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Luminidependens (LD) is an Arabidopsis protein with prion behavior

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Prion proteins provide a unique mode of biochemical memory through self-perpetuating changes in protein conformation and function. They have been studied in fungi and mammals, but not yet identified in plants. Using a computational model, we identified candidate prion domains (PrDs) in nearly 500 plant proteins. Plant flowering is of particular interest with respect to biological memory, because its regulation involves remembering and integrating previously experienced environmental conditions. We investigated the prion-forming capacity of three prion candidates involved in flowering using a yeast model, where prion attributes are well defined and readily tested. In yeast, prions heritably change protein functions by templating monomers into higher-order assemblies. For most yeast prions, the capacity to convert into a prion resides in a distinct prion domain. Thus, new prion-forming domains can be identified by functional complementation of a known prion domain. The prion-like domains (PrDs) of all three of the tested proteins formed higher-order oligomers. Uniquely, the Luminidependens PrD (LDPrD) fully replaced the three of the tested proteins formed higher-order oligomers. To investigate the prion-forming capacity of these proteins in living cells, we took advantage of a comprehensive suite of biochemical and genetic tools previously established in yeast. Most of the well-characterized yeast prions adopt a variety of related, but distinct, self-templating conformations, known as “prion strains.” In genetically identical yeast strains, these prion strains create a spectrum of heritable prion elements, with each exhibiting a different phenotypic strength (21). Because prion proteins in their prion conformations efficiently convert nonprion conformers into the prion state, prion traits are dominant in genetic

Significance

Prion proteins provide the best-understood mode for protein-based molecular memory. Since their discovery in mammals, prions have been identified in diverse organisms including fungi, Aplysia, and Drosophila, but not in the plant kingdom. Applying methods we used to uncover yeast prions, we identified nearly 500 Arabidopsis proteins that harbor potential prion-like domains (PrDs). At least one of these domains, Luminidependends PrD, had some of the classical characteristics of prion proteins when tested experimentally in yeast, making it, to our knowledge, the first protein from the plant kingdom with bona fide prion attributes. Importantly, Luminidependens is involved in the process of flowering, a crucial development course that integrates several internal and external cues, including memories of winter, for its regulation.


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The authors declare no conflict of interest.

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Plants also form memories, recording previous exposure to drought, heat, prolonged cold, and pathogens (14). The basis for these memories is epigenetic, and they are remarkably stable.

For instance, the memory of wintering (known as vernalization) is formed after prolonged exposure to cold and promotes flowering in the spring (15, 16). This memory can persist in a plant grown from the callus of a plant exposed to cold, even if the new plant never experiences the cold (17). The molecular mechanisms underlying these remarkable memory phenomena are not fully understood and could involve prion-like mechanisms. However, no prions have been identified in plants. Using a computational algorithm developed to find yeast prions (5, 18), we identified ~500 Arabidopsis proteins that have distinct prion-like domains (PrDs). These proteins showed a functional enrichment for involvement in flowering.

We focused further investigation on proteins belonging to the autonomous flowering pathway: Luminidependens (LD), Flowering Locus PA (FPA), and Flowering Locus CA (FCA) (19, 20). To investigate the prion-forming capacity of these proteins in living cells, we took advantage of a comprehensive suite of biochemical and genetic tools previously established in yeast. Most of the well-characterized yeast prions adopt a variety of related, but distinct, self-templating conformations, known as “prion strains.” In genetically identical yeast strains, these prion strains create a spectrum of heritable prion elements, with each exhibiting a different phenotypic strength (21). Because prion proteins in their prion conformations efficiently convert nonprion conformers into the prion state, prion traits are dominant in genetic

Prions are proteins capable of adopting profoundly different structures that possess strong functional distinction in which at least one conformation is self-perpetuating (1). Although prions were first identified as the cause of a transmissible neurodegenerative disease in mammals, of greater interest has been their function as a protein-based system of memory (1–3). In yeast, a dozen prion proteins are known (4). Most of these are regulators of transcription, translation, or RNA processing (5). Their self-propagating conformations serve as elements for the inheritance of diverse traits that are passed through the cytoplasm of mother to daughter cells (6, 7). The heritable conformations of prions impart profound phenotypic consequences, which can be beneficial, benign, or detrimental, depending on the genetic background of the strain and the environmental conditions (6).

