Widespread Regulation of Translation by Elongation Pausing in Heat Shock

The MIT Faculty has made this article openly available. Please share how this access benefits you. Your story matters.

<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>As Published</td>
<td><a href="http://dx.doi.org/10.1016/j.molcel.2012.11.028">http://dx.doi.org/10.1016/j.molcel.2012.11.028</a></td>
</tr>
<tr>
<td>Publisher</td>
<td>Elsevier B.V.</td>
</tr>
<tr>
<td>Version</td>
<td>Final published version</td>
</tr>
<tr>
<td>Citable Link</td>
<td><a href="http://hdl.handle.net/1721.1/96262">http://hdl.handle.net/1721.1/96262</a></td>
</tr>
<tr>
<td>Terms of Use</td>
<td>Article is made available in accordance with the publisher's policy and may be subject to US copyright law. Please refer to the publisher's site for terms of use.</td>
</tr>
<tr>
<td>Detailed Terms</td>
<td></td>
</tr>
</tbody>
</table>
Widespread Regulation of Translation by Elongation Pausing in Heat Shock

Reut Shalgi, Jessica A. Hurt, Irina Krykbaeva, Mikko Taipale, Susan Lindquist, and Christopher B. Burge

1Department of Biology
2Department of Biological Engineering
Massachusetts Institute of Technology, Cambridge, MA 02142, USA
3Whitehead Institute for Biomedical Research, Cambridge, MA 02142, USA
4Howard Hughes Medical Institute
Correspondence: cburge@mit.edu
http://dx.doi.org/10.1016/j.molcel.2012.11.028

SUMMARY

Global repression of protein synthesis is a hallmark of the cellular stress response and has been attributed primarily to inhibition of translation initiation, although this mechanism may not always explain the full extent of repression. Here, using ribosome footprinting, we show that 2 hr of severe heat stress triggers global pausing of translation elongation at around codon 65 on most mRNAs in both mouse and human cells. The genome-wide nature of the phenomenon, its location, and features of protein N termini suggested the involvement of ribosome-associated chaperones. After severe heat shock, Hsp70’s interactions with the translational machinery were markedly altered and its association with ribosomes was reduced. Pretreatment with mild heat stress or overexpression of Hsp70 protected cells from heat shock-induced elongation pausing, while inhibition of Hsp70 activity triggered elongation pausing without heat stress. Our findings suggest that regulation of translation elongation in general, and by chaperones in particular, represents a major component of cellular stress responses.

INTRODUCTION

The cellular response to stress involves changes to many levels of gene regulation, including transcription, messenger RNA (mRNA) processing, and translation (Biamonti and Caceres, 2009; Gibson, 2008; Holcik and Sonenberg, 2005). Stress response pathways play important developmental and evolutionary roles, contributing to developmental robustness in varying environments (Akerfelt et al., 2010; Gibson, 2008; Jarosz and Lindquist, 2010; Lindquist, 2009). The heat shock response is one of the best-characterized stress response pathways, during which heat shock proteins (HSPs), a class of molecular chaperones, are upregulated in response to widespread protein misfolding (Richter et al., 2010). Misfolding stress and the heat shock response pathway in particular play specific developmental roles and are implicated in a variety of diseases. Upregulation of chaperones is frequently observed in cancer, and chaperone inhibitors hold promise as antitumor agents (Calderwood et al., 2006; Whitesell and Lindquist, 2005). A number of studies monitoring incorporation of labeled amino acids have observed that protein synthesis is globally inhibited under various types of stresses, including heat shock (Bouche et al., 1979; Lindquist, 1980).

Both conserved and lineage-specific mechanisms are used to repress protein synthesis generally, while facilitating the expression of HSPs and other stress related proteins (Lindquist, 1980, 1981; Miller et al., 1979). Various mechanisms leading to global translational repression and selective upregulation have been reported, generally involving regulation of translation initiation. Primary ways in which cells inhibit translation initiation globally are repression of cap recognition downstream of the mTOR pathway and repression of ternary complex recycling through phosphorylation of the initiation factor eIF2α (reviewed by Sonenberg and Hinnebusch, 2009; Spriggs et al., 2010). Both of these pathways are known to mediate inhibition of translation initiation in the response to heat stress (Duncan and Hershey, 1984; Vries et al., 1997). However, the extent of eIF2α phosphorylation, eIF4E dephosphorylation, and sequestration by 4EBPs that occurs under heat stress is relatively mild, and it has been noted that the regulation of these pathways cannot explain the degree of translational repression observed (Spriggs et al., 2010). Hence, there are still questions about how the full extent of translational repression is achieved during heat shock.

Here we investigated the genome-wide regulation of translation in response to heat shock using ribosome footprint profiling (Ingolia et al., 2009; Ingolia et al., 2011), which maps the locations of ribosomes on mRNAs at nucleotide resolution. Our analysis revealed widespread changes in translational regulation, including an unexpected regulatory response, in which translation elongation is globally paused after translation of ~65 amino acids. Exploration of the mechanism underlying this mode of translational regulation pointed to involvement of the Hsp70 family of chaperones.

