Prion-based memory of heat stress in yeast

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ABSTRACT. Amyloids and amyloid-based prions are self-perpetuating protein aggregates which can spread by converting a normal protein of the same sequence into a prion form. They are associated with diseases in humans and mammals, and control heritable traits in yeast and other fungi. Some amyloids are implicated in biologically beneficial processes. As prion formation generates reproducible memory of a conformational change, prions can be considered as molecular memory devices. We have demonstrated that in yeast, stress-inducible cytoskeleton-associated protein Lsb2 forms a metastable prion in response to high temperature. This prion promotes conversion of other proteins into prions and can persist in a fraction of cells for a significant number of cell generations after stress, thus maintaining the memory of stress in a population of surviving cells. Acquisition of an amino acid substitution required for Lsb2 to form a prion coincides with acquisition of increased thermotolerance in the evolution of Saccharomyces yeast. Thus the ability to form an Lsb2 prion in response to stress coincides with yeast adaptation to growth at higher temperatures. These findings intimately connect prion formation to the cellular response to environmental stresses.

KEYWORDS. actin, amyloid, heat shock, Lsb1, Lsb2, prion, Sup35, ubiquitin, yeast (Saccharomyces cerevisiae)
INTRODUCTION

This extra-view is related to our recent paper describing a new yeast prion that can carry cellular memories of stress. Highly ordered fibrous cross-β protein aggregates (amyloids) and/or oligomers (protofibrils) involved in amyloid generation are associated with a variety of human and animal diseases, including Alzheimer (AD), Parkinson (PD) and Huntington diseases. Amyloids spread by immobilizing the protein molecules of the same sequence into amyloid fibrils and converting them into an amyloid form. Frequently, amyloids can be transmitted between cells within one organism, and sometimes even between organisms, resulting in protein-based infection. The prototype of an infectious amyloid is the prion protein (PrP), associated with transmissible spongiform encephalopathies (TSEs), such as sheep scrapie, mad cow disease, and human Creutzfeldt-Jakob and kuru diseases. Recent data show that many amyloid diseases possess at least some prion-like properties. Many amyloidoses, including the majority of the cases of AD and PD, are idiopathic and not associated with any DNA mutations. Although environmental agents have been linked to amyloidoses, e.g. certain pesticides to PD and certain metal ions to AD, systematic information about environmental factors promoting or inhibiting amyloids formation is lacking, resulting in the absence of clearly defined prophylactic strategies.

Notably, some amyloids play biologically positive functions. Amyloids help in the attachment of bacterial or fungal cells to the substrate and/or to other cells, possess structure-forming properties (e.g. silks), are used for storage (e.g. peptide hormones), and form scaffolds of covalent polymers, such as melanin. Possibly the most fascinating example of the biological consequences of amyloid-style polymerization is the role of self-perpetuating oligomers of CPEB protein in long-term memory. Indeed, amyloids and prions are capable of fixing and propagating change in protein conformation, making them efficient “memory machines.” How then can a prion-like mechanism be used for maintaining a memory of environmental changes?

A simple tractable model system is needed to investigate the impact of environmental factors on amyloids. Endogenous amyloids of the yeast Saccharomyces cerevisiae (termed “yeast prions”) provide such a system. About 10 yeast proteins convert to the amyloid-based prion form in vivo, and many more candidate prion proteins are suspected. Some yeast prions control detectable phenotypic traits, making them an excellent model system for studying general mechanisms of amyloid formation and propagation. At least 1/3 of wild or industrial isolates of yeast possess traits that are inherited in a prion-like fashion. Most yeast prion proteins contain domains (usually Q/N-rich), responsible for prion properties and termed prion domains, or PrDs. Notably, yeast cells can also form multimolecular assemblies that are not heritable but are maintained by a mother cell and carry a “memory” of certain events; memory of the deceptive mating encounter depends on the Whi3 protein. Prion-like QN-rich domains are involved in the formation of a maternally retained super-assembly in case of Whi3. This indicates that stably inherited yeast prions may represent only a tip of the iceberg, providing extreme examples of phenomena involved in a broader spectrum of cellular memories.