Each prion has a unique function depending upon the biological context, with self-templating of the protein-conformational state being the unifying characteristic (8). In mammals, prion-like proteins are involved in signal transduction mechanisms in innate immunity and inflammation (9, 10). In Aplysia and Drosophila, an evolutionarily conserved prion conformation of the cytoplasmic polyadenylation element binding protein serves as a “molecular memory” for the long-lasting maintenance of neuronal synapses (11–13). The diversity of these traits and the conservation of prion formation capability from yeast to human suggests not only that prions are biologically important and frequently beneficial, but also that a world of other prions remains to be discovered.

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crosses (denoted by capital letters). They are also generally inherited cytoplasmically, segregating to most or all meiotic progeny after sporulation (22) (denoted by brackets). Thus, a prion phenotype is denoted as [PRION+], whereas the corresponding nonprion state is denoted as [prion−]. Because it is based on a protein conformation-based genetic element, prion inheritance is sensitive to changes in activity of the proteostasis machinery (23). Collectively, all of these properties distinguish prion-based traits from changes caused by genetic mutations. By testing for the presence of these properties, we show that the prion domain of the plant protein LD can function as a prion capable of forming and perpetuating molecular memory.

**Results**

**Identification and Functional Classification of Arabidopsis Prion Domains.** We previously developed a computational algorithm, PLAAC, to identify prion domains. The algorithm uses a hidden Markov model (HMM) to compare the similarity of a given sequence to the known sequence characteristics of previously identified yeast prions (18). We scored the entire Arabidopsis proteome using this algorithm and identified 474 Arabidopsis proteins that contained PrDs (Fig. 1A and Dataset S1).

Gene Ontology (GO) annotation analysis of the candidate proteins (Dataset S2) showed enrichment in proteins categorized as regulators of transcription ($P = 2.96 \times 10^{-31}$), RNA binding...
Expression of Arabidopsis Prion Domains in Yeast. We fused the PrDs of LD, FPA, FCA, and FY to GFP and expressed the fusion proteins from a galactose-inducible promoter. FY expression was not detectable, but LD, FPA, and FCA were expressed at similar, readily detectable levels (Fig. S1). All three chimeric proteins formed distinct foci, a common characteristic of prion proteins (Fig. 1B). For each protein, multiple initial foci coalesced into one or two larger foci over time (Fig. 1B). As is common for yeast prions, candidate Arabidopsis PrD (cArabPrDs) foci colocalized with Rnq1 aggregates, indicating their localization to the cytoplasmic site known as the “insoluble protein deposit” (24) (SI Results; Fig. S1B). However, the [RnQ+] prion was not required for their localization (Fig. S1C).

Many proteins form foci upon overexpression. The SDS resistance of protein complexes on semidenaturing detergent-aragrose gels (SDD-AGE) can distinguish nonspecific aggregates and cellular organelles from prion-like foci (25). Indeed, the cArabPrD-GFPs formed SDS-resistant oligomers, which were eliminated when the samples were boiled. However, unlike known prions that typically form high-molecular-weight amyloid fibrils, the oligomers formed by cArabPrD-GFPs are low molecular weight (Fig. 1C).

