Identification of a Tissue-Selective Heat Shock Response Regulatory Network

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Abstract

The heat shock response (HSR) is essential to survive acute proteotoxic stress and has been studied extensively in unicellular organisms and tissue culture cells, but to a lesser extent in intact metazoan animals. To identify the regulatory pathways that control the HSR in Caenorhabditis elegans, we performed a genome-wide RNAi screen and identified 59 genes corresponding to 7 positive activators required for the HSR and 52 negative regulators whose knockdown leads to constitutive activation of the HSR. These modifiers function in specific steps of gene expression, protein synthesis, protein folding, trafficking, and protein clearance, and comprise the metazoan heat shock regulatory network (HSN). Whereas the positive regulators function in all tissues of C. elegans, nearly all of the negative regulators exhibited tissue-selective effects. Knockdown of the subunits of the proteasome strongly induces HS reporter expression only in the intestine and spermatheca but not in muscle cells, while knockdown of subunits of the TRiC/CCT chaperonin induces HS reporter expression only in muscle cells. Yet, both the proteasome and TRiC/CCT chaperonin are ubiquitously expressed and are required for clearance and folding in all tissues. We propose that the HSN identifies a key subset of the proteostasis machinery that regulates the HSR according to the unique functional requirements of each tissue.

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Introduction

The heat shock response (HSR) has been studied extensively as a cellular response to acute stress such as elevated temperature [1]. The master regulator of the HSR is Heat Shock Factor 1 (HSF1), a stress responsive transcription factor that regulates the inducible transcription of a family of genes encoding heat shock proteins (HSPs), many of which are molecular chaperones. In the absence of a stress signal, HSF1 is inhibited by a negative feedback loop mediated by the molecular chaperones HSP70 and HSP90 [2–7]. Upon heat shock, HSF1 is activated as the equilibrium of chaperones shifts toward association with metastable polypeptides.

Many key aspects of the HSR have been well established at a cellular level in cultured cells and unicellular organisms, yet the HSR has additional features that are only apparent in multicellular organisms. Heat shock inducible promoters contain multiple cis elements and can be differentially expressed across tissues [8–18]. The HSR is intimately associated with numerous tissue-specific and age-dependent human diseases and regulated cell non-autonomously by neuronal control [19,20]. Finally, HSF1 has important roles during development and longevity, and activation of the HSR is attenuated during aging [13,21–24]. However, despite the importance of the HSR in organismal physiology, relatively little is known about its regulation in multicellular organisms and the extent of differential regulation across distinct tissues is unexplored.

A comprehensive genetic analysis of the HSR regulatory pathways has not previously been possible in any system, in part because traditional forward genetic screens are inadequately suited to the identification of genes that regulate the HSR. These approaches depend on the introduction of mutations, which can destabilize the folding of the corresponding proteins, resulting in indirect induction of the HSR due to the expression of misfolded species. Indeed, a forward genetic screen in Drosophila described such mutations in a muscle-specific actin [25,26]. RNAi based genetic screening resolves the limitations associated with traditional genetic screens associated with the HSR and has been used to gain important insights into many regulatory networks including those associated with models of aggregation-prone proteins, longevity, and stress responses [27–34].

In this study, we have used genome-wide RNAi screening to identify factors important for the positive and negative regulation of the HSR in the metazoan Caenorhabditis elegans in order to establish a comprehensive understanding of its regulation on an organismal level. Further, we used a fluorescent reporter to allow for the analysis of regulation in different tissues. This approach
Author Summary

The heat shock response (HSR) is an essential stress response that functions to maintain protein folding homeostasis, or proteostasis, and whose critical role in human diseases is recently becoming apparent. Previously, most of our understanding of the HSR has come from cultured cells and unicellular organisms. Here we present the identification of the heat shock regulatory network (HSN) in Caenorhabditis elegans, an intact, multicellular organism, using genome-wide RNAi screening. We identify 59 positive and negative regulators of the HSR, all of which have a previously established role in proteostasis, linking the function of the HSR to its regulation. Some HSN genes were previously established in other systems, many were indirectly linked to HSR, and others are novel. Unexpectedly, almost all negative regulators of the HSR act in distinct, tissue-selective patterns, despite their broad expression and universal cellular requirements. Therefore, our data indicate that the HSN consists of a specific subset of the proteostasis machinery that functions to link the proteostasis network to HSR regulation in a tissue-selective manner.

reveals a complex network of positive and negative HSR regulators with critical roles in maintenance of proteostasis that confer differential tissue-selective patterns of heat shock gene expression.