In spite of a heated discussion regarding the biological roles of prions, it is now becoming clear that both pathogenic, and beneficial prions can be found in yeast and other fungi. Some yeast prions control phenotypic traits that can be easily monitored. For example, [PrT+] a prion form of the yeast translation termination factor Sup35 (eRF3), causes a partial defect of translation termination due to aggregation of a termination factor. Formation of the prion results in translational readthrough of nonsense-codons (so called nonsense-suppression), allowing prion detection by growth on selective medium and/or by color on complete medium in the specially designed yeast strains. The availability of such convenient assays makes yeast a powerful system for studying the mechanisms of prion formation and maintenance.
HETEROLOGOUS PRION CROSS-SEEDING

Formation of a yeast prion de novo can be induced by a transient overproduction of the prion-forming protein or its PrD. However, typically this process is efficient only if there is another Q/N-rich protein aggregate in the same cell. For example, de novo formation of the Sup35 prion, \([\text{PSI}^+]\) after transient overproduction of Sup35 protein is promoted by the presence of a non-Mendelian element named \([\text{PIN}^+]\), for \([\text{PSI}^+]\) inducibility. This \([\text{PIN}^+]\) element, initially defined in genetic experiments, was later identified as a prion form of Rnq1 protein and named \([\text{RNQ}^+]\). Furthermore, overproduction of other aggregating Q/N-rich proteins can substitute for \([\text{RNQ}^+]\) and confer the \([\text{PSI}^+]\) inducibility (Pin^+ phenotype) to the yeast cells lacking the Rnq1 prion. Some of these proteins are known to form prions by themselves, however such evidence is lacking for others. Increased amyloid formation by one protein in the presence of the amyloid form of another protein has also been reported or suspected in mammalian systems, for instance, for tau and A\(_\beta\) proteins in case of AD.

The molecular foundations of cross-seeding on prion formation by other prion proteins are still insufficiently studied. Proposed models include: a) direct heterologous cross-seeding of one prion by another, and b) sequestration of folding cofactors (such as chaperones) that promotes misfolding and prion formation. Evidence exists for both mechanisms, and it is possible that different prions may act in different ways.

One of Q/N-rich proteins whose overproduction is shown to promote formation of the \([\text{PSI}^+]\) prion by excess Sup35 in strains lacking the \([\text{RNQ}^+]\) prion is the Lsb2 protein, also called Pin3. Lsb2 is a stress-inducible short-lived protein that is degraded via the ubiquitin-proteasome system. Lsb2 interacts with overproduced Sup35 protein and promotes its accumulation and aggregation at actin patches. Apparently Lsb2 can also promote formation of other Q/N-rich prions, as \([\text{RNQ}^+]\) isolates have also been obtained from cultures overproducing Lsb2. Lsb2 is associated with actin cytoskeleton through interaction with the Las17 protein (yeast homolog of WASP) via SH3 domains. Mutations in Lsb2 abolishing this interaction also abolish its ability to form cytologically detectable puncta, promote aggregation of Sup35 and formation of the Sup35 prion. Our new (previously unpublished) data confirm that excess Lsb2 is not able to promote \([\text{PSI}^+]\) induction in the las17\(\Delta\) strain (Fig. 1). Thus, association of Lsb2 with actin via Las17 is crucial for prion induction by Lsb2. Apparently, Lsb2 confines misfolded Sup35 (and possibly some other misfolded proteins) to the cytoskeleton-associated deposits, thus increasing local concentrations of Sup35 and facilitating prion nucleation. Notably, Lsb2 has a close paralog, Lsb1, which also interacts with Las17 but is not capable of promoting prion formation by Sup35.

Prion Cross-Seeding by LSB2 is due to Formation of a Metastable Prion, \([\text{LSB}^+]\)

Despite its ability to cross-seed other proteins into a prion state, earlier studies failed to detect prion formation by Lsb2 itself. In our recent paper, we have shown that transient overexpression of Lsb2 results in generation of a fraction (3–4%) of the cells that retain ability to induce conversion of Sup35 into \([\text{PSI}^+]\) after overexpression of Lsb2 is turned off. This prion-inducing phenotype was inherited through an indefinite number of mitotic divisions, however only in a fraction of the progeny, indicating a low mitotic stability. The prion-inducing phenotype was dominant and inherited in a non-Mendelian fashion through mating and meiosis, being retained by a fraction (17%) of ascospores. Semi-denaturing detergent agarose gel electrophoresis, SDD-AGE confirmed that cultures that inherited the prion inducing phenotype contain a small but detectable fraction of Lsb2 protein in the form of detergent-resistant polymers, similar to those formed (in a larger amount) in the cells containing overproduced Lsb2. We have designated this heritable aggregated state of Lsb2 protein as a metastable prion, \([\text{LSB}^+]\). The low stability of the \([\text{LSB}^+]\) prion could be due at
least in part to the proteolytic instability of Lsb2 protein, resulting in low levels of this protein in non-stressed cells. Indeed, mutants of Lsb2 that are deficient in ubiquitination, form [LSB+] aggregates that are inherited by a larger fraction of cells, compared with the aggregates formed by a wild type protein.\(^1\) This agrees with our finding that the efficiency of [PSI\(^+\)] induction by Lsb2 is increased by a defect in protein ubiquitination.\(^41\) Also, in agreement with previous data\(^45\) we have shown that the replacement of 8Q stretch by 8N stretch in the Lsb2 protein significantly increases prion-inducing efficiency and decreases average size (as typical of more efficiently propagated prions) of the [LSB+] aggregates. This agrees with the findings that an increase in the proportion of N residues versus Q residues favors more efficient prion propagation by QN-rich proteins.

