

Cite this: *Mol. Omics*, 2019,
15, 108Received 22nd December 2018,
Accepted 13th February 2019

DOI: 10.1039/c8mo00283e

rsc.li/molomics

Small open reading frames and cellular stress responses

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Small open reading frames (smORFs) encoding polypeptides of less than 100 amino acids in eukaryotes (50 amino acids in prokaryotes) were historically excluded from genome annotation. However, recent advances in genomics, ribosome footprinting, and proteomics have revealed thousands of translated smORFs in genomes spanning evolutionary space. These smORFs can encode functional polypeptides, or act as *cis*-translational regulators. Herein we review evidence that some smORF-encoded polypeptides (SEPs) participate in stress responses in both prokaryotes and eukaryotes, and that some upstream ORFs (uORFs) regulate stress-responsive translation of downstream cistrons in eukaryotic cells. These studies provide insight into a regulated subclass of smORFs and suggest that at least some SEPs may participate in maintenance of cellular homeostasis under stress.

Introduction

The FANTOM genome annotation consortium initially relied on a 100 amino acid cutoff to distinguish eukaryotic protein coding sequences because a large number of spurious ORFs of shorter lengths occur randomly within long non-coding RNAs.^{1,2} In prokaryotes, a cutoff of 50 amino acids was used.³ However, with the advent of proteogenomic⁴ technologies, thousands of previously unannotated small open reading frames (smORFs)^{3,5} encoding products of fewer than 100 amino acids have been shown to undergo translation in organisms spanning all domains of life, including bacteria, yeast, flies, mouse, and human.^{6–17} With this increase in coding sequence annotation comes a need to determine the functions of smORF-encoded polypeptides (SEPs). Three classes of smORFs have been proposed in eukaryotes,¹⁸ based on RNA “location” and conservation: (1) non-functional intergenic smORFs that may represent newly evolving genes¹⁹ (2) smORFs that encode functional SEPs and (3) translated upstream ORFs (uORFs) encoded in 5′ untranslated regions of mRNA that function as *cis*-translational regulators of downstream coding sequences. Classes 1 and 2 may also be relevant to bacteria.

One-by-one characterization has shown that dozens of functional SEPs play roles in important biological processes, often by regulating the activity of macromolecular complexes.²⁰ Increasing evidence suggests that a subset of smORFs participate in

cellular stress responses.²¹ Cellular stress responses are evolutionarily conserved molecular responses to changes in environment that would otherwise disrupt homeostasis by damaging cellular molecules.²² These stresses can include temperature, reactive oxygen species, hypoxia, nutrient limitation, and other conditions to which cells must respond in order to survive. In this review, we first consider the functions of bacterial SEPs in stress response pathways (Fig. 1A), and secondly consider both functional and regulatory roles of eukaryotic SEPs.

smORFs and bacterial stress responses

Early evidence for the regulated expression of SEPs during cellular stress came from the study of prokaryotes, and a number of stress-response bacterial SEPs have been characterized both at the phenotypic and molecular levels.^{3,21,32} In this section, we discuss SEP expression during various stress responses, then detail the functions and mechanisms of selected stress-response SEPs in both Gram-negative and -positive bacteria.

Regulated smORF expression during cellular stress in bacteria

Bacterial responses to extracellular stress are governed both transcriptionally and post-transcriptionally.^{33–36} Transcriptional responses are mediated by dedicated transcription factors, such as σ^S /RpoS in Gram-negative and σ^B /SigB in Gram-positive bacteria, which are required for the general stress response (reviewed in ref. 35 and 36, respectively). Post-transcriptional regulatory mechanisms include small regulatory RNAs (sRNA),³⁴ RNA conformational changes,³⁷ and RNA binding proteins; unique among bacterial stress responses, the cold shock response is largely mediated by post-transcriptional mechanisms.³⁸ These transcriptional and post-transcriptional responses govern alterations