Results

Genome-Wide Screens for HSR Regulators

The genetic network upstream of HSF1 and the HSR was identified using a genome-wide RNAi screen in transgenic \textit{C. elegans} expressing the heat shock (HS)-inducible fluorescent reporter \textit{phsp70::gfp} constructed from the promoter of the \textit{C12C8.1} gene \cite{[13]}. Expression of this reporter is not detected under ambient growth conditions of development and adulthood (Figure 1A) and is induced strongly by HS (Figure 1B). The threshold sensitivity of the screen was established using RNAi knockdown of \textit{hsf-1} to suppress HS-induction of the reporter as a reference control for positive regulators (Figure 1C), and RNAi knockdown of \textit{hsp-1}, a member of the HSP70 family that negatively regulates the HSR, resulting in constitutive expression of the reporter as a reference control for negative regulators (Figure 1D).

Genetic modifiers of the HSR were identified by visual scoring of the \textit{phsp70::gfp} reporter upon RNAi-mediated knockdown. A representative subset from each functional class was validated by analysis of endogenous \textit{hsp70} gene expression using qRT-PCR (Figure 1E and 1F). We also extended our analysis to another heat shock gene, \textit{hsp-16.2}, a member of the small HSP family. Consistent with the HSR reporter results, HS-dependent induction of \textit{hsp70} and \textit{hsp-16.2} were reduced upon \textit{hsf-1} knockdown. Likewise, the basal expression of \textit{hsp70} and \textit{hsp-16.2} were increased upon \textit{hsp-1} knockdown. These experiments established the utility of the \textit{phsp70::gfp} reporter and RNAi as a methodology for the identification of both positive and negative regulators of the HSR in \textit{C. elegans}.

Having established the criteria for two classes of HSR genetic modifiers, we performed a genome-wide RNAi screen for genes whose knockdown blocked HS-dependent reporter induction, and for genes whose knockdown resulted in constitutive expression of the reporter. These screens were performed by RNAi feeding using a library containing RNAi constructs targeted against approximately 86% of genes in \textit{C. elegans} \cite{[35]}.

Identification of HSR Positive Regulators

The screen for positive regulators of the HSR identified genes with properties similar to \textit{hsf-1}, whose knockdown suppressed induction of the HSR reporter. To ensure that decreased fluorescence of the reporter did not arise from indirect effects, such as transgene silencing, we performed a counter-screen against suppression of a \textit{phsp-4::gfp} reporter, an ER stress-inducible gene that is not dependent on HSF1 \cite{[36,37]}. This led to the identification of seven positive regulators that are conserved to humans and function in chaperin remodeling, RNA processing, and protein synthesis (Figure 1E, Table 1, Table S1). None of these genes has been previously linked to HSR regulation, although each has been either associated with HS or implicated in the HSR. For example, \textit{dep-66} is a subunit of the NuRD complex, of which other subunits in this complex have been shown to interact with human HSF1 \cite{[38]}. Our data provide evidence that the HSF1-NuRD interaction has functional consequences on the regulation of the HSR. Likewise, \textit{Mi-2}, a subunit of several complexes including NuRD, has been shown to affect the levels of HS genes in \textit{Drosophila} \cite{[39]}. Among the other positive regulators are genes associated with mRNA splicing and translation, biosynthetic processes that are highly sensitive to HS stress. \textit{F09D1.1} is a homologue of USP39, which has been implicated in recycling of the triple-snRNP complex, a step of splicing that is particularly sensitive to temperature. \textit{phi-8} and \textit{phi-11} are subunits of Splicing Factor 3, which has been shown to regulate alternative splicing, \textit{snr-3} is an sm protein which is expected to have a general role in mRNA splicing, and \textit{ets-2} is an elongation factor 2-like protein predicted to have a general role in translational elongation. Finally, as expected, \textit{hsf-1} was identified in the screen as predicted for its central role in the HSR.

Identification of HSR Negative Regulators

The screen for negative regulators of the HSR identified genes whose reduced expression resulted in the constitutive expression of the \textit{phsp70::gfp} reporter. To ensure that these regulators activated the HSR in an HSF1-dependent manner, we employed a subsequent counter-screen using a hypomorphic \textit{hsf-1} mutant \cite{[40]}. Candidate negative regulators were also tested for their ability to constitutively activate endogenous heat shock genes by qRT-PCR (Figure 1F). This strategy led to the identification of fifty-two genes that have the functional properties of negative regulators of the HSR (Table 2, Table S1).

Each of these negative regulators of the HSR function in specific steps of proteostasis and affect either gene expression, protein folding, trafficking, and clearance, and are conserved to humans. Among the regulators that affect protein folding are three prominent molecular chaperone machines corresponding to \textit{HSP70} (\textit{hsp-1}), \textit{HSP90} (\textit{daf-21}) and TRiC/CCT (\textit{cct-1}, \textit{cct-2}, \textit{cct-3}, \textit{cct-4}, \textit{cct-5}, \textit{cct-6}, \textit{cct-7}, and \textit{cct-8}) and three chaperones (\textit{sqt-1}, \textit{unc-45}, and \textit{cym-11}). \textit{HSP70} and \textit{HSP90} are predicted from previous studies that identified them as negative regulators of HSF1 and the HSR. Likewise, a role for the TRiC/CCT chaperin in the regulation of the HSR has been suggested from studies on a small molecule that interacts with TRiC/CCT and induces human HSF1 \cite{[41]}. Regulation of the HSR by chaperonins is functionally conserved in bacteria, as downregulation of the prokaryotic chaperonin GroEL induces the HSR in \textit{E. coli} \cite{[41,42]}. The selectivity of these genes representing three chaperone machines and three chaperones as regulators of the HSN is unexpected given that \textit{C. elegans} expresses nearly 200
chaperone genes, which suggests a high degree of selectivity for chaperone regulation of the HSR.