### Effects of Environmental Stresses on Yeast Prions

Propagation of yeast prions in cell division occurs via repetitive cycles of fibril fragmentation and growth.\(^20,46,47\) Prion fragmentation depends strictly on the cellular chaperone disaggregation machinery, composed of the Hsp104, Hsp70 and Hsp40 proteins.\(^20,23,48-52\) Certain environmental conditions destabilize prions and cause prion loss, possibly due to alterations in the chaperone balance.\(^53,54\) Different prions, and even different variants (“strains”) of a prion formed by one and the same protein may exhibit different responses to stress. Probably the best studied case of prion destabilization by an environmental stress is an effect of a short-term mild heat shock (39–42°C) on some “weak” variants of the Sup35 prion, [PSI\(^+\)].\(^55\) Yeast [PSI\(^+\)] cells, exponentially growing at 25–30°C, then heat shocked for a short period of time (30–60 min) and shifted back to normal temperature, produce a small fraction of colonies that have lost prion and a large fraction (30–40%) of mosaic colonies, pointing to high frequency of prion loss in cell divisions following stress treatment. If yeast cells are incubated at high

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**FIGURE 1.** Association with Las17 is required for prion induction by Lsb2. A. Lsb2 cannot induce formation of prion [PSI\(^+\)] by overexpression of Sup35 in the strain depleted of Las17 (las17\(\Delta\)). Depletion of Las17 does not affect formation of [PSI\(^+\)] by overexpression of Sup35 in the presence of prion aggregates of Rnq1 protein ([RNQ\(^+\)]). [PSI\(^+\)] formation is manifested by growth on SD-Ade medium. B. Total levels of Lsb2 protein are slightly decreased in Las17 strain as detected by western blotting. Different time of Lsb2 induction from plasmid copper inducible promoter is indicated. C. Wild type Lsb2 and more stable ubiquitination deficient mutants (K80; K80R, K41R; P124A, P125A) do not form detergent resistant amyloid-like aggregates in the strain depleted of Las17 as detected by SDD-AGE. Lsb2 W91S mutant unable to bind Las17 does not form amyloid-like aggregates in the presence of Las17 (WT).
temperature for longer periods of time (2–4 hrs), prion destabilization is decreased. Maximal destabilization coincides with the period of maximal imbalance between the Hsp104 chaperone (that is present at very low background levels in exponential cultures and quickly accumulated during heat shock) and the Hsp70-Ssa chaperone (that takes a longer time to achieve a significant fold increase over relatively high background levels). This agrees with data showing that the stoichiometric Hsp104/70/40 complexes promote prion fragmentation and proliferation, while a selective increase in Hsp104 leads to $[PSI^+]$ loss.\textsuperscript{56,57} Analysis of individual cells indicates that prion loss is associated with an asymmetric distribution in post-heat shock cell divisions,\textsuperscript{43,55,58} that was explained either by asymmetry in distribution of prion aggregates per se\textsuperscript{20,55} or by asymmetric distribution of the chaperone Hsp104.\textsuperscript{58} These explanations are not mutually exclusive, as Hsp104 is known to be associated with asymmetrically distributed aggregates after stress,\textsuperscript{59} and excess Hsp104 promotes asymmetric partitioning of the $[PSI^+]$ prion units.\textsuperscript{60}

Importantly, Lsb2 protein is induced during the short-term heat shock,\textsuperscript{41,61} while its paralog Lsb1 is released from the association with ER membrane into a cytosol via heat shock inducible proteolytic processing.\textsuperscript{43} In the absence of either Lsb1 or Lsb2, heat shock mediated destabilization of $[PSI^+]$ is significantly increased.\textsuperscript{43} Moreover, cells lacking both Lsb proteins do not show an amelioration of prion destabilization during a longer exposure to high temperature. These data indicate that Lsb proteins antagonize the effect of stress on prion aggregates and/or asymmetric distribution of prion/chaperone complexes in cell divisions after stress.