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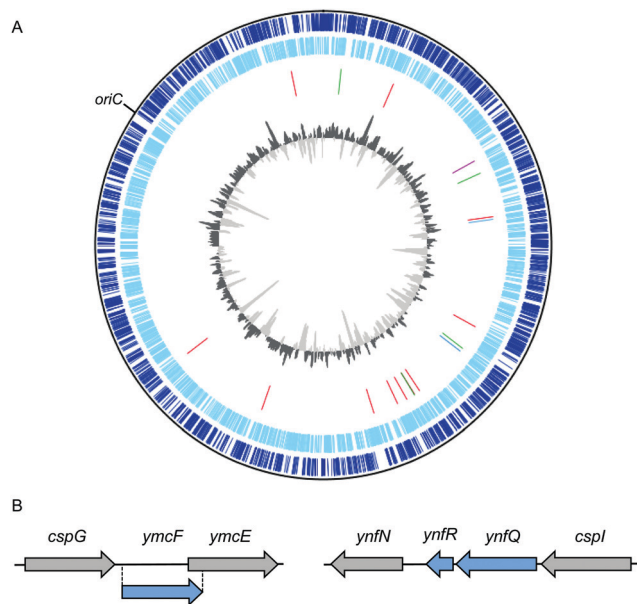


Fig. 1 Locations of stress-response associated small open reading frames (smORFs) in the *E. coli* str. K-12 substr. MG1655 genome. (A) Map of the *E. coli* str. K-12 substr. MG1655 genome. Tracks from the outside to inside represent: (1) annotated coding sequences within the forward strand (dark blue). (2) Annotated coding sequences within the complement strand (light blue). (3) Stress-responsive smORFs discussed in this review, by color: cold shock (blue),^{23,24} heat shock (red),^{21,25} antibiotic stress-inducible (purple),²⁶ and nutrient sensing (green).^{27–30} (4) Percent GC plot with above average GC content in dark gray and below average GC content in light gray. Genome sequence, annotated coding sequences, and stress-responsive smORFs (tracks 1–3) were uploaded to DNAPlotter version 1.0³¹ and selected to construct the map using NCBI RefSeq assembly accession: GCF_000005845.2. (B) Scale diagrams of the *cspG* and *cspl* genomic regions; previously annotated genes are depicted as gray arrows and recently discovered cold-inducible smORFs are depicted as blue arrows.^{23,24}

to the transcriptome, proteome, and metabolome that are required to re-establish homeostasis. Regulated expression of smORFs after exposure to a cellular stress has therefore led to the hypothesis that the encoded SEPs may function in the corresponding stress response.

The seminal observation by Storz and colleagues that ~40% of a set of 51 newly discovered *Escherichia coli* (*E. coli*) smORFs exhibited differential expression during stress responses, including heat shock, oxidative stress, and low pH, provided the first strong evidence that smORFs function during stress.²¹ Interestingly, some of these smORFs are post-transcriptionally regulated, such as *yobF* during heat shock. Importantly, subsequent phenotypic analysis in *E. coli* showed that deletion of three of these smORFs, *yqcG*, *ybhT*, and *yobF*, renders cells sensitive to envelope stress, and the *yobF* deletion strain was severely sensitive to acid stress.³² However, the molecular or biochemical function of YobF in response to heat shock, cell envelope stress, and acid stress has not yet been defined.

Proteomic and genomic approaches have subsequently been applied to identify additional temperature stress-regulated SEPs in *E. coli* K-12. Quantitative proteomics of small membrane proteins revealed an unannotated peptide mapping to a putative

smORF, *gndA*, that is encoded within the *gnd* gene in an alternative reading frame (and is therefore independent at the amino acid level).²⁵ Genomic tagging revealed that GndA expression is only detectable during heat shock. In parallel studies, three novel cold-inducible SEPs have been reported (Fig. 1B). Quantitative proteomics of *E. coli* K-12 revealed peptides YmcF and YnfQ which are specifically induced by cold shock.²³ These peptides map to two unannotated, intergenic sequences downstream of cold shock genes *cspG* and *cspI*, respectively. YmcF and YnfQ are upregulated by cold shock by up to a factor of 10, and exhibit 66% sequence identity, suggesting possible functional overlap. Interestingly, both of these cold-inducible smORFs initiate at AUU start codons, consistent with regulated expression.³⁹ Subsequent work by Hemm and coworkers identified an additional 21 amino acid smORF, *ynfR*, downstream of *ynfQ*, that is also cold-inducible.²⁴