Other negative regulators of the HSR correspond to components of trafficking including subunits of the signal recognition particle (SRP) and other secretory pathway genes (F55C5.8, F08D12.1, F38A1.8, R186.3, F38E11.5 and T14G10.5, hsp-3, T24H7.2, let-607) and mitochondrial import (hsp-6, T09B4.9). Consistent with this, knockdown of SRP subunits in yeast and E. coli has been shown to induce the HSR [43,44] and our study now extends these observations to metazoans. Clearance components include ubiquitin associated (phi-32, uba-1, C53A5.6) and proteasomal subunits (pas-4, pas-5, pas-6, pas-2, pas-3, pas-4, pas-5, pas-6, pas-7, rpt-1, rpt-3, rpt-4, rpt-5, rpt-6, rpm-1, rpm-2, rpm-6, rpm-7, rpm-8, and rpm-11). Inhibition of the proteasome by small molecules has previously been shown to induce the HSR [45,46]. It is intriguing that only the proteasome, and not autophagy or other proteases, functions as a regulator of the HSR, given the large number of components involved in protein clearance. The final class of regulators are involved in protein synthesis (dars-1) and gene expression (W04A4.5, pyp-1, and mdt-15). Microarray results confirm the induction of HSR genes upon mdt-15 knockdown [47]. While the pyp-1 subunit of the NuRF chromatin remodeler has not been previously linked to the HSR, other subunits of NuRF have been suggested to positively affect HSR gene expression [48]. Because we identified only one of 171 predicted E3-ligases (C53A5.6) and only one of 33 predicted tRNA

### Table 1. Positive Regulators of the HSR.

| Cosmid   | Gene       | Description       |
|----------|------------|-------------------|
| ZK328.2  | eftu-2     | EF-2 like         |
| C26C6.5  | dcp-66     | NuRD subunit      |
| F09D1.1  | F09D1.1    | USP39             |
| T08A11.2 | phi-11     | Splicing Factor 3B, subunit 1 |
| T13H5.4  | phi-8      | Splicing Factor 3A, subunit 3 |
| T28D9.10 | snr-3      | SNRPD1            |
| Y53C10A.12 | hsf-1     | Heat Shock Factor |

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Table 2. Negative regulators of the HSR and hsp70:gfp reporter induction.