RESULTS

Ribosomes Accumulate in the First 200 Bases of ORFs after Severe Heat Shock

To better understand translational regulation genome-wide during chronic and acute heat stress in mammalian cells, we
used ribosome footprint profiling to globally map the locations of individual ribosomes along mRNAs (Ingolia et al., 2009; Ingolia et al., 2011), in conjunction with RNA sequencing (RNA-seq) to assess mRNA abundance. Ribosome footprint and RNA-seq libraries were prepared from NIH 3T3 mouse fibroblasts under normal growth conditions (control, 37°C) and after 8 hr of mild (42°C, HS8M, chronic heat stress) or 2 hr of severe heat stress (44°C, HS2S, acute stress) (Figures S1A–S1E available online). Neither of these conditions induced significant cell death, and cells appeared to fully recover 24 hr after withdrawal from heat stress (Figure S1F). Changes in levels of total and ribosome-associated mRNA were observed for many genes following both mild and severe heat stress (Figure S1A). HSPs showed a significant upregulation in their translation levels under both conditions, as measured by footprint RPKM (Figures S1G–S1I). Analytical polysome profiling revealed a substantial decrease in the proportion of heavy polysomes and the accumulation of monosomes in both mild and severe heat stress (Figure 1A),

Figure 1. Heat Shock Induces a Global Increase in Ribosome Occupancy around 5' Ends of Open Reading Frames
(A) Polysome profiles of mouse fibroblasts (3T3 cells) under normal growth conditions (blue) or after mild heat shock (42°C for 8 hr, green) or severe but not lethal heat shock (44°C for 2 hr, red). The polysome region was defined as 5-somes and higher. P:M (polysome to monosome) ratio is 0.79 in control, 0.18 in HS8M, and 0.08 in HS2S. Profile is representative of three replicate experiments, where relative P:M ratios in HS8M were 22.5% ± 0.9% of control values, and relative P:M ratios in HS2S were 7.2% ± 2.7% of control.
(B) Normalized footprint density along mRNAs (see Experimental Procedures).
(C) Raw footprint count per position (smoothed) across the Serpine1 transcript (ENSMUST00000041388, 5' UTR and first 500 nt of the transcript are shown).
(D) Distribution of r5' LR values in HS8M (green) and HS2S (red). Population means are indicated.
(E) Changes in 5' LR values after heat shock.
See also Figure S1 and Table S1.
consistent with previous observations and with a global reduction in translation (McCormick and Penman, 1969). Notably, lighter polysome fractions representing two to four ribosomes actually increased somewhat in severe heat stress (Figure 1A), discussed further below.

The distribution of ribosome footprint sequence reads along individual mRNAs, or in a “metagene” analysis of reads mapping to all mRNAs, revealed a more complex picture (Figures 1B and 1C). While changes in response to 8 hr of mild heat shock were generally modest, dramatic changes in relative ribosome occupancy occurred in response to 2 hr of severe heat shock in four distinct regions. These included increases in the 5’ untranslated region (UTR) and the “initiation region” (reads indicating positioning of the ribosomal P site in the first 15 bases of the open reading frame [ORF]), and an ~1.7-fold increase in the next ~180 nucleotides (nt) after the initiation region, which shifted to a ~1.7-fold decrease in the remainder of the ORF (Figures 1B and S1J). The overall shape of this distribution remained unchanged whether gene-level normalization was used (as in Figures 1B, S1J, and S1K) or simply plotting raw read counts (Figure S1L). Many individual genes that had sufficiently high read coverage displayed a distribution similar to the metagene distribution, as illustrated for Serpine1 (Figure 1C; additional examples are shown in Figure S1M). Similar biases in relative ribosome occupancy after severe heat shock were observed in a biological replicate experiment described below. These wholesale changes in ribosome occupancy along mRNAs imply major shifts in translational regulation in response to heat shock. The accumulation of ribosomes in the ~200 nt after the initiation site relative to the remainder of the ORF was particularly notable because it implied extensive postinitiation regulation of translation. Such remainder of the ORF was particularly notable because it

**Translation Elongation Pausing in Severe Heat Stress**

Multiple regulatory mechanisms might produce such a global increase in 5’ LR, including (1) transient (reversible) pausing of translation elongation around codon 65 on most of the gene’s mRNAs (resulting in accumulation of ribosomes 5’ of the pause), (2) irreversible stalling of translation elongation around codon 65 on a subset of the gene’s mRNAs (also resulting in accumulation of ribosomes upstream), (3) acceleration of translation elongation after codon 65, or (4) premature termination of translation around codon 65 on a subset of a gene’s messages. To discriminate between these potential explanations for the increase in 5’ LR after severe heat shock, we performed analytical polysome profiling after inhibition of translation initiation with the drug Harringtonine, which blocks initiation after subunit joining by preventing elongation during the first round of peptide bond formation (Huang, 1975; Ingolia et al., 2011). Since treatment with Harringtonine inhibits newly initiating ribosomes while allowing previously engaged ribosomes to continue translating through termination, this drug enables the fates of elongating ribosomes to be followed under different conditions.

As expected, under control conditions, treatment with Harringtonine resulted in a decrease in heavy polysomes and an increase in monosomes that was readily apparent after 1 min and more complete after 3 min (Figure 2A). Under severe heat shock conditions, 1 min of Harringtonine treatment had only a minimal effect on the polysome profile, but 3 min treatment resulted in a distinct reduction in heavy polysomes, suggesting that most ribosomes are not irreversibly stalled (Figures 2A and S2A). An acceleration of translation after codon 65 would predict a more rapid rather than slower collapse of polysomes in heat shock conditions, and premature termination would also not be expected to slow the collapse of polysomes. The slower kinetics of this collapse are therefore most consistent with the model in which ribosomes are transiently paused during translation elongation in heat shock conditions.