SEPs are stress-inducible in diverse bacterial species. For example, three smORFs (*sbrABC*) recently discovered in *Staphylococcus aureus* are expressed in a SigB-dependent manner.⁴⁰ *sbrA* and *sbrB* encode SEPs that are 26 and 38 amino acids, respectively, while *sbrC* may encode a sRNA. In a second case, transcriptomic analyses of the photosynthetic cyanobacterium *Synechocystis* sp. PCC 6803 revealed three SEPs, Nsir6, Hlr1, and Norf1, that were induced by stress conditions, including transfer of the cyanobacteria from light to darkness.⁴¹ The *nsir6* and *hlr1* transcripts (nitrogen stress-induced RNA 6 and high light inducible RNA 1) were previously annotated as noncoding RNAs.

Antibiotic stress

Antibiotics activate several bacterial stress pathways and can induce the stringent response via (p)ppGpp signaling.⁴² Certain antibiotics and therapeutics such as ciprofloxacin and mitomycin C induce the SOS response.⁴³ A key antibiotic stress response linked to development of resistance is expression of drug efflux pumps.²⁶ The 49 amino acid membrane-bound AcrZ interacts with the AcrAB–TolC drug efflux pump, which exports some classes of antibiotics to confer resistance (Fig. 2a).²⁶ For example, strains lacking *acrZ* are sensitive to chloramphenicol and tetracycline, but not to erythromycin or rifampicin. While the mechanism of AcrZ is not fully characterized, AcrZ interacts directly with AcrB, which is hypothesized to lead to a conformational change in AcrB and export of specific antibiotics.²⁶

Nutrient sensing and utilization

Specific pathways have evolved to maintain homeostasis during nutrient stress, which can arise from either nutrient limitation or accumulation.³ An early report linking smORF expression to nutrient status showed that the 227 nt *sgrS* sRNA in *E. coli* is expressed during glucose 6-phosphate accumulation.²⁷ *sgrS* also encodes the 43-amino acid SEP SgrT (Fig. 2b).²⁷ The bifunctional *sgrS/sgrT* gene inhibits the glucose permease PtsG at both the RNA and protein level. Under conditions of high intracellular glucose 6-phosphate, the *sgrS* sRNA inhibits translation of the *ptsG* mRNA, while the SgrT SEP binds to PtsG and inhibits glucose uptake. Overexpression of SgrT renders cells incapable



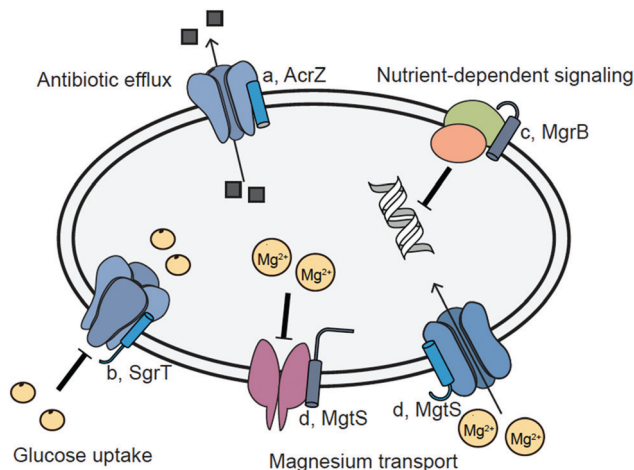


Fig. 2 Putative functions of selected membrane-bound bacterial stress-responsive microproteins. (a) AcrZ enhances export of specific antibiotics by the drug efflux pump AcrAB–ToIC;²⁶ (b) SgrT expression is induced by high intracellular levels of glucose 6-phosphate to inhibit glucose uptake;²⁷ (c) MgrB regulates PhoPQ (green and orange circles) in low intracellular Mg²⁺, decreasing expression of PhoPQ-dependent genes;²⁸ (d) MgtS increases Mg²⁺ uptake and prevents Mg²⁺ export.^{29,47}

of growth on glucose.⁴⁴ Interestingly, preliminary studies suggest that SEPs may be linked to monosaccharide utilization in other organisms, such as *Brucella abortus*, in which three recently identified, stress-inducible, membrane-localized SEPs increase cell growth rate on L-fucose.⁴⁵