| Cosmid   | Gene   | Description                        | S | I  | M   |
|----------|--------|------------------------------------|---|----|-----|
| F26D10.3 | hsp-1  | HSP70 Chaperone                    | ● | ●  | ●   |
| C47EB.5 | daf-21 | HSP90 Chaperone                    | ● | ●  | ●   |
| T05C12.7 | cct-1  | CCT/TRIC Chaperone                 | ○ | ○  | ●   |
| T21B10.7 | cct-2  | CCT/TRIC Chaperone                 | ○ | ○  | ●   |
| F54A3.3  | cct-3  | CCT/TRIC Chaperone                 | ○ | ●  | ●   |
| K01C8.10 | cct-4  | CCT/TRIC Chaperone                 | ○ | ○  | ●   |
| C07G2.3  | cct-5  | CCT/TRIC Chaperone                 | ○ | ●  | ●   |
| F01F1.8  | cct-6  | CCT/TRIC Chaperone                 | ○ | ●  | ●   |
| T10B5.5  | cct-7  | CCT/TRIC Chaperone                 | ○ | ●  | ●   |
| Y55F3AR.3| cct-8  | CCT/TRIC Chaperone                 | ○ | ●  | ●   |
| R05F9.10 | sgt-1  | TPR cochaperone                    | ○ | ●  | ●   |
| F30H5.1  | unc-45 | TPR cochaperone                    | ○ | ○  | ●   |
| T01B7.4  | cyn-11 | Cyclophilin cochaperone            | ○ | ●  | ●   |
| C36B1.4  | pas-4  | Proteasome 205 subunit             | ● | ●  | ●   |
| F25H2.9  | pas-5  | Proteasome 205 subunit             | ● | ●  | ●   |
| CD4.6    | pas-6  | Proteasome 205 subunit             | ● | ●  | ●   |
| C47B2.4  | pbs-2  | Proteasome 205 subunit             | ● | ●  | ●   |
| Y38A8.2  | pbs-3  | Proteasome 205 subunit             | ● | ●  | ●   |
| T20F5.2  | pbs-4  | Proteasome 205 subunit             | ● | ●  | ●   |
| K05C4.1  | pbs-5  | Proteasome 205 subunit             | ● | ●  | ●   |
| C02F5.9  | pbs-6  | Proteasome 205 subunit             | ● | ●  | ●   |
| F39H11.5 | pbs-7  | Proteasome 205 subunit             | ● | ●  | ●   |
| C52E4.4  | rpt-1  | Proteasome 195 subunit             | ● | ●  | ●   |
| F23F12.6 | rpt-2  | Proteasome 195 subunit             | ● | ●  | ●   |
| F23F11.8 | rpt-3  | Proteasome 195 subunit             | ● | ●  | ●   |
| F56H1.4  | rpt-4  | Proteasome 195 subunit             | ● | ●  | ●   |
| Y49E10.1 | rpt-6  | Proteasome 195 subunit             | ● | ●  | ●   |
| T22D1.9  | rpn-1  | Proteasome 195 subunit             | ● | ●  | ●   |
| C23G10.4 | rpn-2  | Proteasome 195 subunit             | ● | ●  | ●   |
| F57B9.10 | rpn-6  | Proteasome 195 subunit             | ● | ●  | ●   |
| F49C12.8 | rpn-7  | Proteasome 195 subunit             | ● | ●  | ●   |
| R12E2.3  | rpn-8  | Proteasome 195 subunit             | ● | ●  | ●   |
| K07D4.3  | rpn-11 | Proteasome 195 subunit             | ● | ●  | ●   |
| F57B10.1 | let-607 | Transcription Factor (ER)          | ● | ●  | ●   |
| C15H9.6  | hsp-3  | HSP70 Chaperone (ER)               | ● | ●  | ●   |
| F38A1.8  | SRP receptor α subunit             | ● | ●  | ●   |
| R186.3   | SRP receptor β subunit             | ● | ●  | ●   |
| F55C5.8  | SRP subunit                        | ● | ●  | ●   |
| F08D12.1 | SRP subunit                        | ● | ●  | ●   |
| F25G6.8  | SRP subunit                        | ● | ●  | ●   |
| F38E11.5 | COPI β subunit                     | ● | ○  | ●   |
| T14G10.5 | COPI γ subunit                     | ○ | ○  | ●   |
| T24H7.2  | HSP70 Chaperone (ER)               | ○ | ○  | ●   |
| C37H5.8  | hsp-6  | HSP70 Chaperone (mito)             | ○ | ○  | ●   |
| T09B4.9  | TIM44 subunit (mito)               | ○ | ○  | ●   |
| C47E12.5 | uba-1   | E1 ubiquitin ligase                | ○ | ○  | ●   |
| F52C6.3  | phi-32  | Ubiquitin                          | ○ | ●  | ●   |
| C53A5.6  | E3 ubiquitin ligase                | ○ | ○  | ●   |
| B0464.1  | dars-1  | Asp tRNA Synthetase                | ○ | ○  | ●   |

Table 2. Cont.

| Cosmid     | Gene  | Description      | S | I  | M   |
|------------|-------|------------------|---|----|-----|
| W04A4.5    | Integrator subunit | ○ | ●  | ○   |
| C47E12.4   | pyp-1 | NuRF subunit     | ○ | ●  | ○   |
| R12B2.5    | mtd-15 | Mediator subunit | ○ | ●  | ○   |

S = Spermatheca, I = Intestine, M = Muscle, ○ = Induction, ● = No Induction.

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HSP Negative Regulators Exhibit Distinct Tissue-Selective Patterns

A striking feature of the negative regulators of the HSR is that the HSR reporter in not uniformly induced across all tissues, but rather displays tissue-selective expression patterns in the intestine, muscle, and spermatheca (Figure 2, Table 2). Of the negative regulators, only knockdown of HSP70 and HSP90 induced expression of the reporter in all three tissues. By comparison, knockdown of the three chaperones and all eight subunits of the TRIC/CCT chaperone machine induced the reporter only in the muscle. In contrast, downregulation of subunits of the proteasome and many secretory pathway genes induced the reporter only in the intestine and spermatheca, but not in the muscle. Knockdown of the remaining genes induced the reporter only in the intestine. These patterns are unlikely to be due to RNAi artifacts because the tissue-selective patterns of reporter induction were similar for all subunits within specific complexes, yet non-overlapping between different complexes (i.e., all proteasomal subunits induced in intestine and spermatheca, and all TRIC/CCT subunits induced in the muscle).

Tissue-selective expression of the HSR reporter was unexpected as nearly all of the negative regulators are ubiquitously expressed components of essential cellular machines. For example, even though proteasomal subunits do not induce the HSR in muscle, it has been shown in C. elegans that most, if not all, proteasomal subunits are expressed in muscle and that RNAi knockdown of proteasomal subunits yields muscle specific phenotypes such as stabilization of a ubiquitin-GFP reporter in the muscle and early onset aggregation of a polyQ disease model expressed only in the muscle [28,51]. These results also suggest that depletion of subunits from complexes such as the proteasome does not induce the HSR simply by misfolding other subunits in that complex since these effects would not be expected to have tissue selectivity.