To confirm that the reduced collapse rate of polysomes under heat stress after Harringtonine treatment was due to altered elongation rates, we used the translational inhibitor puromycin, which causes dissociation of ribosomes that are actively translating. Treatment with puromycin yielded a similar reduction in polysomes in heat-stressed cells as under control conditions (Figure S2B), indicating that under severe heat stress, as in control conditions, most ribosomes are translating and are neither irreversibly stalled nor prematurely terminating. Together, these data support that the slowed kinetics of polysome collapse result from altered elongation rates.

Ribosome pausing was centered around codon 65 based on the metagene analysis, but such an averaged representation could hide variability between genes. To characterize this phenomenon in more depth, we calculated the inferred location of ribosome pausing on a subset of the gene’s mRNAs (also resulting in accumulation of ribosomes upstream), (3) acceleration of translation elongation after codon 65, or (4) premature termination of translation around codon 65 on a subset of a gene’s messages. To discriminate between these potential explanations for the increase in 5’ LR after severe heat shock, we performed analytical polysome profiling after inhibition of translation initiation with the drug Harringtonine, which blocks initiation after subunit joining by preventing elongation during the first round of peptide bond formation (Huang, 1975; Ingolia et al., 2011). Since treatment with Harringtonine inhibits newly initiating ribosomes while allowing previously engaged ribosomes to continue translating through termination, this drug enables the fates of elongating ribosomes to be followed under different conditions.

To confirm that the reduced collapse rate of polysomes under heat stress after Harringtonine treatment was due to altered elongation rates, we used the translational inhibitor puromycin, which causes dissociation of ribosomes that are actively translating. Treatment with puromycin yielded a similar reduction in polysomes in heat-stressed cells as under control conditions (Figure S2B), indicating that under severe heat stress, as in control conditions, most ribosomes are translating and are neither irreversibly stalled nor prematurely terminating. Together, these data support that the slowed kinetics of polysome collapse result from altered elongation rates.

Ribosome pausing was centered around codon 65 based on the metagene analysis, but such an averaged representation could hide variability between genes. To characterize this phenomenon in more depth, we calculated the inferred location and statistical significance of shifts in ribosome footprint density along individual genes using a modified Kolmogorov-Smirnov (KS) test. We reasoned that if translation of an mRNA is paused in heat shocked cells, ribosomes should accumulate upstream of the pause, so the density of footprint reads should be significantly higher between the AUG and the location of the pause relative to the remainder of the ORF; this can be determined by
The KS test (Supplemental Experimental Procedures). For example, this test identified a highly significant shift in footprint density at nucleotide 189 (codon 63) of the Vimentin ORF, a position we call the "KS location," suggesting that ribosomes pause in this vicinity and accumulate upstream (Figure 2B).

Genome-wide KS analysis indicated that 75% of the translated mRNAs had a significant elongation pause under severe heat shock conditions at a false discovery rate (FDR) cutoff of 5%. The location detected by the KS analysis beyond which footprint density began to plateau mostly occurred between...
Translation Elongation Pausing in Heat Shock

Translation elongation pausing is induced during heat shock. This phenomenon is observed in mouse mRNAs with a median estimated position of 263 nt. Thus, individual gene analysis indicated an elongation pause in the first few hundred bases for the majority of mouse mRNAs in response to severe heat shock, a phenomenon we call “5' ribosome pausing.” A ribosome paused in the first couple of hundred bases of an ORF might be expected to cause accumulation of a few ribosomes upstream, likely running as a light polysome, consistent with the observed increase in the light polysome fraction seen in severe heat shock (Figure 1A).

While our analyses suggested that the vast majority of genes experience 5' ribosome pausing in severe heat shock, the increase in 5' LR was not universal. One gene set of interest was the set of transcripts that were not significantly changed (based on the KS test). Functional enrichment analysis of this set detected enrichment for transcriptional regulators (GO term “transcription factor activity,” p = 3.5 × 10⁻⁶, FDR < 1%), which might contribute to the transcriptional response to heat shock. We used a more stringent set of criteria to define “escapers” as mRNAs whose 5' LR did not significantly change (or decreased) after severe heat shock (Supplemental Experimental Procedures). We did not observe HSPs to be among the escapers, suggesting that the upregulation of translation of HSPs noted above (Figures S1G–S1I) has other causes, e.g., increased mRNA levels and/or increased translation initiation. However, these criteria did identify 187 mRNAs that escaped translational pausing, including several transcription factors (Table S2). Interestingly, the Atf4 and Atf5 transcription factor genes were both among the escapers, with r5 LR values of 1.18 and 1.05, respectively, far below the mean value of around 3 (Figures 2D and S2D). ATF5 is translationally induced after a variety of stresses, including heat shock (Zhou et al., 2008). The homologous ATF4 factor is a major regulator of oxidative, endoplasmic reticulum (ER), and amino acid starvation stress (Harding et al., 2000) but has not been previously connected to the heat shock response. Both of these factors are known to be transcriptionally upregulated under stress conditions via a mechanism involving translation of upstream ORFs (uORFs) (Harding et al., 2000; Zhou et al., 2008). Our observations suggest that escape from the global elongation and initiation blocks are used to regulate distinct subsets of genes.