LDPrD Provides a Faithful, Protein-Based Phenotypic Switch. As a defining prion characteristic, alternate prion conformations must self-perpetuate, templating other proteins of the same type to the prion state. In yeast cells, this process produces a heritable phenotypic switch that is stably perpetuated across hundreds of generations. To assess whether the cArabPrDs have this characteristic, we took advantage of the modular nature of prion domains (26) and the well-characterized [PSI+] phenotype (6, 27). When the N-terminal PrD of Sup35 is replaced by the PrD of another prion, the resulting chimeras can form a heritable element that behaves like [PSI+]. Any such chimeras (PrD-Sup35C) can be readily assessed for their prion-forming ability in strains carrying an adel1-14 allele, which harbors a premature stop codon. In the nonprion state, the chimeric Sup35 protein is fully functional, causing ribosomes to stop translation at the stop codon. Such cells are unable to grow in the absence of adenine and form red colonies on complete medium (due to the accumulation of a synthetic by-product). When the chimeric protein switches to the prion state, the C-terminal termination activity is sequestered, ribosomes are read through the stop codon, and functional Ade1 is produced. Cells can then grow in the absence of adenine and produce white colonies (5, 10, 28–30).

To test this prion characteristic, cArabPrD-Sup35 chimeras were individually expressed in cells lacking endogenous Sup35. The strains harboring cLDPrD- or cFCAPrD-Sup35C did not grow in medium without adenine (Fig. 2A, Left) and produced red colonies in complete media, indicating that these chimeric Sup35 proteins were functional. However, cells harboring cFPAPrD-Sup35C grew in adenine-deficient medium and formed white colonies, indicating that most of this chimeric protein was already in a nonfunctional state. We were therefore unable to test its prion-forming ability any further.

To test the capacity of cLDPrD- and cFCAPrD-Sup35C to undergo a prion switch, we took advantage of the fact that transient overexpression of prion proteins is sufficient to induce the conformational switch (31). This switch in conformation then self-perpetuates, even when expression levels are returned to normal. Transient overexpression of cLDPrD-GFP switched the strain harboring cLDPrD-Sup35C into a prion state, as evidenced by the ability to grow on adenine-deficient medium (Fig. 2A, Right). Moreover, we observed a spectrum of prion strains with this protein, ranging from strong (white) to weak (dark pink) in complete medium (Fig. 2B). However, cFCAPrD did not provide a prion switch because it was unable to grow on adenine-deficient medium, even after overexpression (Fig. 2A, Right). Because LDPrD-Sup35C could stably exist in either a distinguishable [prion−] or a variety of [PRION+] states, we henceforth focused our efforts on characterizing the [PRION+] states provided by LDPrD ([LD+]).

In addition to the adel1-14 read-through, we confirmed that [LD+] cells had the ability to read through other premature stop codons. We introduced a GFP construct with a premature stop codon in both [LD+] and [ld−] strains. GFP expression was obtained only in the [LD+] cells. This result was assessed by fluorescence microscopy and FACS analysis (Fig. 2C).

We confirmed that the phenotype obtained in the [LD+] strain was solely dependent upon the LDPrD (SI Results; Fig. S2 A and B). Further, [LD+] was maintained with high fidelity through sequential transmissions, successive streaks on plates, and hundreds of generations of growth in liquid medium, although we observed spontaneous but rare loss of [LD+] (<5%; Fig. S2C). These data firmly established that the LDPrD could maintain a heritable self-perpetuating state.
Materials and Methods

LD+ strains have strong prion-like characteristics. We next examined the dominance of the [LD+] phenotype by analyzing matings and the cytoplasmic heritability of [LD+]. Indeed, both the diploids, formed by mating [LD+] with [ld−] cells, and the [ld−] haploids that were cytoduced with [LD+] cells grew on adenine-deficient medium and appeared white in complete medium (Fig. 3A and B). Because the diploids exhibited sporulation defects, our ability to analyze any inheritance patterns in their meiotic progeny was restricted. However, our results established that the [LD+] phenotype is indeed protein-based, fulfilling a key criterion of prion biology. Propagation of prions in fungi relies on the generation of smaller “seeds” that are generated by the disaggregate, Hsp104 (32–35). Traditional manipulations of Hsp104 that cure most prions were ineffective against [LD+]. Transient inhibition of Hsp104 activity using chemical inhibition with 5 mM GdnHCl (Fig. S3A), genetic perturbation using a dominant-negative variant of Hsp104 that interferes with the wild-type Hsp104 activity (Fig. S3B), and overexpression of Hsp104 (Fig. S3C) did not affect the [LD+] phenotype. In contrast, all of these methods eliminated the [PSI+] phenotype generated by Sup35 (Fig. S3 D–G). Although most known yeast prions are dependent on Hsp104, recently emerging evidence indicates that other molecular chaperones like Hsp70 and, potentially, Hsp90 may play Hsp104-independent roles in prion biology (9, 36, 37). Therefore, we exploited a dominant-negative Hsp70 variant (Ssa1-K69M or Hsp70(Y204)) that eliminates the yeast prion [GAR+] (37). We transiently expressed Hsp70(DN) in [LD+] strains and assayed the [LD+] phenotype after restoring Hsp70 activity (by removing the plasmid harboring Hsp70(DN)). Most [LD+] strains (65%) reverted to [ld−] strains (Fig. 4A). This phenomenon was especially common among intermediate strains.