To further investigate the tissue-selective patterns of HSR regulation, we examined the expression of two additional reporters that are inducible by heat shock and dependent on HSF1. The pshp-16.2::gfp reporter is inducible in the intestine, muscle and excretory system, and is dependent on HSF1 and DAF-16 (Figure S1) [32,53]. The pckb-2::gfp reporter is inducible only in the intestine and is also activated by the unfolded protein response, an ER stress response [54]. Knockdown of HSR negative regulators revealed highly overlapping patterns of tissue-specific induction with all three reporters (Table 3). In the muscle, there was a highly consistent pattern of induction between the C12C8.1 and hsp16.2 reporters, with two genes inducing both, seven inducing neither, and only a single gene showing differential induction. In the intestine, HSR negative regulators gave identical patterns of induction for the C12C8.1 and the ckb-2 reporters, with nine out of
ten causing induction, yet only a smaller subset, three out of ten, also induced the hsp16.2 reporter. Given the differences in the regulation and function of the three genes, the three reporters demonstrate remarkably consistent patterns of tissue-selective HSR induction.

We next validated the tissue-selective effects using pharmacological inhibitors and mutants. We found that incubation of L4-staged worms with MG132, a pharmacological inhibitor of the proteasome, caused induction of the phsp70::gfp reporter in the intestine and the spermatheca, but not in the muscle tissue (Figure 3A). This pattern matches that seen with RNAi knockdown of proteasomal subunits, providing further support that the tissue-selective effects are unlikely to be an RNAi artifact. Most of the negative regulators are essential, however we were able to test the effects of mutations in T24H7.2, an ER localized HSP70, and the cochaperones unc-45, sgt-1, and cyn-11, and found using qRT-PCR that the expression of endogenous HSR genes was induced (Figure 3B). We further demonstrated that this induction was tissue-selective using qRT-PCR analysis on dissected intestinal cells. We found that T24H7.2 mutant animals, but not unc-45 mutant animals, induced endogenous hsp70 in the intestine (Figure 3C). The tissue-selective induction of endogenous genes in the intestine by these mutations matched the induction of the phsp70::gfp reporter by RNAi knockdown, thus providing a validation of both the use of RNAi and the fluorescent reporter.

Analysis of the HSR Regulatory Network

The genes that we identified form a genetic regulatory network of the HSR in C. elegans. To characterize the relationship between these regulators, we utilized an interaction network from previous physical, genetic, and predicted interaction data [55]. A network

Figure 2. Tissue-selective induction of the phsp70::gfp reporter by knockdown of negative regulators. Nomarski and fluorescent images corresponding to whole animals and fluorescent images of the spermatheca, intestine, and muscle tissue are shown. The boundary of the animals, intestine and spermatheca taken from Nomarski images are added as a visual aide to some images. (A–E) RNAi knockdown of daf-21 leads to induction of the reporter in all three tissues. (F–J) RNAi knockdown of cct-1 causes induction only in muscle. (K–O) knockdown of F38A1.8 causes induction in the intestine and spermatheca. Images are taken at different exposures to maximize fluorescence of each image. Scale bars of whole animal images correspond to 100 µm, while scale bars of the images depicting specific tissues correspond to 50 µm. Asterisks denote only autofluorescence.

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Table 3. Tissue-selective patterns for multiple HSR reporters.

| Gene  | Description | hsp70 | ckb-2 | hsp16.2 |
|-------|-------------|-------|-------|---------|
| hsp-1 | HSP70       | ●     | ●     | ●       |
| daf-21| HSP90       | ●     | ●     | ●       |
| cct-1 | CCT/TRIC    | ○     | ●     | ○       |
| pas-4 | Proteasome  | ●     | ○     | ○       |
| let-607| Transcription Factor | ○ | ○ | ○ |
| F38A1.8| SRP receptor α | ● | ○ | ○ |
| hsp-6 | HSP70 (mito) | ● | ● | ○ |
| C53A5.6| E3 ubiquitin ligase | ● | ● | ○ |
| dars-1| tRNA Synthetase | ● | ● | ○ |
| pyp-1 | NuRF        | ●     | ●     | ●       |

I = Intestine, M = Muscle, ● = Induction, ○ = No Induction.

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representation of the interaction data, in which HSR regulators are nodes and interactions between them are edges, reveals that HSR negative regulators are enriched in interactions with other HSR negative regulators: 39 of 52 negative regulator genes are connected in a single interaction network (Figure 4). We next applied a community detection algorithm to determine the structure of this interaction network [56,57]. This analysis shows that the network is composed of three distinct modules, indicated by the shapes of the nodes. The modular structure of this network is unlikely to have arisen by chance since it does not appear in randomized networks containing the same number of nodes and connections (p<10^-4). The three modules are primarily composed of clearance, cytoplasmic protein folding, and gene expression and protein synthesis components, respectively. While it is unsurprising that proteasome or protein folding subunits cluster together into distinct modules, the existence of the third module is entirely unexpected. The modules identified using the interaction data (node shapes) correspond closely with the observed tissue patterns of reporter induction (node colors) thus providing additional validation of both the specificity of tissue expression and network structure. These results further suggest that the underlying functional modules give rise to the tissue-specific patterns of HSR induction.