**Ribosome Association of Hsp70 Chaperones Is Altered during Heat Shock**

We next sought to explore potential mechanisms underlying the general phenomenon of 5' ribosome pausing. The near-universal scope and relatively specific location of the pause suggested the involvement of a general aspect of translation elongation, which might be dependent more on the nascent peptide than on genespecific properties. Nascent peptides are bound by a variety of factors such as SRP (discussed below) and Hsp70 chaperones (Kramer et al., 2009), whose regulation is also a hallmark of the heat shock response. The cytosolic Hsp70 (70 kilodalton [kDa] heat shock protein) chaperones HSC70 and HSP70—the constitutive and inducible forms of cytosolic Hsp70, respectively—are among the major chaperones in the cell that mediate protein folding. During translation, these proteins associate with ribosomes and interact with nascent chains as they emerge from the ribosome exit tunnel (Beckmann et al., 1990; Nelson et al., 1992). This function of Hsp70 family proteins has been shown to be important for cell growth both in yeast and in mammals (Jaiswal et al., 2011; Nelson et al., 1992). Heat stress places a substantial burden on chaperones as a result of the misfolding of existing cytosolic proteins, and, in response, cells upregulate stress-specific homologs of these chaperones, including HSP70 (encoded by the Hspa1a and Hspa1b genes in mice), which supplements the levels of the constitutive HSC70, encoded by Hspa8.

To assess the association of Hsp70 chaperones with the ribosome after heat shock, we fractionated polysome gradients from control or heat-shocked cells and measured the amount of HSC70/HSP70 that comigrated with polysomes using a pan-Hsp70 antibody. We detected a consistent 2- to 4-fold reduction in the association of HSC70/HSP70 with monosomes and polysomes in severe heat shock (Figures 3A and 3B). Relatively little comigration of HSC70/HSP70 was observed in the polysomal portion of the gradient following dissociation of ribosomes by EDTA treatment, supporting the specificity of the detected association (Figure S3A). Significant depletion of HSC70/HSP70 was detected in several fractions ranging from monosomes to 5-somes (Figure 3B). mRNAs with ribosomes paused at typical locations (around 200 nt) are expected to migrate more as lighter polysomes or monosomes, depending on whether additional ribosomes accumulate upstream of the pause. Reduced association of Hsp70 chaperones with translating ribosomes could affect the ribosome in various ways, including the exposure of nascent chains.

**Significantly Paused mRNAs Have Features Associated with Hsp70 Interaction**

HSC70/HSP70 bind nascent chains generally but have higher affinity for hydrophobic peptides, and hydrophobic peptides are expected to be more reliant on interaction with HSC70/HSP70 for proper folding (Hartl et al., 2011). The first 20–25 residues of proteins tend to be more hydrophobic than the rest of the protein, particularly for proteins containing a signal peptide (von Heijne, 1981), but also to a lesser extent for other proteins (Figures 3C and S3C). We found that genes with significant 5' ribosome pausing encoded more hydrophobic N termini than did nonsignificantly paused genes (Figure 3C, t test p value = 3.8 × 10⁻⁷). Hydrophobicity of the N terminus also differed between mRNAs grouped by their KS location. Those with KS locations between codons 44–171 were associated with higher hydrophobicity of the N terminus than the (smaller) sets of messages with more distally or more proximally located pause sites (Figure S3E), suggesting that amino terminal hydrophobicity contributes to pausing in this region. Hsp70 is known to have a particularly high affinity for stretches of five hydrophobic residues flanked by four positively charged amino acids on each side (Rüdiger et al., 1997). Analysis of the binding site content of protein N termini with the scoring system proposed by Rudiger and colleagues revealed that significantly paused mRNAs had better average Hsp70 binding site scores, reflecting greater propensity for Hsp70 binding (Figure 3D), suggesting that these peptides would be more dependent on Hsp70 binding. Consistently, the N termini of nonsignificantly paused mRNAs encoded...
Figure 3. Association of HSC70/HSP70 with Polysomes Is Altered after Heat Shock

(A) Western blot of HSC70/HSP70 (with pan-HSP70 antibody; see the Experimental Procedures for details) protein levels (middle panel) across a polysome gradient in control and severe heat shock cells (HS2S, top panel). RPL10A was used to assess ribosome abundance in each fraction (middle panel). Levels of
Translation Elongation Pausing in Heat Shock

Molecular Cell

Modulation of Hsp70 Chaperone Activity Regulates Translation Elongation Pausing

To test the potential role of Hsp70 chaperones in heat shock-induced elongation pausing, we first analyzed thermotolerance. Chaperone expression is induced by mild heat shock (Kelley and Schlesinger, 1978; Lindquist, 1980, 1981) (Figure S3B), explaining why mild heat shock treatment prior to severe shock has protective effects on cells in the phenomenon known as thermotolerance (Gerner and Schneider, 1975; Henle et al., 1978). Thermotolerance is dependent on protein synthesis and enables enhanced translation under severe heat shock conditions (Lindquist, 1980; Petersen and Mitchell, 1981). To assess how pretreatment of chaperones influences ribosome density patterns, we performed a thermotolerance experiment, pretreating cells with 8 hr of mild heat shock prior to 2 hr of severe heat shock, along with a biological replicate of the control, mild, and severe treatments. Pretreatment of cells with mild heat shock resulted in partial rescue of the polysome collapse that occurred during severe heat shock (Figure 4A), consistent with the expected relief from translational inhibition. Replicate ribosome footprint data yielded metagene profiles under control, mild, and severe heat shock conditions (Figure 4B) similar to those observed in Figure 1. Remarkably, pretreatment with mild heat shock almost completely rescued the 5′ ribosome pausing observed after 2 hr of severe heat shock (Figures 4B and S4A). These observations suggest that 5′ ribosomal pausing is regulatable and is reduced by factors induced by mild heat stress such as chaperones.