Hsp90 has been implicated in a prion-like phenotype underlying innate immunity: Aggregates of the prion-like MAVS protein were eliminated by chemical inhibition of Hsp90 (9). We transiently inhibited Hsp90 activity by passing [LD+] and [ld−] strains in media containing either a concentration of 10 or 20 μM radicicol. In contrast to dominant-negative Hsp104 overexpression, the majority (80%) of the [LD+] strains altered their phenotype to the [ld−] state, although some of them showed this effect only at a higher dose of radicicol (Fig. 4B). These results

Fig. 3. The [LD+] state is dominant and can be inherited through the cytoplasm. (A) Dominance of the [LD+] state. (A, i) Cartoon depiction of the crosses. The α-mating type of [LD+] was crossed with the α-mating type of [ld−] and the resulting diploids were assessed for Ade-read-through in adenine-deficient medium and color phenotype in complete medium (A, ii). (B) [LD+] phenotype can be cytoduced. (B, i) Cartoon depiction of generation of cytoductants. Various modifications were made to both the α-mating type of [LD+] and the α-mating type of [ld−], as indicated in the figure and described in Materials and Methods. (B, ii) The buds from the heterokaryons were grown in different selective media to ensure the strain accuracy: [LD+] were able to grow in glycerol, whereas [ld−] could not due to defective mitochondria; only [ld−] grew on YPD+NAT because [ld−] has an antibiotic marker in the nucleus; only cytoductants grew in medium with both glycerol and NAT. The phenotypes of the cytoductants were assessed in medium lacking adenine and complete medium.

Fig. 4. The [LD+] state can be perturbed by transient inhibition of molecular chaperones. (A) Transient inhibition of Hsp70 by expression of Hsp70(DN) (Ssa1(DN)). (A, i) Cartoon depiction of transient inhibition of Hsp70. (A, ii) Sup35C assay. Transient inhibition of Hsp70 affected some [LD+] strains because they no longer grew in medium lacking adenine and appeared red in complete medium. (B) Transient inhibition of Hsp90 by radicicol. (B, i) Cartoon depiction of transient inhibition of Hsp90. (B, ii) Sup35C assay. Transient inhibition of Hsp90 using 10 and 20 μM radicicol affected most [LD+] strains because they no longer grew in medium lacking adenine and appeared red in complete medium.
raise the possibility that the inheritance of [LD+] may be blocked by transient inhibition of Hsp70 or Hsp90.

**Discussion**

Here, to our knowledge, we report the first evidence that at least one plant protein has the capacity to serve as a prion. The LD protein, which functions in the autonomous flowering pathway, contains a domain that is similar in sequence to the prion domains of yeast proteins and can fully substitute for prion function in yeast. The LD prion domain enters into a higher-order, self-perpetuating structure that can change the function of an associated protein domain in a stable and heritable way. That is, it is capable of forming protein-based molecular memories.

Plants are known to possess the capacity to form memories in response to diverse environmental factors, including changes in temperature and day length (38). One such memory is vernalization, which develops upon exposure to prolonged cold and is the mechanism by which a plant recognizes and remembers the passing winter (15, 16). Epigenetic changes in response to this exposure prime the plant for flowering in the spring (39). The phenomenon is, in a sense, similar to long-term potentiation of a neuronal synapse, which has recently been shown to involve the prion-like assembly of RNA-binding proteins (12, 30, 40).