To further probe the genetic properties of the HSN, we investigated the relationships between the positive and negative regulators to provide a systems-level pathway analysis. We tested whether depletion of positive regulators (that decrease reporter induction by heat shock) would suppress reporter induction mediated by depletion of negative regulators. We found that knockdown of each positive regulator prior to knockdown of the negative regulator hsp-1 (HSP70) decreased induction of the reporter (Figure 5A). This indicates that the positive HSR regulators are epistatic to HSP70. These data are consistent with a model in which the positive regulators of the HSR act at or downstream of chaperone-mediated regulation of the HSR. Similar results were obtained for other negative regulators including daf-21 (HSP90), pas-4 (proteasome), C53A5.6 (E3 ubiquitin ligase), let-607 (ER transcription factor), F38A1.8 (SRP), hsp-6 (mitochondrial HSP70), and dars-1 (tRNA synthetases).

These results, in addition to the tissue-independent nature of the positive regulators and their association with biosynthetic processes, further distinguish the roles of the positive regulators from the negative regulators. RNAi knockdown is not equivalent to genetic ablation, so these relationships correspond to sensitivities rather than absolute dependencies. Therefore, we tested the effects of depletion of the positive regulators in a strain containing a deletion of the negative regulator unc-45, causes induction in the intestine and spermatheca (arrows), similar to RNAi knockdown of proteasomal subunits. The scale bar corresponds to 200 μm. Asterisks denote only autofluorescence. B) Mutations in T24H7.2, sgt-1, cyn-11, and unc-45 cause induction of the HSR in whole worms measured using qRT-PCR. C) Mutation of T24H7.2, but not unc-45, causes induction in the intestine relative to N2 control animals, measured by qRT-PCR analysis of hsp70 in dissected intestinal tissue. Averages shown are from at least two biological replicates.

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Discussion

Our findings indicate that the heat shock regulatory network (HSN) enables the HSR to sense and respond to a wide range of disruptions in proteostasis, thus providing a direct link between the function of the HSR and its regulation. The four functional clusters of the HSN each identify a small subset of the entire proteostasis machinery that functions in gene expression and protein synthesis, folding, trafficking, and clearance. The negative regulators fall within each of these functional categories, whereas the positive HSR regulators are more restrictive and cluster only to gene regulation and protein synthesis. Previous studies on the mitochondrial stress response have revealed that depletion of
Figure 4. Network analysis of HSR regulators. Shown is a network with HSR negative regulator genes depicted as nodes and interactions as edges. Node shape denotes grouping corresponding to a community detection algorithm based on the structure of the interaction network. Node color corresponds to the tissue-specific phsp70::gfp reporter induction. Cartoons of worms depicting the tissue specificity appear next to nodes containing those colors.

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specific subunits of electron transport chain complexes leads to induction of the mitochondrial stress response; however in our screen we did not identify subunits of macromolecular complexes as regulators of the HSR, thus revealing differences in how these two compartments detect and respond to a proteostatic imbalance [58].

The negative regulators of the HSN displayed a surprising extent of tissue-selective effects on HS gene expression, which may arise from differences in the expression levels or activity of the regulators between tissues. One clear example of differential tissue specific expression of HSR regulators occurs in activated B cells, that rely heavily upon the secretory pathway and exhibit high expression levels of secretory pathway components (for a review, see [59,60]). But as described in the results section, nearly all components of the HSN are broadly expressed. For example, TRiC/CCT is required for the folding of actin and tubulin, which are expressed in every cell [61]. However, the specialized function of muscle tissue could necessitate a stronger dependency for actin and myosin, which in turn explains the functional requirement for TRiC/CCT and account for the enhanced sensitivity of muscle to TRiC/CCT depletion. In addition to differential sensitivity to the regulators, our results indicate that each tissue exhibits distinct profiles of HS-inducible genes, which likely arises from tissue-specific factors that influence HS gene inducibility. Together, these data indicate that in addition to its unique proteome and specialized function, each tissue may contain a distinct complement of the proteostasis machinery, a differential sensitivity to disruption of proteostasis networks, and a distinct response to proteostasis disruption.

Figure 5. Epistasis analysis of HSR regulators. The effects of HSR positive regulator knockdown on induction of the reporter by negative HSR regulator knockdown were measured using the phsp70::gfp reporter. (A) Images showing the results from double RNAi with each positive regulator and the negative regulator hsp-1. In each case, knockdown of the positive regulator decreased reporter fluorescence compared to knockdown of hsp-1 alone. (B) Quantitation of the effects of HSR positive regulator knockdown using RNAi on induction of endogenous HSR genes by the HSR negative regulator T24H7.2 mutant reveals that the positive regulators are epistatic to T24H7.2.