Bacterial SEPs are also inducible and functional during divalent metal ion stress. When intracellular Mg²⁺ is low, PhoPQ upregulates gene expression, including the smORF *mgrB*.⁴⁶ *E. coli* MgrB (Fig. 2c), a 47-amino acid SEP, interacts with PhoQ to inhibit its autophosphorylation and activation.²⁸ Induction of the SEP MgtS (Fig. 2d) is also observed in a PhoPQ-dependent manner. MgtS co-purifies with the Mg²⁺ ATPase MgtA, leading to its stabilization and increased Mg²⁺ import.²⁹ This membrane-bound SEP also interacts with the PitA cation-phosphate transporter to prevent Mg²⁺ export.⁴⁷ In contrast, accumulation of Mn²⁺ can be toxic to cells.⁴⁸ The SEP MntS is repressed by the manganese-dependent transcriptional regulator MntR at high manganese, and overexpression of MntS leads to increased manganese sensitivity.³⁰ MntS may function to increase intracellular Mn²⁺ at low Mn²⁺ concentrations.⁴⁹

Prli42 and the *Listeria monocytogenes* stressosome

The stressosome is a ~1 MDa cytosolic complex that regulates the general stress response in Gram-positive bacteria.⁵⁰ The stressosome senses extracellular stress and, through a previously undefined mechanism, initiates intracellular signaling to activate SigB. Cossart and colleagues recently utilized an N-terminalomics approach to identify Prli42, a membrane-associated, 31-amino acid SEP that binds to the stressosome subunit RbsR and anchors RbsR to the membrane.⁵¹ Loss of Prli42 or the Prli42–RbsR interaction renders cells sensitive to oxidative stress and decreases expression of virulence factors in *Listeria*, suggesting that Prli42 is required for signaling by the

stressosome during stress and host infection. Prli42 therefore provides a model of a SEP–protein interaction that regulates stress-response signaling in bacteria.

smORFs and eukaryotic stress responses

Upstream smORFs (uORFs) and translational regulation during stress

Translational regulation of the proteome is an important component of eukaryotic stress responses and may occur more rapidly than transcriptional responses; more expression-level changes occur at the protein level (several thousand genes) than at the mRNA level (hundreds of genes) during stresses such as glucose and oxygen deprivation.⁵² Generally, global protein translation is downregulated during cellular stress, while translation of a subset of stress-response proteins remains constant or increases.^{53–55} A specific class of eukaryotic smORFs – upstream ORFs (uORFs) – play a role in stress-dependent translational regulation of downstream cistrons.^{56–58} Recent global profiling studies in yeast, plants and mammals^{9,13,59,60} have shown that uORF translation is widespread, especially following cellular stress.⁶¹ Ribosome profiling of oxidatively stressed yeast results in rapid accumulation of ribosomes on transcripts bearing uORFs following five minutes of hydrogen peroxide exposure.⁶² This observation is paralleled in human cells affected by oxidative stress,⁶³ as well as oxygen and glucose deprivation.⁵²

The prevailing model of uORF-mediated translational regulation holds that translating a uORF prevents scanning and/or re-initiation at the downstream coding sequence. Re-initiation is dependent on the distance between the uORF and downstream cistron.^{64,65} While uORFs were initially reported to act as *cis*-translational inhibitors of downstream coding sequences within the same mRNA,^{56,66,67} it has become clear that uORFs can either down- or upregulate downstream protein translation depending on the uORF start codon. AUG-initiated uORFs typically compete for translation with their downstream ORFs under normal growth conditions.^{68,69} In contrast, uORFs initiating with near-cognate (non-AUG) start codons are more likely to exhibit positively correlated translation with downstream coding sequences.⁷⁰ Non-AUG initiated uORFs may also play a role in upregulating downstream proteins previously thought to undergo non-canonical initiation under stress conditions or global translational arrest, as demonstrated during nutrient starvation and meiosis.^{70,71} However, the presence or sequence of a uORF is not sufficient to predict translational regulation during stress.