Various environmental stresses were also reported to increase \textit{de novo} formation of yeast prions, such as $[URE3]$, $[RNQ^+]$, $[PSI^+]$, and $[MOT3^+]$.\textsuperscript{20,22,23,62-65} However, the molecular basis of the effects of environmental conditions on prion formation remains unknown in most cases. As Lsb2 is a stress-inducible protein, we have investigated if its prion-inducing properties are influenced by stress.

$[LSB^+]$ Prion Is Induced by Stress

Indeed, our results (1) show that about 0.1% of colonies produced by yeast cells that were heat shocked at 39°C exhibited a metastable $[PSI^+]$-inducibility phenotype, indicative of the $[LSB^+]$ prion. The frequency of such colonies was increased up to 0.6% in the strain containing the 8Qto8N derivative of Lsb2, but they were not detected in the $lsb2\Delta$ strain. These data confirm that the $[LSB^+]$ prion can be generated (without an artificial protein overproduction) as a result of the physiological increase of Lsb2 levels in response to heat stress.

Thus, Lsb2 serves as a sensor of stress, which can maintain a cellular memory of stress in subsequent generations via converting into a heritable (but metastable) prion form. It should be noted that while Lsb2 was proposed to play a role in endocytosis,\textsuperscript{66} this role is obviously minor and dispensable. In contrast, our data show that Lsb2 facilitates the assembly of other misfolded proteins at specific cytoskeleton-associated sites.\textsuperscript{41} This mechanism may have arisen as a protective tool intended to minimize the pathogenic effects of inducing misfolded proteins throughout the cell, and may also help prevent degradation of essential proteins under unfavorable conditions, as proposed previously.\textsuperscript{53} Indeed, analysis of individual cells shows that heat shock induced cell death is increased in the absence of both Lsb1 and Lsb2.\textsuperscript{43} As assembly of other aggregates occurs through interaction with punctate structure formed by aggregated Lsb2, this process is likely to be facilitated by the conversion of Lsb2 into a prion form. In addition, prion aggregates of Lsb2 are likely more proteolytically stable, compared with a non-prion protein, as judged from analogy with other known prions.\textsuperscript{57-69} Thus, cells bearing the $[LSB^+]$ prion likely contain more Lsb2 protein compared with $[lsb^-]$ cells, and also contain it in the form that is ready for performing its stress-related functions. The biological consequence of the formation of the metastable $[LSB^+]$ prion is a subpopulation of cells, which “remember” the stress and are therefore, may be better prepared for
the return of stress conditions. This increases the possibility of survival of the population as a whole in such a case.

Notably, the protective aggregate assembly mediated by Lsb2 in a prion form may generate self-perpetuating amyloid forms (prions) of other proteins as a by-product, as we observe in case of \([\text{PSI}^+]\) induction, and possibly, for \([\text{RNQ}^+]\).\(^{41}\) Indeed, increased \textit{de novo} formation of the \([\text{PSI}^+]\) prion has been detected in certain stress conditions, including incubation at high temperature.\(^{64}\) It is possible that induction of the transient \([\text{LSB}^+]\) prion contributes to this phenomenon. Other prions triggered by \([\text{LSB}^+]\) could be pathogenic, however the fast degradation of Lsb2 protein after stress (or after adaptation to stress conditions) and the metastable nature of the \([\text{LSB}^+]\) prion ensure induction of such prions would occur only in a fraction of the population and therefore minimize potential negative effects. On the other hand, stress-dependent induction of prion formation increases overall phenotypic variability and may generate prion-controlled traits that become enhance survival of a subpopulation of cells. Known examples of yeast prion increasing survival in certain stressful conditions include: \([\text{MOT3}^+]\),\(^{22}\) that increases resistance of yeast cells to ethanol stress, and \([\text{MOD}^+]\), that increases resistance to fluconazole.\(^{70}\)

