoc et al., 2007; Querol-Audi et al., 2013) and 11 subunits in the fission yeast Schizosaccharomyces pombe (Sha et al., 2009; Zhou et al., 2005). elf3 appears to encircle the 40S ribosome to serve as a scaffold orchestrating the recruitment of other elfs involved in mRNA binding, scanning, and AUG recognition (Erzberger et al., 2014; des Georges et al., 2015; Querol-Audi et al., 2013). For certain mRNAs, the elf3-dependent initiation mechanism involves direct interactions with RNA stem-loop structures or methylated adenosines within the 5’ UTR (Lee et al., 2015; Meyer et al., 2015).

Upregulation of elf3 subunits is frequently observed in human cancers (Hershey, 2015). Overexpression of some subunits can drive de novo holo-complex formation and modest increases in protein synthesis along with cell transformation (Zhang et al., 2007), although the specific mechanisms leading to transformation remain unknown. A recent study suggested that elf3 promotes the synthesis of proteins related to cell proliferation and demonstrated that elf3-mediated synthesis of c-JUN promotes cell migration (Lee et al., 2015).

Not all elf3 subunits are essential, however, suggesting that some subunits have regulatory functions such as mediating the translation of subsets of mRNAs under specific conditions (Choudhuri et al., 2013; Grzmil et al., 2010; Kim et al., 2007; Zhou et al., 2005). For example, elf3e was first identified as a gene disrupted by integration of mouse mammary tumor virus during breast tumorigenesis (Asano et al., 1997; Marchetti et al., 1995), and elf3e is downregulated in several human cancers (Butitta et al., 2005; Hershey, 2015; Marchetti et al., 2001; Suo et al., 2015). Downregulation of elf3e induces epithelial-mesenchymal transition in breast epithelial cells (Gillis and Lewis, 2013) and transdifferentiation of human mesenchymal stem cells into carcinoma-associated fibroblasts (Suo et al., 2015), but the molecular mechanisms underlying the apparent tumor suppressor function of elf3e remain unknown.

As in breast cancer, elf3e is dispensable in fungi, including fission yeast (Smith et al., 2013; Zhou et al., 2005). Cells deleted...
for the gene encoding eIF3e (also known as Yin6p or Int6p) or its binding partner eIF3d (Moe1p) show a 25% reduction in global protein synthesis and growth, and are hypersensitive to stress conditions (Bandyopadhyay et al., 2000, 2002). Although these observations suggested potential mRNA-selective functions, the specific effect of eIF3e on protein synthesis and its role in breast tumorigenesis have remained elusive. We show here that eIF3e and eIF3d form a specificity module for the efficient synthesis of components of the mitochondrial electron transport chain (ETC) and that lack of eIF3d and eIF3e leads to a metabolic switch from respiration to glycolysis, similar to what is frequently observed in cancer cells undergoing the Warburg effect. The data implicate the eIF3d-eIF3e module of eIF3 in a translational circuit to uphold the metabolic balance that may be disrupted in human cancer.

RESULTS

Cells Deleted for eIF3 Subunits e and d Are Deficient in Initiation of Translation

Using sucrose density gradient centrifugation, we observed that eIF3e-deleted cells have a disturbed polysome profile with accumulation of light polysomes (less than five) at the expense of heavy polysomes (more than five; Figure 1A). A strain deleted for eIF3d, encoding a direct interactions partner of eIF3e (Yen and Chang, 2000), has the same growth and stress phenotype (Bandyopadhyay et al., 2002) and showed a polysome profile identical to eIF3e-deleted cells (Figure 1A). The abnormal profile suggested that the mutants are defective in recruiting ribosomes to mRNAs. To determine the efficiency of translation initiation, polysome run-off was induced by removing glucose from the culture medium for 15 min, followed by glucose re-addition to induce initiation of translation. Although wild-type cells efficiently re-established polysomes containing more than five ribosomes within 10 s of glucose re-addition, the bulk of polysomes formed in eIF3e mutant cells within the same period contained fewer than four ribosomes (Figure 1B). This defect, which was even more pronounced at 30 s after glucose re-addition, indicated a reduced efficiency in recruiting ribosomes to mRNAs.

Proteomic Analysis of Ribosome-Associated Proteins Points to a Role of eIF3e in the Synthesis of Mitochondrial ETC Components