To identify potential Arabidopsis prions, which might be involved in some forms of plant memory, we scanned the proteome using a computer algorithm trained on known yeast prions. Using yeast cells as “living test tubes” for protein folding in a eukaryotic cytoplasm, we showed that three domains from Arabidopsis proteins (LD, FPA, and FCA) had some prion-like properties. All of these proteins are in the autonomous flowering pathway (39), which participates in flowering by responding to yet-unknown internal cues. All three aggregated and formed SDS-resistant assemblies. However, in our test system, only the candidate prion domain of LD could stably and heritably adopt either a soluble conformation or a higher-order prion conformation. Indeed, as is the case for most yeast prion proteins, the LD PrD formed a variety of self-templating conformations with distinguishable phenotypic consequences.

Unlike classical yeast prions, [LD+] was not curable by Hsp104. Instead, our results suggest that Hsp70 and Hsp90 may play a role in maintaining this prion. Also, instead of high-molecular-weight amyloids, the LD-PrD-GFP is similar in size to low-molecular-weight oligomers. We have previously shown that at least one fungal prion, [GAR+], does not form amyloid or require Hsp104; instead, this prion is Hsp70-dependent (36, 37). Therefore, it is possible that [LD+] belongs to this noncanonical class of prions.

In Arabidopsis, prolonged exposure to cold causes heritable chromatin modifications, including histone methylation and silencing of FLC, a key repressor of flowering (41–45). But how, at a molecular level, does a plant differentiate between a single cold night and winter? Stable, self-perpetuating changes in the conformational states of certain proteins, such as those studied here, might be involved, LD, FCA, and FPA are predicted to bind nucleic acids. FCA and FPA both contain RNA-recognition motifs (46, 47) and are known to play important roles in RNA 3′ processing and transcription termination (48). LD contains a homedomain that can bind DNA and regulate transcription (20). Because all three of these proteins are able to form SDS-resistant assemblies, altered activities in their assembled forms could influence the chromatin modifications that are known to be involved in flowering decisions.

Whether the LD protein undergoes a biologically significant prion-like conformational change that might play a role in the flowering decision in plants remains to be seen. However, roles for self-perpetuating prion conformations in chromatin remodeling are highly likely. Indeed, at minimum, one yeast chromatin-remodeling factor, the Swi subunit of the SWI/SNF chromatin-remodeling complex, can form a prion that epigenetically alters its chromatin remodeling functions and thereby imparts heritable changes in the biological state of the organism (49). The role that prion-like conformational changes might play in plant biology and, in particular in the many forms of plant memory, is currently unknown but highly intriguing.

On another note, it is interesting to reflect on the fact that in the mid-20th century, some Soviet scientists used the plant memory phenomenon of vernalization as a strong argument against the “gene-based” concept of inheritance (50). Given this fact, it is ironic that the concept of vernalization may, indeed, have a heritable non-DNA element involved in its perpetuation. Of course, the capacity for proteins to stably switch states in a somatically heritable way is itself dictated by DNA sequences that encode proteins capable of such unusual, autonomous, self-perpetuating biological switches. Our work suggests that much broader roles for prions exist and that this is a subject ripe for further exploration.

**Materials and Methods**

**Computational Detection of Arabidopsis Prion Candidates.** Each Arabidopsis protein was searched for the presence of PrDs by using a HMM that parsed each protein into prion-like and non-prion-like (background) regions. Any protein containing a PrD of at least 60 consecutive prion-like amino acid residues received a nonzero COREscore (Dataset S1). This COREscore is the sum of a new individual log-likelihood ratios for each amino acid residue contained within this domain. The COREscores of the candidate 474 Arabidopsis PrDs are presented in Dataset S1.

**GO Annotation Analysis.** We performed GO annotation analysis using the Database for Annotation, Visualization and Integrated Discovery (DAVID) Bioinformatics Resource 6.7 platform. We used Arabidopsis proteome as the background for functional category enrichment (Dataset S2).