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A

Positive Regulator RNAi

Control  hsf-1  eftu-2  dcp-66

Negative Regulator RNAi: hsp-1

phi-11  F09D1.1  snr-3  phi-8

phsp70::gfp

B

Relative mRNA levels

Control  dcp-66  eftu-2  hsf-1  phi-8  phi-11  F09D1.1  snr-3

Positive Regulator RNAi

Figure 5. Epistasis analysis of HSR regulators. The effects of HSR positive regulator knockdown on induction of the reporter by negative HSR regulator knockdown were measured using the phsp70::gfp reporter. (A) Images showing the results from double RNAi with each positive regulator and the negative regulator hsp-1. In each case, knockdown of the positive regulator decreased reporter fluorescence compared to knockdown of hsp-1 alone. (B) Quantitation of the effects of HSR positive regulator knockdown using RNAi on induction of endogenous HSR genes by the HSR negative regulator T24H7.2 mutant reveals that the positive regulators are epistatic to T24H7.2.

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Figure 6. HSR regulatory network model. Each HSR regulator, denoted by common terminology, is indicated as a box and is grouped according to its presence in a multi-subunit complex or functional pathway (i.e., the proteasome or secretory pathway). Positive or negative effects on HSR regulation are indicated by either a green arrow or red line respectively. Positive regulators are further separated from negative regulators by grey shading in the background. At the center of the network, HSF1 integrates signals from the various regulators and establishes a coordinated HSR.

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would be predicted to suppress protein aggregation. Instead, there is substantial overlap (twenty out of fifty-two genes) between the negative regulators of the HSR and a separate genome-wide screen for enhancement (early onset) of polyQ aggregation in muscle (Table S1). The common gene set includes the TRIC/CCT chaperonin (6), HSP70, mitochondrial HSP70, the proteasome (10), ubiquitin, and the E1 ubiquitin ligase. Therefore, it is likely that knockdown of these genes leads to both a disruption of proteostasis and activation of the HSR. Consistent with this model, there is almost no overlap with another genome-wide screen for suppression of polyQ aggregation in muscle. In contrast, knockdown of positive regulators, which suppresses the HSR, would be predicted to cause early-onset polyQ aggregation, and indeed, five of the seven positive HSR regulators have this phenotype and none suppress polyQ aggregation.

The paradigm for HSR regulation has previously focused on HSF1 and the negative feedback loops consisting of the HSP70 and HSP90 chaperones. Our results reveal that the network that regulates the HSR is much larger and corresponds to at least fifty-nine genes of this newly defined HSN. Many of these genes have been previously linked to HSR regulation in other systems, including prokaryotes, suggesting that this regulatory network is likely conserved through evolution. The precise mechanistic links between many of these genes and the HSR and other components of the HSN remains to be defined, and it will be important to investigate whether the tissue-selective regulation of the HSR is also conserved. Nevertheless, the identification of these genes in a comprehensive genetic screen for HSR regulators not only validates their functional properties but also reinforces the evolutionary conservation of the HSR. In summary, the systems-level identification and characterization of the HSR regulatory network described in this paper provides several important insights into regulation of the HSR during stress and provides a basis for future analysis of HSR regulation during development, ageing, and human disease.

Materials and Methods

Nematode Strains and Cultures

Nematodes were handled and analyzed using standard laboratory techniques and cultured at 20°C (62). Worms were synchronized by bleaching with hypochlorite (NaOCl) and hatching overnight in M9. Where indicated, intestines were dissected from living animals in M9 media. All nematode strains were derived from the N2 Bristol wild-type strain. The following strains and cultures were used: (1) AM446 rol-6(su1006); RB1694 rol+(e907) I; (2) JH1005 zc59 [phsp-4::gfp]; (3) CL2070 dvl-1(n70) [phsp-16.2::gfp]; (4) BC14636 dpy-5(e907) I; (5) CB13872[Ce.B0285.3::gfp+;Ceh361]; (6) PS3551 hsf-1(yu441); (7) AM658 hsf-1[yu441]; rol-6 (su1006); (8) RB1694 T2H17.2(ok2107) II; (9) RB703 unc-4(e3(ok468) III; (10) RB1053 rolF9.10(ok1000) II; and (11) VC1372, rol-21(ek178) II [13,37,40,52,63].

RNAi

Genome-wide RNAi screening was performed using a bacterial feeding approach with a library targeting approximately 86% of the C. elegans genome (MRC Geneservice, Cambridge, U.K.). Bacterial cultures were grown overnight in LB with 5 μg/ml tetracycline and 50 μg/ml ampicillin and induced with 1 mM IPTG for four hours. To avoid L1-stage developmental arrest associated with essential genes, L1 larvae were allowed to develop for 19 hours on plates containing OP50 bacteria prior to exposure to RNAi.