To gain further insight into the apparent initiation defect of eIF3e mutant cells, we sought to assess the state of translating ribosomes. Cell lysate from wild-type and eIF3e-deleted cells was digested with RNase, and 80S ribosomes and their associated proteins and nascent polypeptides were isolated and analyzed by liquid chromatography and tandem mass spectrometry (“80S proteomics”; Figure 1C). In five independent experiments, 1,622 proteins were quantified, of which 261 showed a statistically significant difference (p ≤ 0.05) between the wild-type and eIF3e mutant cells (Data S1).
Gene ontology (GO) enrichment analysis (Huang et al., 2009) revealed that the group of 130 proteins that were downregulated in the 80S fraction of eif3e-deleted cells fell into three main categories: ribosome biogenesis, transmembrane transport, and oxidative phosphorylation, in particular subunits of mitochondrial respiration complexes III, IV, and V (Figure 1D; Figure S1A). The eIF3 complex (GO:5852) was also enriched because of minor depletion of eIF3b and h and the virtually complete absence of eIF3d from 80S particles isolated from eif3e-deleted cells (Figure S1B). This finding is consistent with our previous demonstration that S. pombe eIF3 complexes missing subunit e are also devoid of eIF3d (Sha et al., 2009) and suggests that eIF3d and eIF3e are recruited into the complex as a module. At the same time, these data complement our previous evidence that a stable eIF3 complex can exist in the absence of eIF3d and eIF3e (Sha et al., 2009; Zhou et al., 2005).

The list of 131 proteins whose abundance was significantly increased in 80S complexes isolated from eif3e-deleted cells was enriched for amino acid and lipid biosynthesis, transmembrane transport, and glycolysis and its two downstream metabolic pathways alcohol fermentation and tricarboxylic acid (TCA) cycle. Only minor differences were found in ribosomal proteins (Figure S1C), other initiation factors (Figure S1D), and ribosome-associated chaperones (HSP70s, CCT complex), suggesting that the overall 80S structure is intact in the absence of eIF3e.

Cells Deleted for eif3e Are Defective in Synthesizing ETC Proteins
To assess whether the differences in the 80S-associated proteome observed between wild-type and eif3e mutant cells reflected differences in the synthesis of nascent proteins, we acquired a series of datasets on global protein dynamics employing stable isotope labeling techniques coupled with mass spectrometry. Differences in translation rates as determined in a pulsed stable isotope labeling with amino acids in cell culture (SILAC) experiment (Selbach et al., 2008) correlated significantly ($r = 0.40$, $p < 0.001$) with the differences observed in the 80S-associated proteome (Figure 2A). In addition, of the translation rates determined for 2,597 proteins, 228 were significantly reduced in eif3e mutant cells, whereas 187 were significantly increased. The lists of proteins with significantly altered rates of synthesis-enriched GO terms that overlapped with those obtained for proteins with altered 80S association shown in Figure 1D, indicating that the synthesis of components of the mitochondrial respiration chain is downregulated in eif3e-deleted cells, whereas amino acid biosynthesis and glucose metabolism are upregulated (Figure S2A; Data S1). Identical GO terms were also enriched in the set of 694 proteins whose steady-state levels significantly differed in eif3e mutant cells, as determined by quantitative mass spectrometry (Figure S2B; Data S1).

We next directly measured changes in the synthesis rates of 80S-associated proteins in wild-type and eif3e mutant cells by labeling nascent proteins through growing cells in medium containing heavy nitrogen ($^{15}$N), followed by purification of 80S complexes and mass spectrometry. Observing changes in the $^{14}$N/$^{15}$N ratios of the identified peptides allowed us to assess
the rate of synthesis of 232 proteins associated with 80S complexes (Data S1), of which 27 represented metabolism-related proteins that we had found to be altered in their 80S association in eif3e mutant cells. Plotting the protein synthesis rates versus the steady-state levels of these 80S-associated proteins revealed a highly significant correlation ($r = 0.64$, $p < 0.001$; Figure 2B), again indicating that mitochondrial proteins with reduced 80S ribosome association in eif3e mutant cells correspond to those with a reduced rate of synthesis. Conversely, increased association of glycolysis and TCA cycle proteins appears to reflect increased synthesis.

Although there was an overall positive correlation between a protein’s association with 80S complexes and its corresponding mRNA level ($r = 0.45$, $p < 0.001$; Figure 2C), only 19 of the 261 proteins (7.3%) differentially associated with 80S ribosomes in wild-type versus eif3e mutant cells showed a statistically significant change at the mRNA level. In addition, decreased 80S association of 35 mitochondrial proteins was not correlated with their mRNA levels (Figures 2D and 2E). The transcriptomic differences between wild-type and eif3e mutant cells (402 mRNAs; Data S1) did not enrich any of the metabolism-related GO terms found in the proteomic datasets (80S proteome, pulsed SILAC (pSILAC), steady-state dataset; Figure S2C). Instead, there was marginally significant enrichment of the terms transmembrane transport and cellular stress response (Figure S2C). In addition, thiamine biosynthesis was enriched in the set of 218 mRNAs that were upregulated in eif3e mutant cells (Figure S2C). Comparison with our parallel proteomic datasets revealed that the corresponding biosynthetic enzymes as well as the thiamine transporter Thi9p were also upregulated at the protein level (Data S1), suggesting that eif3e mutant cells have an increased demand for thiamine, a vitamin essential for glucose oxidation at multiple levels. In summary, the integrated analysis of 80S proteomics, pSILAC, steady-state protein, and mRNA datasets strongly suggested that the highly selective changes in the levels of metabolism-related proteins observed in the absence of eIF3e are primarily driven by changes at the level of translating specific classes of transcripts.