** Yeast Strains, Vectors, and Antibodies.** Yeast strains used in this study were YMJ509 and YMJ584 (5). They were grown on standard synthetic medium lacking standard amino acids/bases and containing either d-glucose (SD) or d-galactose as carbon source. The PrDs from LD, FPA, FCA, and FY were synthesized from GeneArt (Life Technologies). These domains were then used to generate yeast entry clones by using the pZz21 donor plasmid (Addgene).

The above entry clones were used to generate various destination plasmids by using 424GAL1cubBFpG and 415AD1H1cubSup35C (51): 424GALArbPrD-GFP (ArabPrD-GFP) and 415ADH1ArabPrD-SUP35C (ArabPrD-SUP35C). Expression of carA GFP-GFP was controlled by using 2% (wt/vol) galactose for induction.

** GO Annotation Analysis.** We performed GO annotation analysis using the Database for Annotation, Visualization and Integrated Discovery (DAVID) Bioinformatics Resource 6.7 platform. We used Arabidopsis proteome as the background for functional category enrichment (Dataset S2).

**Cell Cytometry Analysis.** Yeast cultures were grown in synthetic dropout medium to log phase and then transferred to a 96-well plate for analysis in an EasyCyte flow cytometer (Guava Technologies). GFP fluorescence was monitored using the green channel. High and low GFP gates were set such that > 95% of [LD+] and [LD−] cells were included in the low-GFP gate. Percentage of cells in these gates were compared for all strains.

**Read-Through Assays.** adet-14 read-through was assessed as detailed in ref. 5. Briefly, the ArabPrD-Sup35C chimeras were individually expressed in the YRS100 strain (a derivative of YMJ584), where the endogenous Sup35 protein has been deleted, but the activity of this essential protein is covered by 416GPDSup35C. Then a plasmid shuffle was performed by plating the cells in 5-fluoroorotic acid (5-FOA). Therefore, the cArabPrD-Sup35C (LD) was the only source of Sup35 activity in these cells.

To assess GFP read-through, [LD+] and [LD−] strains were transformed with a fusion protein containing a GFP marker preceded by a stop codon (S2).
Generating α-Mating Type Strains. Yeast have two mating types: MATa and MATα. Only crosses from opposite mating type yield diploids. To mate [L−]+ strain with [L−] strain, we switched the mating type of [L−] strain from α-type to χ-type. To do so, we transiently overexpressed a Homothallic switching endonuclease (HO) gene (on a Ura3 marker) for 2-4 h. The HO plasmid was counterselected by plating single colonies on 5-FOA. α-type colonies were then identified by crossing with mating type tester strains (DS88a or MB4u).

Generating Strains for Cytoduction. The [L−] strain was made kar1 (karyogamy) mutant, so that it was unable to undergo karyogamy during mating. Next, we introduced an antibiotic-resistant (NAT+, resistant to nourseothricin) nuclear marker in the [L−] strain by disrupting the nonessential HO locus. This strain was then converted into respiration-deficient petite strain (−), so that it was unable to grow in medium containing nonfermentable carbon sources, like glycerol (Fig. 3B). Heterokaryons were established by crossing corresponding “donor” [L−] and the “recipient” [L−] strains. Cytoductants were selected by growing the buds from the heterokaryons in medium containing nourseothricin and where glycerol was the only carbon source.

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Supporting Information

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SI Results

Expression of cArabPrDs in Yeast. We achieved maximal expression of the cArabPrD-GFPs after 7 h of induction, as demonstrated by both fluorescence microscopy and immunoblot analysis (Fig. 1B and Fig. S1A). We used this time point for further experimentation.

GO Annotation Analysis. Like known yeast prions, proteins categorized as regulating transcriptional ($P = 2.97 \times 10^{-31}$) or RNA metabolic processes ($P = 3.72 \times 10^{-12}$) were highly enriched (Dataset S2). This result suggests that the power of prion-like proteins affecting the phenotype in the absence of genotype alterations may be global, extending this phenomenon even to the plant kingdom.