### Evolutionary Acquisition of Prion-Forming Properties by Lsb2

The Lsb2 paralog, Lsb1 also directly interacts with Sup35 PrD in the yeast 2-hybrid assay\(^{43}\) and forms puncta that colocalize with the Sup35 aggregates.\(^{41,43}\) However in contrast to Lsb2, overproduction of Lsb1 does not promote prion formation.\(^{43}\) Surprisingly, we have found that the difference in a single amino acid residue N213 (Lsb2) vs. S239 (Lsb1) within the otherwise conserved C-terminal region confers the prion-inducing ability to Lsb2 in comparison to Lsb1 (Fig. 2).\(^{1}\) The N213S substitution completely abolished the ability of overproduced Lsb2 to promote \([\text{PSI}^+]\) induction (and therefore, its ability to form an \([\text{LSB}^+]\) prion), while the reciprocal S239N substitution enabled overproduced Lsb1 to become \([\text{PSI}^+]\) inducer. Remarkably, the N to S substitution disrupts an “amyloid stretch” consensus sequence formed by the C-terminal hexapeptide of Lsb2. Hexapeptides conforming to this consensus were shown to be a characteristic

![Prion–inducing activity of Lsb2](image.png)

**FIGURE 2.** Prion–inducing activity of Lsb2 coincides with yeast adaptation to higher growth temperature. Schematic shows phylogenetic relationships among some members of the *Saccharomyces sensu stricto* genus. The preferred growth temperature of each species is indicated.\(^{74,75}\) CLUSTALW-formatted multiple sequence alignment of C-terminal of Lsb1/Lsb2 is shown. Difference in amino acids is indicated in red (Lsb2) and blue (Lsb1). Residue essential for prion induction is bordered. Amyloid stretch hexapeptide is underlined.
feature of the proteins capable of forming amyloids in vitro. Lsb2 and Lsb1 possess 2 additional amyloid stretch hexapeptides, however only the presence of the third, C-proximal amyloid stretch (found only in Lsb2) strongly correlates with the prion-inducing ability (Fig. 2).

Notably, only the Saccharomyces sensu stricto clade contains 2 paralogs of Lsb, while other Saccharomyces species possess only Lsb1 (data available at http://www.yeastgenome.org). Moreover, within the S. sensu stricto clade, only the Lsb2 proteins of S. cerevisiae and its sister species, S. paradoxus contain N residue at the position 213, while Lsb2s of more distantly related species, S. mikatae and S. bayanus bear S (Fig. 2). This indicates that the prion-inducing activity of Lsb2 is a relatively recent evolutionary acquisition. It has been reported previously that the prion forming ability can be confined to some yeast proteins by individual amino acid substitutions in these proteins. We demonstrate that the substitutions producing a prion-forming potential indeed arise and can be fixed in real phylogenetic lineages. It is worth noting that both S. cerevisiae and S. paradoxus are characterized by increased optimal growth temperatures, compared with other species of the S. sensu stricto clade. Thus, acquisition of the prion-forming ability by the Lsb2 protein coincides with the acquisition of increased thermostolerance in evolution of Saccharomyces yeast.

A switch of the S. cerevisiae / S. paradoxus branch to the increased growth temperatures should have resulted in an increased exposure to the high temperature stress conditions. As described above, prion-forming potential of Lsb2 and prion-mediated stress memory could have become beneficial for the population survival in such conditions. Therefore, it is possible that the acquisition of prion-forming capabilities by Lsb2 protein played an adaptive role in yeast evolution.

**CONCLUSIONS AND PERSPECTIVES**

Our data uncover a new mechanism of stress memory that is based on the ability of a stress-inducible protein Lsb2 to acquire a
A metastable prion state in response to stress and to maintain it for the indefinite number of cell divisions but only in a fraction of the clonal population (Fig. 3). The ability of aggregated Lsb2 to promote assembly of other aggregated proteins in the cytoskeleton-associated deposits indicates that the prion-forming potential of Lsb2 may play a protective role during a proteotoxic stress, and that prion-mediated stress memory can produce a subpopulation that is better prepared for the return of stressful conditions. Acquisition of the prion-forming potential by Lsb2 is a relatively recent evolutionary event that coincides with the adaptation of Saccharomyces yeast to higher growth temperatures. Further work will show if such a prion-based stress memory could also be mediated by other proteins. Remarkable similarities between amyloidogenic formations of yeast and higher eukaryotes suggest that prion-based stress memories could be applicable beyond yeast. Understanding general mechanisms of the environmental and physiological modulation of prion aggregation may pave the way for developing the anti-amyloid prophylactic strategies in humans.16,23,73

DISCLOSURE OF POTENTIAL CONFLICTS OF INTEREST

No potential conflicts of interest were disclosed.

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