**Metabolomic Analysis Indicates a Glycolytic Switch in eIF3d- and eIF3e-Deficient Cells**

To assess phenotypic consequences of the translational changes observed in eif3e mutants, we considered the possibility that the abnormal polysome profile may result from a defect in ribosome biogenesis. However, eif3e mutants did not display typical phenotypic features of ribosome biogenesis mutants, such as flocculation (data not shown) and “ribosome halfmers” (Li et al., 2013; Meyer et al., 2007), in the sucrose density gradient profile (Figure S3A). Cells deleted for eif3e did, however, display a 10%–20% reduction in total RNA levels, the bulk of which consists of ribosomal RNA (Figure S3B). Thus, although ribosome biogenesis may be slightly reduced in eif3e mutant cells, these data suggest that this is a consequence rather than the cause of the reduced growth rate of the mutant.

With respect to the large energy requirements of protein synthesis in exponentially growing cells—up to 30% of the cellular ATP utilization (Rolfe and Brown, 1997)—we considered the possibility that failure to maintain the synthesis of mitochondrial respiration chain components might lead to a deficiency in ATP, thereby causing inefficient protein synthesis. Exponentially growing *S. pombe* cells are known to derive ~50%–65% of their energy from mitochondrial respiration (Zuin et al., 2008). Consistent with the proteomic data, cells deleted for eif3e maintained under exponential growth conditions showed an ~60% reduction in mitochondrial oxygen consumption rate (Figure 3A). Similar reductions were observed with strains deficient in two other non-essential subunits, eIF3d and eIF3h (Tif85). Deletion of eIF3j (Hcr1), a sub-stoichiometric subunit thought to have modulatory function (Wagner et al., 2014), had no effect on respiration (Figure 3A).

To determine whether reduced respiration coincides with reduced ATP levels, we generated a comprehensive metabolomic profile of wild-type and eif3e-deficient cells by liquid chromatography (LC)-mass spectrometry (MS). In triplicate experiments, ATP, ADP, and AMP levels did not significantly vary between wild-type and eif3e-deleted cells (Figure 3B), indicating that the translational phenotype of eif3e mutant cells is not due to ATP depletion. We then used the MetaboAnalyst tool suite (Xia et al., 2015) to identify a set of 90 metabolites (of over 8,000 LC-MS peaks) whose levels were significantly changed in eif3e mutant cells (Data S2). Pathway analysis of this metabolite set integrated with the list of significantly altered proteins identified by quantitative proteomics (Data S1) enriched numerous highly significant Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways (Figure 3C). The analysis revealed glucose utilization (i.e., glycolysis, pyruvate metabolism, TCA cycle), amino acid metabolism, and nucleotide metabolism as significantly upregulated KEGG pathways. Likewise, oxidative stress defense functions (glutathione metabolism and pentose phosphate pathway) were upregulated in eif3e-deleted cells, a finding that is consistent with the well-established stress phenotype of eIF3e mutant cells (Bandyopadhyay et al., 2000; Udagawa et al., 2008) as well as with our transcriptomic profile that indicated upregulation of oxidative stress response genes in the mutant (Figure S2C). Consistent with a role of eIF3 in promoting the synthesis of the respiration chain, disruption of the ETC has previously been shown to lead to endogenous oxidative stress in *S. pombe* (Zuin et al., 2008). Biosynthesis pathways of several vitamin coenzymes essential for glucose, amino acid, and lipid metabolism was also found upregulated in cells lacking eIF3e (i.e., biotin, pyridoxal 5’-phosphate, thiamine). The apparent high demand for thiamine of eIF3e mutant cells is consistent with the upregulation of the thiamine transporter Thi9p noted above.

**Cells Deleted for eIF3e and eIF3d Are Dependent on Glucose**

The combined proteomic and metabolomic data suggested that *S. pombe* cells lacking eIF3e function undergo a metabolic switch from respiration (which is downregulated) to glycolysis (which is upregulated) to maintain energy balance. To test this hypothesis, we determined glucose utilization by measuring the depletion of glucose from the medium. Glucose uptake was significantly increased in cells lacking eIF3e (Figure 4A). This finding is consistent with the pronounced induction of the high-affinity glucose transporter Ght5p (Saitoh et al., 2015) (Figure 2G), which was more than 60-fold higher in the 80S fraction.
isolated from eif3e mutant cells (Data S1). Using RT-PCR, we assessed the presence of ght5 mRNA across the fractions of a sucrose density gradient. Although, in wild-type cells, a significant portion of the transcript was present in monosomal fractions, ght5 mRNA was quantitatively sequestered in polysomes of eif3e-deleted cells, indicating strongly increased translation of ght5 mRNA in the mutant (Figure 4B).