Foci Formation Is Not Dependent on [RNQ+]. The amyloid assemblies of some yeast prions, like [PSI+], require inducibility factors to acquire these conformations. The protein Rnq1 acts as an inducibility factor in its prion form, [RNQ+]. For example, overexpression of the Sup35 PrD, NM, does not induce foci in a [rnq−] background (Fig. S1C). However, cArabPrD-GFPs formed foci even in the absence of [RNQ+], distinguishing them from yeast prions like [PSI+]. Therefore, although the cArabPrD-GFPs colocalize with the Rnq1-CFP, they are not dependent on Rnq1 for formation of oligomeric species.

LDPrD Is Responsible for the Protein-Based Phenotypic Switch. We eliminated the remote possibility that the [LD+] phenotype resulted from degradation of the chimeric protein by immunoblot analysis. The LDPrD-Sup35C levels were very similar in both [LD+] and [ld−] cells (Fig. S2A). We also confirmed that the phenotype we obtained was indeed solely due to LDPrD by swapping the LDPrD-Sup35C with an untagged-Sup35C in the [LD+] strain. Without the LDPrD domain, the cells instantly reverted back to a [prion−] state (Fig. S2B).
Fig. S1. Characterization of Arabidopsis prion domains in S. cerevisiae. (A) Immunoblot analysis of cArabPrD-GFPs. The constructs were expressed for various time points and analyzed by α-GFP. α-PGK1 was used as the loading control. (B) Some cArabPrD-GFP foci localize in the insoluble protein deposit (IPOD). Fluorescence microscopy of yeast cells with the cArabPrD-GFP (green) and Rnq1-CFP (red) after 7 h of coexpression is shown. The colocalized aggregates are marked with arrows. (C) Foci formation is not dependent on [RNQ+]. Fluorescence microscopy images of cArabPrD-GFPs in a [rnq−] strain background are shown. NM-YFP is shown as a control.
Fig. S2. Phenotypic switch into the [PRION+] state. (A) Levels of the LDPrD-Sup35C protein are the same in [LD+] and [ld−] cells. The soluble protein levels of LDPrD-Sup35C were similar in both strains as assessed by α-Sup35C antibody. α-PGK1 antibody was used as loading control. (B) The [LD+] phenotype is dependent on the LDPrD. (B, i) Cartoon representation. A reverse plasmid shuffle was performed by which the untagged GPD-Sup35C was transformed into the [LD+] strains and the LDPrD-Sup35C plasmid was removed. This led to faithful translation termination due to the availability of soluble Sup35C protein. (B, ii) The [LD+] strain reverts back into a [psi−]-state, and therefore there is no growth in medium lacking adenine and cells appear red in complete medium compared with the [LD+] strain. (B, iii) The strains were checked for accuracy by growing on respective selective medium, SD-Ura for the presence of GPD-Sup35C and SD-Leu for LDPrD-Sup35C. (C) The [LD+] phenotype is lost randomly at a very low frequency. This result was apparent from the appearance of red color colonies as marked by the arrows among [LD+] cells.
Fig. S3. Perturbations in Hsp104 levels do not affect [LD+] phenotype. (A, i) Cartoon depiction of transient inhibition of Hsp104 activity chemically (5 mM GdnHCl) or genetically (Hsp104 dominant negative, Hsp104\(^{DN}\), or Hsp104 overexpression). (A, ii, B, and C) Sup35C assay. Transient inhibition of Hsp104 activity chemically or genetically did not affect the [LD+] phenotype because the cells appeared white in complete medium. In all cases, the [PSI+] phenotype reverted to a [psi−] phenotype.

Dataset S1. By using the PLAAC algorithm, the entire Arabidopsis proteome was analyzed for prion domains. Sequence ID for each protein with prion domain is provided. The proteins are ranked according to COREscore. Prion-domain sequence is also provided.

Dataset S1

Dataset S2. GO annotation analysis was performed for the proteins with candidate prion domains