To more specifically look at the role of HSC70/HSP70 chaperones in the phenomenon, we explored whether inhibition of Hsp70 activity under control conditions would affect translation elongation. To test this possibility, we subjected cells to a short treatment of a small-molecule drug, VER-155008, that inhibits the ATPase activity of both constitutive and inducible Hsp70 proteins, an activity that is required for their chaperone function (Massey et al., 2010). Treatment with the Hsp70 inhibitor for a short period led to a reproducible reduction in heavy polysomes and an accumulation of monosomes (Figure 4C), supporting the direct involvement of Hsp70 proteins in the control of translation in mammalian systems. Footprint profiling revealed a pronounced accumulation of ribosomes near the 5′ ends of mRNAs in cells treated with the drug for 3 hr in the absence of stress (Figure 4D).

Analysis of individual mRNAs revealed that ~2,500 mRNAs were significantly paused after treatment with Hsp70 inhibitor, the majority of which were also significantly paused under severe heat shock (overlap of 2072, p = 10\(^{-34}\)) with a similar but somewhat broader distribution of inferred KS locations (Figure S4C). Similar to what we observed under severe heat shock (Figure 3C), mRNAs that were significantly paused after Hsp70 inhibitor treatment encoded significantly more hydrophobic N termini than nonpaused mRNAs (p = 5 \times 10^{-7}, Figure S4F). The inferred KS locations were in the vicinity of those seen under severe heat shock for 60% of the mRNAs, comparing the same mRNAs in both experiments (Figures S4D and S4E), while the metagene analysis showed accumulation somewhat 5′ to that observed in heat shock. Differences between the effects of inhibitor treatment and severe heat shock on footprint profiles may reflect incomplete inhibition of Hsp70 activity or functions of Hsp70s that do not require ATP hydrolysis. Similar to the trends observed in severe heat stress, more proximal KS locations correlated with greater N-terminal hydrophobicity, while more distal KS locations were associated with reduced N-terminal hydrophobicity (Figure S4G).

To verify that Hsp70 chaperones directly affect ribosome accumulation and to explore this phenomenon in another species, we overexpressed the inducible Hsp70 (Hspa1a) in human 293T cells. We first subjected 293T cells to 2 hr of severe heat stress. After severe heat shock, 293T cells showed a similar 5′ ribosome accumulation to that observed in mouse 3T3 cells, with a similar pause location at around 200 nt into the ORF based on metagene analysis (Figure 4E), of somewhat smaller
This observation extends the phenomenon of heat stress-induced ribosomal accumulation to a second mammalian species.

Ectopic expression of Hsp70s is generally inhibitory to cell growth (Feder et al., 1992), but we were able to achieve moderate levels of overexpression of HSP70, about 1.5-fold in...
under control conditions (Figure S4H). HSP70-overexpressing in transformed lines, 293T cells express both HSC70 and HSP70 conditions (Figure S4H). We note that, as is commonly observed excess over HSC70/HSP70 levels observed in normal growth conditions (Figure S4H). We note that, as is commonly observed in transformed lines, 293T cells express both HSC70 and HSP70 under control conditions (Figure S4H). HSP70-overexpressing cells were then subjected to 2 hr of severe heat shock. Strikingly, this moderate ectopic expression of Hsp70 resulted in a partial rescue of the elongation pausing following severe heat shock, as seen in Figure 4E (see also Figures S4I and S4J). The observations that Hsp70 inhibitor treatment leads to 5’ ribosome pausing in the absence of stress, and that moderate overexpression of Hsp70 partially relieves heat shock induced elongation pausing, directly implicate Hsp70 chaperone in the regulation of translation elongation during severe heat shock.

The Hsp70-Translational Machinery Interactome: Characterization and Stress-Associated Changes

Hsp70 chaperones are known to bind numerous proteins (Calloni et al., 2012; Kampinga and Craig, 2010). Here, it was of particular interest to characterize interactions between Hsp70s and the translation machinery. Consistent with a recent observation (Jaiswal et al., 2011) that the ribosomal cochaperone DNAJC2 (MPP11) does not influence the ribosomal association of HSC70 (the predominant Hsp70 protein in 3T3 cells in both control conditions and after 2 hr of severe heat shock), knockdown of this factor in 3T3 cells did not appreciably affect the distribution of ribosomes under normal growth conditions (data not shown).