To test whether increased glucose uptake is relevant to cell physiology, we determined growth under glucose-limiting conditions. Although yeast extract growth medium (yeast extract supplements [YES]) typically contains 3% glucose, wild-type cells can maintain growth at glucose concentrations as low as 0.2% (Takeda et al., 2015). However, the growth of eif3e and eif3d mutant cells was inhibited at glucose concentrations below 1.5% (Figure 4C). At 0.2%, both eif3 mutants were inviable (Figure 4C). In Edinborough minimal media (EMM), which contains 2% glucose and in which cells have an increased reliance on respiration (Zuin et al., 2008), eif3 mutant cells grew very poorly (Figure 4D). Growth improved, however, 5- to 25-fold upon raising the glucose concentration to 5% (Figure 4D). Cells deficient in eif3e and eif3d were entirely unable to grow in medium containing non-fermentable glycerol as the carbon source (Figure 4D), confirming that they are severely impaired in respiration.

Figure 3. Metabolic Re-programming in eif3-Deficient Cells
(A) The oxygen consumption rate was determined in the indicated WT and eif3 mutant strains. Two different WT strains were used that are syngeneic with the eif3-deleted strains to their right. Error bars represent SDs of three (WT, eif3e, eif3d, eif3h, eif3j) or four (WT, coq4) independent measurements. p Values correspond to the results of a t test with two-tailed distribution and unequal variance.
(B) eif3e/WT ratios of the indicated metabolites as determined by LC-MS/MS in triplicates and statistical analysis in MetaboAnalyst. Values are the results of a t test.
(C) KEGG pathways enriched in the sets of 90 metabolites and 521 proteins that significantly differed between wild-type and eif3e mutant cells. Plotted are the Z scores for pathway enrichment and pathway topology analysis derived from the MetaboAnalyst 3.0 tool suite (Xia et al., 2015). Because this type of integrated metabolite and protein expression analysis is only available for mammalian data, the putative human orthologs of the significantly changed S. pombe proteins were used for the analysis.

Because eIF3 appears to promote the synthesis of ETC proteins, eIF3 subunits might be induced under conditions of increased respiration. Mitochondrial respiration is upregulated by shifting S. pombe cells to 0.08% glucose, a response that coincides with increased levels of the complex I subunit Cox2p (Takeda et al., 2015), which we found to require eIF3e for efficient synthesis (Data S1). Under the same glucose-limiting conditions, both eIF3 subunits we tested, eIF3e and eIF3m, were upregulated between 30%-40% after 4 hr (Figure 4E), suggesting that eIF3 contributes to the upregulation of respiration upon glucose restriction.

Lack of eIF3e and eIF3d Causes Endogenous Oxidative Stress and Premature Aging
Consistent with the described stress sensitivity of eif3e and eif3d mutant cells (Bandyopadhyay et al., 2000; Udagawa et al., 2008), we found that these cells are sensitive to oxidative stress administered by the presence of 1 mM hydrogen peroxide in the medium (Figure 5A). To test whether poor growth on EMM is due to endogenous oxidative stress, from which respiration-deficient cells are known to suffer (Zuin et al., 2008), we supplemented the medium with the anti-oxidant N-acetyl cysteine (NAC). NAC was able to substantially rescue the growth of eif3e and eif3m mutant cells on EMM, leading to a minimum 25-fold improvement (Figure 5B). These phenotypic data, combined with the mRNA data indicating induction of a stress response signature, strongly suggest that the defect in mitochondrial respiration observed in eIF3-deficient cells leads to leakage of electrons from the ETC and subsequent oxidative stress.
Because *S. pombe* cells with impaired mitochondrial function are known to display defects during postdiauxic stationary phase (Zuin et al., 2008), we determined the survival of *eif3e*- and *eif3d*-deleted cells in stationary phase, which is a read-out of chronological aging (Roux et al., 2006). Studies in *Saccharomyces cerevisiae* established a “respiratory threshold” below which chronological longevity was drastically curtailed (Ocampo et al., 2012). Conversely, enhanced synthesis of ETC proteins was observed in *eif3e* and *eif3d*-deleted cells, suggesting a role for the eIF3 complex in regulating mitochondrial function.

Figure 4. Effect of eIF3 Deficiency on Glucose Uptake

(A) Glucose uptake was measured by determining the depletion of glucose from the medium during the growth of WT and *eif3e*-deleted cells using a YSI 2950 metabolite analyzer. Glucose concentration in the medium was normalized to cell growth (i.e., optical density 600 [OD600]) to account for the slower growth rate of *eif3e*-deleted cells. Data are averages of three biological replicates, with error bars indicating SDs. Statistical significance was assessed with a t test assuming two-tailed distribution and unequal variance.

(B) RNA was purified from sucrose density gradient fractions prepared from WT and *eif3e* mutant cells and employed in RT-PCR reactions with primers specifically amplifying the mRNA encoding the hexose transporter *ght5* and *erg2* as a reference. Only fractions containing mRNA (3–12) were used for RT-PCR. Total RNA was used as a reference. Reactions excluding reverse transcriptase are shown to prove that the band shown is derived from RNA and not DNA.