uORF-mediated regulation of protein translation occurs as a result of changes in the pre-initiation complex. During the integrated stress response, the trimeric eIF2 complex, which is responsible for initiator tRNA delivery to the 40S ribosome, is repressed through phosphorylation of the eIF2 α subunit.⁷² This repression of eIF2 activity has several effects on translation: global protein translation is downregulated,⁷³ AUG-initiated uORFs are skipped by the preinitiation complex, relieving their



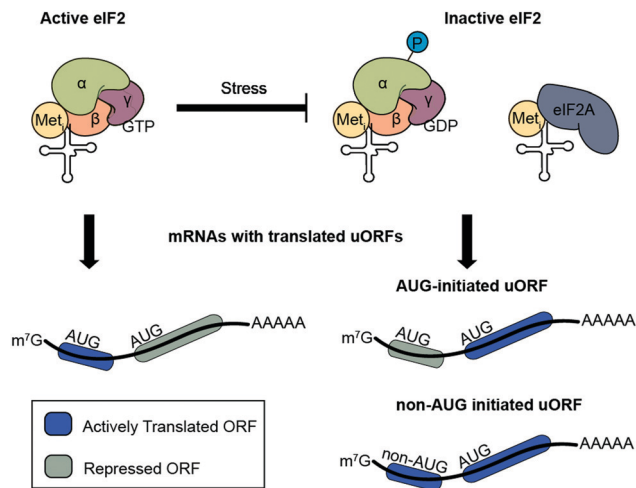


Fig. 3 Regulated translation of upstream open reading frame (uORF)-containing transcripts under cellular stress. Under normal conditions, active eIF2 is abundant and, in the subset of transcripts that contain them, AUG-initiated uORFs are translated, downregulating expression of the downstream ORF. Stress induces phosphorylation of the eIF2 α subunit and results in eIF2 inactivation. Limiting eIF2 concentrations cause ribosomes to bypass AUG-initiated uORFs and drive downstream ORF translation. Simultaneously, weak eIF2 competitor eIF2A can activate translation of non-AUG initiated uORFs in the transcripts that contain them.

inhibition of downstream protein translation,⁷⁴ and the weak eIF2 competitor eIF2A is de-repressed and delivers initiator tRNA to selected sites⁷⁵ including non-AUG codon-initiated uORFs,⁷⁶ driving their translation during stress (Fig. 3). For example, eIF2A drives translation of two uORFs initiating with UUG and CUG start codons and induces expression of the downstream cistron encoding binding immunoglobulin protein (BiP), an ER-resident chaperone vital for the activation of the integrated stress response.⁷⁶ This mechanism also operates in squamous cell carcinoma tumorigenesis, in which eIF2A-dependent translation drives a 1.8-fold increase in uORF occupancy by ribosomes.⁷⁷

uORFs are generally thought to compete for scanning ribosomes, which can then only initiate translation of downstream coding sequences *via* leaky scanning or re-initiation,⁷³ implying that the regulatory function of uORFs should depend only on their translation and therefore be independent of their sequences. In a few cases, however, the specific amino acid sequence of a uORF is required for its regulatory activity.^{78–80} An early report of this phenomenon described a uORF in the 5' untranslated region (UTR) of *DDIT3*, which encodes the CHOP protein, a transcription factor that promotes a switch from stress response signaling to cell death.⁸¹ Translation of the uORF alone is insufficient to recapitulate translational downregulation of CHOP, as introduction of nonsense and missense mutations within the uORF alleviated translational repression of CHOP, whereas silent mutations did not.⁸¹ Further mutational analysis defined an IPI motif within the uORF that promotes ribosome stalling to inhibit CHOP translation *in cis*.⁸² Fungal uORFs in the 5' UTR of arginine biosynthetic genes *ARG2* and *CPA1* also regulate downstream protein production *in cis* in a sequence-dependent manner *via* ribosome stalling.^{83–87}

Taken together, these studies show that uORF translational regulation plays a key role in proteomic reprogramming during cellular stress responses. While several uORFs have been reported to sequence-specifically induce ribosome stalling, translated products of uORFs have generally been assumed to lack function at the polypeptide level (though the uORF-encoded MIEF1 micro-protein, which binds to and regulates the mitochondrial ribosome, presents a counterexample⁷⁴). In contrast, conserved smORFs encoded in dedicated transcripts have been proposed to be functional,²⁰ and a number of these smORFs are involved in mediating stress responses.⁷⁶