To explore potential ways in which Hsp70 chaperones may influence elongation pausing, we next sought to characterize direct interactions between Hsp70s and the translation machinery using the high-throughput LUMIER with BACON assay (LUMIER with Bait control, Experimental Procedures) (Taipale et al., 2012). Using this assay, we examined the interaction of 504 clones representing a broad panel of 377 translation-related genes (Table S3) in a 293T cell line stably expressing Renilla-tagged HSC70. We found that HSC70 interacts with about half of these factors to varying extents (213 genes, above the 75th percentile of the estimated background luminescence, Figure S5A).
and with 109 of them strongly (at the 99.9th percentile of background, Figures S5A and S5B). The known Hsp70 client proteins HSF1 (Abravaya et al., 1992) and p53 (Zylicz et al., 2001) were among the top interactors (ranked 4 and 33 in their luminescence score), as were a handful of cochaperones which were included as positive controls. We next asked whether these interactions changed in response to stress. The LUMIER assay is based on Renilla luciferase tagging, and Renilla luciferase activity is markedly reduced after heat shock (data not shown). Thus, for application of this method, we instead treated cells with another proteotoxic stress, the proteasome inhibitor bortezomib. Of course, some aspects of the cellular responses to bortezomib and heat stress may differ. But the resulting interaction data showed a global reduction in the interaction of HSC70 with ribosomal proteins (Figure S5C), paralleling the reduced ribosomal association of Hsp70 observed in severe heat shock.

To explore potential changes in the interactions of the endogenous Hsp70 chaperones with the translational machinery following heat stress, we used a lower-throughput communo- precipitation (coIP) assay. This assay was applied after 2 hr of severe heat shock to each of a set of 48 translation-related proteins, including primarily translation factors and ribosomal proteins (and several positive controls) representing a spectrum of basal HSC70 interaction scores (Figures 5B, 5C, and S5D and Table S4). In this assay, transfected FLAG tagged proteins were pulled down and protein expression quantified by anti-FLAG antibody, while interaction with the endogenous Hsp70 was measured with a pan-Hsp70 antibody. Because the FLAG tags were located at the C terminus, this assay measures interaction with mature rather than nascent proteins. Interaction scores were then calculated as the ratio of [Hsp70]/[FLAG], controlling for effects on expression. HSF1, the heat shock transcription factor, had reduced interaction after heat stress (Figures 5B and 5C), consistent with its phosphorylation, trimerization, and association with the translational machinery (Björk and Sistonen, 2010)

Heat shock is known to cause misfolding of pre-existing proteins, which are then detected by Hsp70 and other chaperones for refolding. If the chaperones are unable to refold those proteins, they are targeted for degradation, thus preventing accumulation and aggregation of misfolded proteins in the cell (Hartl et al., 2011). We classified each of the tested proteins according to the change in its Hsp70 interaction score (increased or decreased), and protein levels (stable or destabilized, as measured by FLAG), after heat shock. Several proteins showed a large reduction in their overall protein levels while their interaction with Hsp70 chaperones remained unchanged or was elevated, resulting in an increase in their interaction scores (Figure 5B). This group is expected to consist of proteins that become misfolded and degraded as a result of severe heat shock, and hence show an increase in Hsp70 interaction. Among those are the release factor GSPT2 (eRF3 component), and the initiation regulator RPS6KB2.

Only two proteins showed a consistent upregulation in their interaction with Hsp70 without a respective decrease in expression: RPL22 and RPL4 (Figure 5B). Interestingly, both these proteins protrude into the interior of the ribosome exit tunnel and have been shown to play a role in gene-specific elongation stalling in bacteria and eukaryotes (Wilson and Doudna Gate, 2012).

The largest class of changes included proteins that were relatively stable and showed reduced Hsp70 interactions (Figure 5B). We expect this group to reflect regulatory interactions. Hsp70 was found to interact with five elongation factors under normal growth conditions (Figure 5A), out of which EEF1A1 was found to have reduced Hsp70 interactions during heat stress. Several ribosomal proteins also showed this type of reduced interaction with Hsp70 after heat shock, further supporting the reduced association of Hsp70 with ribosomes during heat stress. In particular, RPL23A, which is located on the outside of the exit tunnel, was downregulated in its interaction with Hsp70 after severe heat shock (Figures 5B and 5C). This supports the notion that Hsp70 chaperones are present in proximity to the exit tunnel under normal growth conditions, and their association is reduced under severe heat shock.

DISCUSSION

In this study we identify an unanticipated aspect of the cellular response to heat stress, in which ribosomes accumulate at the 5’ ends of open reading frames of most mRNAs, apparently as a result of temporary pausing of translation elongation (Figure 6). Under heat stress, cellular priorities shift from growth to cytoprotective functions, including prevention of protein misfolding and aggregation. Reduced HSC70/HSP70 association with ribosomes and altered interactions with the translational machinery were observed after 2 hr of severe heat stress. These changes may serve several functions, including reversible inhibition of translation elongation that, together with inhibition of translation initiation, may help to reduce cellular protein production and the associated burden on the cellular chaperone machinery. Reversibility of the elongation pause by induction of chaperones would allow cells to rapidly accelerate protein synthesis and growth once they have effectively adapted to the stress associated with protein misfolding, or after a return to nonstress conditions.