(C) Growth of *eif3e*- and *eif3d*-deleted cells under limiting glucose conditions. 5-fold serial dilutions of the indicated strains were plated onto YES medium containing decreasing glucose concentrations, and growth was scored after 2–4 days.

(D) Growth of *eif3e*- and *eif3d*-deleted cells on EMM containing increasing concentrations of glucose or on glycerol as the sole carbon source.

(E) Effect of glucose restriction on the levels of eIF3e and eIF3m proteins. Strains harboring alleles of *eif3e* and *eif3m* modified with protein A epitope tags (Zhou et al., 2005) were shifted from medium containing 3% glucose to medium containing 0.08% glucose for the indicated times. Cell lysates were loaded in duplicates and blotted with anti-protein A antibodies to detect eIF3e and eIF3m, respectively. Bands obtained with anti-PSTAIR antibodies were used as a loading reference (bottom). Blots from biological replicates were quantified with the LI-COR Biosciences Image Studio Lite package, and results corrected for the loading reference were plotted. Statistical significance was assessed with a t test assuming two-tailed distribution and unequal variance. *p < 0.05.*
components prolongs the lifespan in *Drosophila* under conditions of dietary restriction (Zid et al., 2009). Accordingly, we found that, after 5 days in the postdiauxic phase, respiration-deficient *S. pombe* cells deleted for *eif3e* or *eif3d* had a 4- to 5-fold reduction in viability, as determined by the ability to form colonies (Figure 5C). Likewise, using exclusion of propidium iodide as a marker of cell integrity, cells deleted for *eif3e* or *eif3d* showed an ~5-fold reduction in survival (Figure 5D). In summary, these data indicate an important metabolic homeostatic function of the eif3d-eif3e module in cellular physiology and aging.

**eIF3e Promotes the Synthesis of Electron Transport Proteins in Human Breast Cells**

Because *S. pombe* Int6p and human eIF3e are functional orthologs (Crane et al., 2000), we asked whether human cells also rely on eIF3e for maintaining ETC integrity. Knockdown of eIF3e with two distinct small interfering RNAs (siRNAs) in MCF7 breast cancer cells and in non-tumorigenic MCF10A cells led to a marked downregulation of UQCRB and ATP5H (Figures 6A and 6B), two complex III and V subunits whose *S. pombe* orthologs, ubiquinol-cytochrome-c reductase complex subunit 6 (SPCC737.02c) and F0-ATPase subunit D (SPBC29A10.13) require eIF3e for efficient synthesis (Data S1). The complex II subunit SDHB was also downregulated in eIF3e-depleted MCF10A cells, whereas the complex I subunit NDUF8 is not affected by knockdown of eIF3e (Figure 6D; Figure S5D). eIF3 also bound the mRNA encoding the glycolytic enzyme GAPDH, whose *S. pombe* orthologs are upregulated in the absence of eIF3e (Data S1). In summary, these data indicate that the function of eIF3e in maintaining the synthesis of mitochondrial ETC components may be conserved both in yeast and mammalian cells.

However, we could not detect a specific interaction of eIF3 with SDHB mRNA (Figure 6E) even though SDHB protein and reporter activity were decreased in eIF3e knockdown cells (Figures 6A and 6D; Figure S5A). Conversely, although we detected robust interaction with mRNA encoding the proteasome subunit PSMB6 (Figure 6E; Figure S5D), eIF3 also bound the mRNA encoding the glycolytic enzyme GAPDH, whose *S. pombe* orthologs are upregulated in the absence of eIF3e (Data S1). In summary, these data indicate that the function of eIF3e in maintaining the synthesis of mitochondrial ETC components may be conserved both in yeast and mammalian cells.

However, we could not detect a specific interaction of eIF3 with SDHB mRNA (Figure 6E) even though SDHB protein and reporter activity were decreased in eIF3e knockdown cells (Figures 6A and 6D; Figure S5A). Conversely, although we detected robust interaction with mRNA encoding the proteasome subunit PSMB6 (Figure 6E; Figure S5D), a finding consistent with recent photoactivatable ribonucleoside-enhanced crosslinking and immunoprecipitation (PAR-CLIP) data (Meyer et al., 2015), its 5’ UTR did not mediate detectable eIF3e dependence in the reporter assay (Figure 6D). It is therefore likely that mechanisms in addition to mRNA binding mediate the effect of eIF3 on translation.