Functional stress-response smORFs in eukaryotes

Characterization of SEPs that function in eukaryotic cellular and organismal stress responses is dramatically accelerating. Several recent reports have implicated SEPs in response to infection and innate immunity. First, ribosome profiling of influenza virus-infected human lung cancer cells identified 19 novel smORFs in long non-coding RNAs (lncRNAs) and other non-coding RNAs that were either up- or downregulated during infection.⁸⁸ Among these, a SEP translated from the host gene for miR-22, *MIR22HG*, was upregulated during infection with both wild-type influenza and NS1-mutant influenza that is rapidly cleared from cells due to interferon responses, suggesting that the *MIR22HG* SEP may respond to cellular stress due to viral particle exposure. More recently, ribosome profiling was applied to identify differential translation of lncRNA-encoded smORFs in lipopolysaccharide (LPS)-treated mouse macrophages.⁸⁹ An LPS-upregulated smORF within the lncRNA *Aw112010* encodes a CUG-initiated SEP that drives interleukin-12 beta expression. Characterization of a knockout mouse demonstrated that the *Aw112010* SEP is essential for mucosal immunity during both *Salmonella* infection and colitis. While the molecular mechanisms of the *MIR22HG* and *Aw112010* SEPs remain uncharacterized, these studies provide a link between SEP expression and infection in cells and *in vivo*.

The SEP humanin has been reported to protect cells from stress-induced apoptosis. Humanin was first discovered in 2001 as a neuroprotective factor in Alzheimer's disease, conferring neuronal resistance to apoptosis by a disease variant of the amyloid precursor protein.⁹⁰ Humanin has subsequently been reported to play additional intracellular roles in suppressing apoptosis *via* Bax binding and inactivation.⁹¹ While these functions suggest that humanin is protective against apoptosis downstream of cellular stress, it remains unclear how humanin is produced in cells, as its coding sequence may map to either mitochondrial or genomic DNA.⁹¹

Extensive work has identified SEPs that participate in muscle regeneration following injury. DWORF,⁹² a 34-amino acid SEP that localizes to the sarcoplasmic reticulum membrane, was identified in a lncRNA exhibiting heart- and muscle-specific expression (Fig. 4a). DWORF is downregulated at the protein and mRNA level during ischemic heart failure.⁹² DWORF normally functions to increase Ca²⁺ uptake into the sarcoplasmic reticulum *via* interaction with the Ca²⁺-ATPase SERCA and displacement of three other polypeptide inhibitors.^{93–95}