The genome-wide screen was performed in 96-well liquid cultures containing approximately 10 animals, 50 μl M9, 5 μg/ml cholesterol, 5 μg/ml tetracycline, 50 μg/ml ampicillin, 0.4 mM IPTG, 0.1 μg/ml fungizone, and 75 μl of RNAi bacterial suspension and grown at 20°C for 60 hours in a temperature-controlled shaker. For the heat shock screens, the animals were sensitized by exposure for two hours at 24°C, 24 hours before screening for reporter induction. The heat shock conditions are at 31.5°C for two hours followed by 24 hours of recovery at 20°C prior to screening for stress-induced fluorescence. Screening was performed using Leica MZ16 FA fluorescence microscope equipped with a GFP2 filter.

Validation and analysis of the regulators from the primary screen were done using solid RNAi plates containing nematode growth medium (NGM) agar with 5 μg/ml tetracycline, 50 μg/ml ampicillin and 1 mM IPTG and seeded with RNAi bacteria. Synchronized worms grown on OP50 bacteria for 19 hours were incubated on RNAi plates for 48 hours before analysis of induction (negative regulators) or wrapped in parafilm and heat shocked in a water bath at 33°C for 1 hour and then recovered for 24 hours prior to analysis (positive regulators). Worms were immobilized in levamisole and imaged using either a BD Pathway 435 High-content Bioimager (BD Biosciences) or a Zeiss Axiovert 200 fluorescent microscope. A gene was scored as positive only if >20% of animals demonstrated induction. Epistasis analysis was performed by knockdown of each positive regulator as before followed by double RNAi of the positive and negative regulators together.

Each RNAi construct was validated by sequencing. Functional information on the identified genes was collected using WormBase [64].

Drug Assay

Pharmacological inhibition of the proteasome was conducted using transgenic animals carrying the phsp70::gfp reporter grown on standard NGM plates seeded with OP50 bacteria. L4 larval stage animals were incubated with 100 μM MG132 (AG Scientific) in 0.5% DMSO or 0.5% DMSO alone for 2–3 hours and returned to plates. Fluorescence was scored the next day in young adult animals.

Fluorescence Imaging and Tissue Identification

Transgenic animals carrying the fluorescent reporter were mounted on 3% agarose pads, immobilized with 2 mM levamisole and viewed using the Zeiss Axiovert 200. Animals were imaged using 10X/0.25 A-Plan and 100X/1.4 oil DIC Plan-APOC-HROMAT objectives. Images were captured using a Hamamatsu digital camera (C4742-98) with Axiovision Release 4.7 software. Tissue-identification was based on nematode anatomy and tissue morphology using images from the C. elegans atlas [63]. A tissue was scored as positive only if >20% of animals demonstrated induction.

qRT–PCR

RNA was isolated from whole animals lysed by vortexing for twenty minutes after addition of TRIzol (Invitrogen) and DNA was removed using a DNA-free Kit (Ambion) according to standard protocols. cDNA synthesis was performed using an iScript cDNA Synthesis Kit (BioRad) and qRT–PCR was performed using an iQ SYBR Green Supermix Kit (BioRad) using provided protocols and run on a BioRad iCycler. 18S RNA was used as a normalization control.
Network Analysis
We utilized a graph partitioning scheme that separates the network into groups of nodes, which collectively maximizes the density of within-partition edges in the network [56,57]. The significance of the number of interactions between the negative regulators was tested by comparing their density to the density of interactions predicted genome-wide in C. elegans. The significance of the modularity of the HSR negative regulator network was tested by sampling Monte Carlo realizations in which we exchanged pairs of edges, maintaining the degree distribution of the network.

Supporting Information

Figure S1 Tissue-selective induction of the hsp-16.2 reporter (phsp-16.2::gfp) by knockdown of negative regulators. Nomarski and fluorescent images corresponding to whole worms and fluorescent images of the muscle tissue, excretory system, and intestine of the phsp-16.2::gfp reporter strain are shown. The boundaries of the animals taken from Nomarski images were added as a visual aid to some images. (A–E) In the absence of heat shock, the empty vector control showed no induction above background fluorescence. (F–J) Heat shock induces the reporter in all three tissues. (K–O) RNAi knockdown of hsp-1l leads to induction of the reporter only in excretory system and intestine; (P–T) knockdown of daf-21 leads to induction in only muscle; and (U–Y) knockdown of cct-1 leads to induction only in muscle, and (Z–AD) knockdown of F38A1.8 leads to induction only in the intestine. Images are taken at different exposures to maximize fluorescence of each image. Scale bars of whole animal images correspond to 100 μm, while scale bars of the images depicting specific tissues correspond to 50 μm. Asterisks denote only autofluorescence.

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Author Contributions
Conceived and designed the experiments: EG KR RM. Performed the experiments: EG DMC KR. Analyzed the data: EG DMC KR PDR RM. Contributed reagents/materials/analysis tools: PDM. Wrote the paper: EG KR RM.

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