**DISCUSSION**

**Control of Energy Metabolism at the Level of mRNA Translation**

Compared with transcriptional pathways (Scarpulla et al., 2012), little is known about the control of mitochondrial biogenesis at
Recent studies implicated the mTORC1/4E-BP/eIF4E axis in augmenting the synthesis of ETC subunits as well as mitochondrial ribosomal proteins (Gan-din et al., 2016; Morita et al., 2013; Truitt et al., 2015). The findings presented here identified the eIF3 holo-complex as another critical promoter of the synthesis of ETC components and mitochondrial ribosomal proteins. Lack of this function, as apparent in cell deleted for eif3d and eif3e, causes respiratory deficiency and oxidative stress, a compensatory upregulation of glycolysis, strong dependence on glucose, and a reduced lifespan (Figure 7). These biochemical and cell biological observations are bolstered by negative genetic interactions of deletion of eif3e with the deletion of mitochondrial proteins as well as suppression of the eif3e mutant phenotype by overexpression of glycolytic and TCA cycle enzymes (Data S3). In response to glucose limitation, cells upregulate eIF3e and eIF3m, and probably the entire eIF3 holo-complex, in an apparent attempt to maximize the synthesis of respiration chain components. eIF3 function may therefore be a central node in the plasticity of cellular energy metabolism.

Transcript-Selective Control by eIF3: eIF3d-eIF3e as a Specificity Module

Our results show that cells deficient in eIF3e and eIF3d are not globally defective in protein synthesis because they are translating the compensatory response mRNAs encoding glycolytic enzymes and the glucose transporter Ght5p with greater efficiency than wild-type cells. Our findings thus strengthen our previous finding that S. pombe harbors two distinct eIF3 complexes distinguished by the presence or absence of the eIF3d-eIF3e dimer (Sha et al., 2009; Zhou et al., 2005). Consistent with this model, our quantitative mass spectrometry data show...
that a quasi-stoichiometric eIF3 complex containing all subunits except eIF3d is stably associated with 80S complexes in cells deleted for eIF3e (Figure S1B). The stability of this alternative complex, referred to as eIF3<sup>d/e</sup> (Figure 7), likely accounts for the viability of cells lacking eIF3d or eIF3e, whereas most other eIF3 subunits are essential. Together with the exact overlap of their mutant phenotypes, their direct physical interaction, and their mutual dependency for proper protein levels and recruitment into the eIF3 holo-complex (Sha et al., 2009; Yen and Chang, 2000), it is likely that eIF3d and eIF3e together serve as a specificity module to drive the synthesis of mitochondrial proteins.

How might the eIF3d-eIF3e module regulate specific mRNAs? Using RNA immunoprecipitation and microarray hybridization (RIP-CHIP), we have reported previously that eIF3 subunits interact with specific sets of mRNAs (Zhou et al., 2005). Re-examination of these datasets prompted by the 80S proteomics data presented here revealed a significant enrichment of the gene ontology term “mitochondrial part” (GO:0044429) within the list of the 100 mRNAs associating most abundantly with eIF3e (Data S4). Indeed, 12 of the top 100 mRNAs encode mitochondrial proteins, including mitochondrial ribosomal proteins as well as complex I, II, IV, and V subunits (Data S4). In this work, we found that the interaction of eIF3 with mRNAs encoding ETC components is conserved in human cells. Likewise, recent PAR-CLIP data (Lee et al., 2015; Meyer et al., 2015) confirmed direct binding of eIF3 subunits to unique sets of mRNAs that we found to be highly enriched in the gene ontology terms “mitochondrion organization” (GO:0007005, false discovery rate [FDR] = 10<sup>-22</sup>) and “respiratory electron transport chain” (GO:0022904, FDR = 10<sup>-20.2</sup>) (Data S5). In conjunction with our demonstration that knockdown of eIF3e in human breast cells leads to downregulation of several ETC proteins, eIF3 likely promotes the synthesis of mitochondrial ETC components through direct interaction with their encoding mRNAs.

The efficient synthesis of Ght5p and glycolytic enzymes in eIF3e-deleted cells ostensibly relies on the alternative eIF3<sup>d/e</sup> complex, which we and others have shown to account for ~75% of the total eIF3 population in <i>S. pombe</i> (Ray et al., 2008; Sha et al., 2009; Zhou et al., 2005). Remarkably, the translational efficiency of ght5 mRNA is even higher in the absence of eIF3e than in its presence because it is quantitatively sequestered into polysomes in eIF3e-deleted cells (Figure 4B). This suggests that homo-eIF3, which contains subunits d and e, actively represses ght5 mRNA translation. Our previous RIP-CHIP data (Zhou et al., 2005) revealed that eIF3e interacts with mRNAs for which we now show either decreased (e.g., ETC components) or increased (e.g., glucose transporters) translation in the absence of eIF3e (Data S4). Significantl, Lee et al. (2015) recently reported that eIF3 binding to two distinct mRNAs can promote or inhibit their translation. Regardless, through dual stimulatory and inhibitory activities that remain to be defined molecularly, transcript-specific functions of eIF3 subunits appear to fine-tune energy metabolism at the level of protein synthesis.