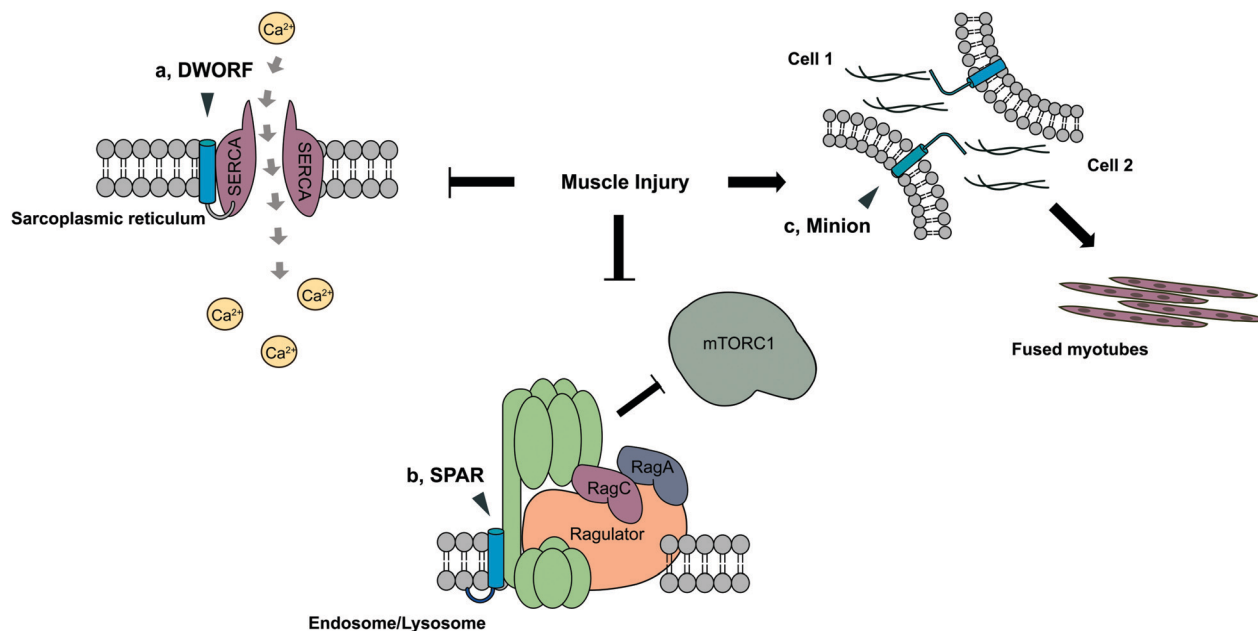


Fig. 4 Microproteins influence muscle regeneration following injury. (a) In uninjured muscle, DWORF binds the SERCA calcium pump and increases calcium flow into the sarcoplasmic reticulum. (b) In uninjured muscle, SPAR binds the ragulator v-ATPase and prevents mTORC1 activation. (c) Following injury, Minion mediates myoblast fusion.

Decreased contractility observed during heart failure can be caused by reduced Ca²⁺ levels in the sarcoplasmic reticulum resulting from insufficient activity of the SERCA pump.⁹⁶ Activation of SERCA through DWORF overexpression restored calcium levels and heart contractility in a mouse model of heart disease.⁹⁷ Another example is SPAR,⁹⁸ a SEP encoded by lncRNA *LINC00961* which is downregulated upon muscle injury (Fig. 4b). SPAR normally localizes to the endosome/lysosome membrane to promote association between lysosomal v-ATPase, Ragulator, and Rag GTPases, preventing mTORC1 activation. Upon muscle injury, SPAR downregulation promotes mTORC1 activation and muscle regeneration. Conversely, Minion⁹⁹ or Myomerger,¹⁰⁰ is a SEP which is transcriptionally upregulated in muscle tissue regeneration and development (Fig. 4c). Skeletal muscle development and regeneration following injury proceeds through temporally regulated stem cell activation and differentiation, myoblast fusion and subsequent maturation into myofibers,^{101,102} CRISPR/Cas9 knockdown of Minion results in defects in myoblast fusion, while homozygous mutants are unviable, most likely due to the inability to form multinucleate myotubes. In summation, differential expression of a suite of SEPs is required for response to injury in both cardiac and skeletal muscle.

Conclusion

Mounting evidence supports regulatory (in eukaryotes) and functional (in both prokaryotes and eukaryotes) roles for smORF translation in cellular stress responses. A future direction will be elucidation of the functional, molecular, and phenotypic roles of dozens of yet-uncharacterized SEPs that have been identified as differentially regulated during various stress conditions in a wide variety of organisms. While dozens of SEPs have been implicated

as differentially expressed at the RNA or protein level during stress responses, post-translational regulation of SEPs, especially *via* post-translational modifications (PTMs), has remained largely unaddressed. Given the importance of PTMs in stress signaling,^{73,103} identification of stress-regulated PTMs may be informative in elucidation of SEP functions. Finally, it is tempting to speculate that the small size of smORFs allows rapid translation, consistent with a need for rapid response to external stressors; measurements of the dynamics and abundance of SEP expression relative to the rate of production of known stress response proteins could test this hypothesis. Taken as a whole, the growing literature demonstrating roles for SEPs in cellular stress provides one testable hypothesis for characterization of newly discovered smORFs, and has also improved our understanding of the full complement of regulatory factors in stress response pathways.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported in part by the Searle Scholars Program, the Leukemia Research Foundation, the NIH (R01GM122984), and Yale University West Campus start-up funds (to S. A. S.). A. K. was supported in part by an NIH training grant (5T32GM06754).

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