We observed elongation pausing in heat stress in both mouse and human cells. These findings, together with a much earlier report pointing to changes in elongation rates with heat shock in Drosophila cells (Ballinger and Pardue, 1983), suggest that the mechanism may be a deeply conserved feature of the response to proteotoxic stress. Our data show that the levels and activity of Hsp70s impact elongation pausing. A variety of possible mechanisms for these effects can be imagined, including altered interactions of Hsp70 with elongation factors, Hsp70-dependent regulation of ribosomal proteins in the exit tunnel, or aberrant exposure of nascent peptides in heat stress. Multiple chaperones are known to interact with ribosomes and with nascent peptides in both prokaryotes and eukaryotes (Kramer et al., 2009), and it is possible that additional ribosome-associated chaperones play a role in translation elongation under normal or stress conditions.

Analyzing the Hsp70-translational machinery interactome, we found that Hsp70 interacted with five elongation factors under normal growth conditions, and that its interaction with EEF1A1 was reduced after severe heat stress. This raises the possibility that Hsp70’s altered interaction with EEF1A1 in heat stress might...
Molecular Cell
Translation Elongation Pausing in Heat Shock

Figure 6. Heat Shock-Induced Translation Elongation Pausing Is Modulated by Hsp70 Chaperones

During severe heat shock, ribosomes often pause at 5’ ends of mRNAs after translating about 65 amino acids (Figures 1 and 2), resulting in accumulation of ribosomes upstream and depletion of ribosomes downstream of this point; after severe heat shock, HSC70/HSP70 association with the ribosome is reduced (Figures 3A and 3B). As Hsp70 proteins interact with nascent peptide chains as they emerge from the exit tunnel (Beckmann et al., 1990; Nelson et al., 1992), nascent chains might be exposed. More significantly paused mRNAs tend to encode N termini with stronger Hsp70 requiring characteristic (Figures 3C and 3D). Modulation of Hsp70 levels, by thermotolerance or overexpression, rescues heat shock-induced elongation pausing, while inhibition of Hsp70 activity leads to pausing in the absence of stress (Figure 4). Hsp70 chaperones interact with various translation factors and ribosomal proteins, and this interaction is regulated after heat shock (Figure 5). HSC70/HSP70 may affect ribosome dynamics via its interaction with elongation factors, regulation of ribosomal proteins in the exit tunnel, or through exposure of nascent chains.

- Ribosomes pause at ~200 nt in most ORFs
- Reduced association of Hsp70 chaperones with ribosomes
- Elongation pausing associated with hydrophobic N-termini
- Modulation of Hsp70 levels/activity affects elongation pausing

influence elongation rates, though whether and how this could contribute to elongation pausing is unclear. A handful of initiation factors also had decreased interactions with Hsp70 under heat stress, suggesting a possible connection between Hsp70 and regulation of translation initiation under heat shock beyond its previously reported role. However, any link to initiation factors is likely distinct from Hsp70’s role in elongation pausing.

Our analysis identified RPL4 and RPL22, the only two ribosomal proteins that extend into the wall of the ribosome exit tunnel, forming a “constriction” (Wilson and Doudna Cate, 2012), as the only two stable proteins in the Hsp70-translational machinery interactome that showed a consistent increase in their overall extent of Hsp70 interaction in heat shock. The exit tunnel itself was long considered a passive region in the ribosome. More recently, evidence has emerged that structural changes involving RPL22 and RPL4 can occur inside the exit tunnel, leading to regulatory effects on elongation (Berisio et al., 2003; Wilson and Doudna Cate, 2012). For example, the bacterial L22 protein interacts with the nascent peptide of SecM (Nakatogawa and Ito, 2002) and the eukaryotic RPL4 protein interacts with the nascent peptide of a uORF of the CMV protein gp48 (Bhushan et al., 2010), both leading to greater constriction of the exit tunnel and stalling of elongation. One possibility is that during heat shock Hsp70 chaperones regulate the conformation of the exit tunnel via interactions with RPL22 and/or RPL4.

The overall association of HSC70/HSP70 with ribosomes was downregulated after 2 hr of severe heat stress (Figures 3A and 5C). A number of ribosomal proteins have previously been shown to efficiently incorporate into ribosomes when C-terminally tagged, e.g., RPL22 (Sanz et al., 2009), RPL23A (Inada et al., 2002; Ross et al., 2007), RPL18, and RPL16 (Halbeisen et al., 2009), suggesting that our assay often detected interaction with proteins incorporated into ribosomes. The extent of pausing correlated with hydrophobicity of the N termini and with presence of Hsp70 binding motifs. Hsp70 proteins have evolved to recognize exposed hydrophobic patches, particularly when flanked by basic residues, as a sign of a misfolded or aggregated protein. In the absence of ribosome-associated Hsp70, nascent peptides with stronger Hsp70 binding motifs might be more dependent on Hsp70 and thus have a greater tendency to misfold or aggregate, potentially impacting the efficiency of translation. More distant KS locations were associated with reduced N-terminal hydrophobicity (Figures 3C, 3D, and S3E), consistent with ribosomes being able to translate further downstream before pausing. These observations suggest that exposure of nascent peptides might impact elongation. Finally, RPL23A (Rpl25 in yeast), which serves as the docking site for nascent chain binding accessory factors (Kramer et al., 2009), interacted strongly with Hsp70 under normal growth conditions, and showed reduced interactions under heat stress (Figures 5B and 5C). This observation suggests that Hsp70 is present in the vicinity of the exit tunnel under normal growth conditions, but much less so during heat stress. Indeed Hsp70 has been proposed in the past to play a role in aiding in the passage of nascent peptides through the exit tunnel (Nelson et al., 1992).
Together, the above observations suggest that exposed nascent peptides emerging from the ribosome might be involved in the process of elongation pausing (Figure 6).