**eIF3 and Cancer Metabolism**

As with other eIFs, upregulation of eIF3 subunits is frequently observed in human cancers (Hershey, 2015). Overexpression of individual subunits can drive de novo holo-complex formation and modest increases in global protein synthesis along with cell transformation (Zhang et al., 2007). With the re-emerging focus on the essential role of the mitochondrial ETC in cell proliferation, transformation, cancer stem cell maintenance, metastasis, and...
drug resistance (Birsoy et al., 2015; De Luca et al., 2015; LeBlue et al., 2014; Martinez-Outschoorn et al., 2014; Sullivan et al., 2015; Tan et al., 2015; Truitt et al., 2015; Wolf, 2014), it is conceivable that the ETC-promoting function of eIF3 discovered in this report underlies, at least in part, its tumor-promoting activity. In this model, eIF3 would be limiting for mitochondrially produced building blocks that fuel cancer cell anabolism and thus represent an intriguing cancer drug target.

Unlike most eIF3 subunits that are upregulated in cancer, eIF3e mRNA has been found to be downregulated in bulk samples of human breast and lung cancers (Butittta et al., 2005; Marchetti et al., 2001). A recent study clarified that eIF3e protein downregulation in breast cancer occurs not in the epithelial carcinoma cells but in stromal cancer-associated fibroblasts (CAFs) (Suo et al., 2015). In fact, knockdown of eIF3e is sufficient to confer CAF-like properties to normal human mammary fibroblasts (Suo et al., 2015). In situ studies of human breast cancer samples have revealed that CAFs have low oxidative phosphorylation (OXPHOS) activity but are highly glycolytic, whereas epithelial cancer cells show the opposite metabolic profile, a phenomenon referred to as the “reverse Warburg effect” (Martinez-Outschoorn et al., 2014; Whitaker-Menezes et al., 2011). In this scenario, CAF catabolism and glycolysis will provide lactate and pyruvate to fuel the oxidative metabolism known to drive the tumor cells (Birsoy et al., 2015; De Luca et al., 2015; Martinez-Outschoorn et al., 2014; Sullivan et al., 2015; Tan et al., 2015; Wolf, 2014). Thus, based on our discovery of eIF3 as a promoter of ETC synthesis, the downregulation of eIF3e as well as all other eIF3 subunits (Finak et al., 2008; data not shown) specifically in CAFs may underlie metabolic synergy in the breast cancer microenvironment. If so, eIF3 inhibition as a therapeutic strategy may need to be approached with caution, at least in breast cancers showing the reverse Warburg profile.

EXPERIMENTAL PROCEDURES

Yeast Strains

Strains deleted for eIF3d and eIF3e were described before (Yen and Chang, 2009). The sks2 gene was replaced with the NAT gene conferring resistance to nourseothricin. S. pombe strains were maintained in YES or EMM unless otherwise noted.

Sucrose Density Gradient Profiling

Cell lysates prepared in polysome lysis buffer (20 mM Tris-HCl [pH 7.5], 50 mM KCl, 10 mM MgCl2, 1 mM DTT, and 100 μg/ml cycloheximide) were separated on 10%–50% (w/v) sucrose gradients. For immunoblotting, fractions were mixed with SDS sample buffer. For RNA isolation, collected fractions were precipitated in 1 volume of isopropanol at –80°C, followed by purification via the RNeasy kit (QIAGEN).

Preparation of 80S Complexes for LC-MS/MS Analysis

Wild-type and eIF3e-deleted cells in exponential growth phase were lysed in a buffer containing 20 mM Tris (pH 8), 140 mM KCl, 5 mM MgCl2, 100 μg/ml cycloheximide, 1% Triton X-100, 200 μg/ml heparin, and 1 mM PMSF and digested with RNase A. The lysates were loaded onto 25% sucrose cushions, and ribosomal pellets were analyzed by liquid chromatography-tandem mass spectrometry (LC-MS/MS) (Brill et al., 2009).

For analysis of newly synthesized components, wild-type and eIF3e-deleted cells were switched from YES to EMM containing 15N-NH4Cl as the nitrogen source for 4 hr, followed by purification of 80S complexes and LC-MS/MS analysis. Statistically significant differences in the rate of synthesis of 80S-associated proteins between wild-type and eIF3e mutant cells were derived from calculating 15N-WT and 15N-eIF3e protein ratios from 15N-labeled peptides using IP2-Integrated Proteomics Pipeline software.

Quantitative Proteomic Profiles

Total lysates of wild-type and eIF3e-deleted cells were digested with trypsin, and peptides were separated by 2D-LC and analyzed on an Orbitrap Velos Pro mass spectrometer (Thermo Fisher Scientific) operated in positive data-dependent acquisition mode. All mass spectra were analyzed with MaxQuant software version 1.5.0.25 (Cox and Mann, 2009). MS/MS spectra were searched against the Pombase protein sequence database ASM294 (version 2.23) using settings described in the Supplemental Experimental Procedures.