The prevalence of elongation pausing in the heat stress responses of both mouse and human cells make it an intriguing phenomenon to explore in other stresses and organisms. Elongation pausing could facilitate translational repression under certain conditions, such as proteotoxic stresses, while induction of chaperones might counteract stress-induced elongation pausing. For example, elevated levels of chaperones are often observed in cancer (Mossor and Morimoto, 2004; Whitesell and Lindquist, 2005). This phenomenon may therefore be relevant to a broad spectrum of stresses and diseases.

**EXPERIMENTAL PROCEDURES**

**Heat Shock and Drug Treatment Conditions**
Mouse fibroblast 3T3 cells were plated at low density and then treated with mild (8 hr at 42°C) or severe heat shock (2 hr at 44°C) to induce chronic or acute heat stress responses, respectively (Supplemental Experimental Procedures). Harringtonine treatments were performed at 0.1 mM harringtonine for 3 min, or 1 min of 0.1 mM harringtonine followed by 2 min of 0.1 mg/ml cycloheximide (CHX), which were added to the media. Puromycin treatment was done at 1 mg/ml for 3 min. Treatment with the Hsp70 inhibitor VER-155008 (Massey et al., 2010) (Tocris Bioscience) was performed at 20 μM for 3 hr.

**Ribosome Footprint Profiling**
Ribosome footprinting was performed as described by (Ingolia et al., 2009) with modifications described in the Supplemental Experimental Procedures.

**Mapping of Footprint Reads**
Ribosome footprint reads were trimmed from the 3’ end from 36 bases to 32 bases, and multiple trailing adenosine (A) bases were removed. Next, footprint reads of size 22–32 bases were mapped to the mouse genome (mm9) or the human genome (hg18) with Bowtie (Langmead et al., 2009). Reads mapping uniquely to exonic positions were subsequently mapped to Ensembl gene and transcript annotation files (downloaded from UCSC) using custom python and Perl scripts (Supplemental Experimental Procedures).

**Polysome Profiling**
3T3 cells were treated with 0.1 mg/ml cycloheximide (CHX) for 5 min, then lysed (Supplemental Experimental Procedures). Profiles were normalized by loading of equal amounts of RNA OD units of all samples in a single experiment. To compare between experiments, we used relative P.M ratio, calculated as the P.M ratio in the given treatment divided by that in the corresponding control condition and expressed as a percentage.

**Protein Association with Polysome Fractions**
After ultracentrifugation, 24 equal volume fractions were collected and pooled as described in Figure 3A, top panel, and precipitated by addition of ice cold acetone and overnight incubation at –20°C. Significance of the reduction in ribosomal association of HSC70/HSP70 was calculated by t test over five replicates using MATLAB. Further details provided in the Supplemental Experimental Procedures.

**Antibodies**
Antibodies are detailed in the Supplemental Experimental Procedures.

**Footprint Density Plots and Tests of Pause Significance and Location**
All uniquely mapping exonic footprint reads were mapped to Ensembl mRNA transcripts with their genomic annotations in the ensGene table from the UCSC database. For each transcript, a profile was generated as a vector containing the number of reads for which the 5’ end mapped 12 nt 5’ of each position of the transcript, and then normalized by dividing by the mean number of reads per position along the first 450 nucleotides of the CDS, similarly to Ingolia et al. (2009). Presented in the figures are the averaged profiles for the filtered unique set of transcripts in each condition (see the Supplemental Experimental Procedures for further details on this and on KS analysis).

An elevation of ribosome density at the 5’ end of yeast genes, which was gradually decreasing up to ~600 bases of the ORF, has been observed in yeast under normal growth conditions in the first footprinting study (Ingolia et al., 2009). This elevation has been attributed to effects of transfer RNA abundance on translational efficiency (Qian et al., 2012; Tullier et al., 2010).

**Hydrophobicity Profiles and Hsp70 Binding Site Scoring**
Hydrophobicity profiles and Hsp70 binding site scoring are described in the Supplemental Experimental Procedures.

**Hsp70 Translation Machinery Interactome**
LUMIER with BACON assay was done as in Taipale et al. (2012), with a 293T cell line stably expressing Renilla-tagged HSC70 (Hspa8). CoIP with endogenous Hsp70 chaperones was performed as in Taipale et al. (2012). Further details are provided in the Supplemental Experimental Procedures.

**ACKNOWLEDGMENTS**
We thank Wendy Gilbert and members of her lab for helpful advice and suggestions, use of equipment throughout this study, and for critical reading of the manuscript. We thank Eric Wang for help with mapping of footprint data and for useful discussions, Sandro Santagata and Luke Whitesell for helpful advice, and Daniel Treacy for help with footprint library preparation. R.S. is an Awardee of the Weizmann Institute of Science-National Postdoctoral Award Program for Advancing Women in Science. This work was supported by an EMBO long-term fellowship and the Machiah foundation (R.S.), NIGMS fellowship number F32GM095060 (J.A.H.), and grants from the NIH (C.B.B.).

Received: March 27, 2012
Revised: September 27, 2012
Accepted: November 30, 2012
Published: January 3, 2013

**REFERENCES**
Molecular Cell

Translation Elongation Pausing in Heat Shock