For pulsed SILAC, wild-type and eIF3e-deleted cells were grown to exponential phase in YES medium, washed, and switched to EMM containing 30 mg/ml isotopically labeled 13C6 15N2 lysine (K8). Cell lysate was prepared after 2 hr and analyzed by 2D-LC-MS/MS as described above. MaxQuant was used for quantifying K8-labeled peptides. Statistically significant differences in the rate of synthesis of individual proteins between wild-type and eIF3e mutant cells were derived from calculating K8-WT and K8-eIF3e protein ratios from K8-labeled peptides.

RNA-Seq Analysis

Total RNA was extracted by phase separation using RNA-Bee (Amsbio) and chloroform, and samples were treated with the Ribominus eukaryotic kit (Invitrogen) to deplete rRNA. The rRNA-depleted sample was subjected to enzymatic fragmentation and used for cDNA library construction. The cDNA was PCR-amplified and sequenced on the SOLiD 4 platform. Differentially expressed genes were called using the edgeR package.

Metabolomic Profiling by LC-MS

Metabolomic analysis was performed as described previously (Phuskal et al., 2013). Briefly, wild-type and eIF3e-deleted cells were quenched in methanol and disrupted using a Multi-Beads Shocker (Yasui Kikai). Proteins were removed by centrifugal filtration, and samples were concentrated by vacuum evaporation. Finally, each sample was re-suspended in 40 μl of 50% acetonitrile, and 1 μl was used for LC-MS analysis on a Paradigm MS4 high-performance liquid chromatography (HPLC) system (Michrom Bioresources) coupled to an LTQ Orbitrap mass spectrometer (Thermo Fisher Scientific). The data were analyzed using MetaboAnalyst (http://www.metaboanalyst.ca/). Pathway enrichment analysis was done by hypergeometric test, topology analysis was based on degree centrality, and a combined gene-metabolite mode was used.

Studies in Mammalian Cells

eIF3e was knocked down in MCF7 and MCF10A cells using siRNA transfection. The levels of eIF3e and ETC proteins were analyzed by immunoblotting as described in the Supplemental Experimental Procedures. mRNA levels were determined by qPCR. For reporter gene assays, constructs fusing the 5’ UTRs of ETC-encoding mRNAs were transfected into MCF10A cells upon knockdown of eIF3e. Luciferase activity was normalized to luciferase mRNA measured by qPCR. eIF3-mRNA interactions were tested by immunoprecipitation of eIF3 complexes with eIF3e and eIF3c antibodies, followed by mRNA extraction and detection by qPCR.

Statistical Analysis

Statistical analysis of replicate datasets was performed with Microsoft Excel. Typically, data were averaged, SDs were calculated, and statistical significance was assessed using the T-test function, assuming two-tailed distribution and unequal variance. The statistical analysis of proteomic and transcriptomic data-sets is detailed in the Supplementary Experimental Procedures section.

ACCESSION NUMBERS

The accession number for the RNA-seq data reported in this paper is GEO: GSE80349. The accession numbers for the proteomic datasets reported in this paper are Pride: PXDD04137, PXDD04138, PXDD04146, and PXDD04147.
SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, five figures, two tables, and two data files and can be found with this article online at http://dx.doi.org/10.1016/j.celrep.2016.07.006.

AUTHOR CONTRIBUTIONS

M.S. performed RT-PCR experiments including sucrose density gradient fractions, stress sensitivity, aging experiments, and all Luciferase reporter assays. D.S. performed the elf3L knockdown experiments in human cells and the RNA-IP experiments. J.S.S. did polysome profiling, prepared 80S complexes for mass spectrometry and RNA samples for RNA-seq, and performed glucose sensitivity experiments. T.P. and M.Y. performed the metabolomic studies. S.B. and E.H. measured respiration in various elf3-deleted strains. K.M. and A.R.C performed all LC-MS/MS experiments. F.Q. performed the bioinformatics analysis of RNA-seq data. D.A.W. conceived the project; performed the pSILAC experiment; determined the effect of glucose withdrawal on elf3L levels; performed the analysis of global proteomic, transcriptomic, and metabolomic datasets; prepared the figures; and wrote the manuscript.

ACKNOWLEDGMENTS

We thank Eric Chang for yeast strains, elf3d antibodies, and discussions. Francois Bachand is thanked for Rps2p antibodies. This work was supported by Grant GM105802 (to D.A.W.). Parts of this work were funded by P30 Grants CA030199 and GM085764. D.A.W. is a scholar of the Foreign 1000 Talent Program funded by the Government of the People’s Republic of China.

Received: November 2, 2015

Revised: May 19, 2016

Accepted: July 3, 2016

Published: July 28, 2016

REFERENCES


Bandyopadhyay, A., Lakshmanan, V., Matsumoto, T., Chang, E.C., and Mastra, U. (2002). Moe1 and splint6, the fission yeast homologues of mammalian translation initiation factor 3 subunits p66 (elf3d) and p48 (elf3e), respectively, are required for stable association of elf3 subunits. J. Biol. Chem. 277, 23660–23677.


