Abnormal calcium homeostasis and protein folding stress at the ER
A common factor in familial and infectious prion disorders

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Prion-related disorders (PrDs) are caused by the accumulation of a misfolded and protease-resistant form of the cellular prion, leading to neuronal dysfunction and massive neuronal loss. In humans, PrDs have distinct etiologies including sporadic, infectious and familial forms, which present common clinical features; however, the possible existence of common neuropathogenic events are not known. Several studies suggest that alterations in protein folding and quality control mechanisms at the endoplasmic reticulum (ER) are a common factor involved in PrDs. However, the mechanism underlying ER dysfunction in PrDs remains unknown. We have recently reported that alterations in ER calcium homeostasis are common pathological events observed in both infectious and familial PrD models. Perturbation in calcium homeostasis directly correlated with the occurrence of ER stress and higher susceptibility to protein folding stress. We envision a model where alterations in ER function are central and common events underlying prion pathogenesis, leading to general alterations on protein homeostasis networks.

The presence of abnormal protein inclusion in the brain is a common pathologic feature of many neurodegenerative diseases, including Alzheimer’s disease (AD), Parkinson’s disease, Huntington’s disease, Amyotrophic lateral sclerosis and Prion-related Disorders (PrDs). PrDs, also known as transmissible spongiform encephalopathies, are fatal neurodegenerative diseases affecting humans and other mammals, characterized by a spongiform degeneration of the brain and progressive neuronal loss. At the biochemical level, PrDs are associated with the accumulation and deposition of the abnormal and misfolded form of the cellular prion protein (PrPc), termed PrPRES (for protease resistant form). Depending on its etiology, PrDs can be divided in three main forms, familial, sporadic and infectious. Familial PrDs including Creutzfeldt-Jackob disease (CJD), fatal familial insomnia (FFI) and Gerstmann-Sträussler-Scheinker syndrome (GSS), are all linked to mutations in the gene encoding PrPc, PRNP and represent approximately 10% of total cases. In humans, the infectious form is very rare and accounts for less than 1% of total cases, highlighting the new variant Creutzfeldt-Jackob disease (vCJD). The sporadic CJD underlay around 90% of total PrD cases in humans.2,3 The mechanism triggering the misfolding of PrPc in infectious forms is very unusual. The most accepted “protein-only” hypothesis postulates that infectious prion pathogenicity results from a direct interaction between the infectious/misfolded PrP form, termed PrPRES, and the native folded and fully matured PrPc. This interaction is predicted to induce a conformational change of its primarily α-helical structure to an insoluble β-sheet conformation, a process that occurs cyclically on an autocatalytic manner as we and others have demonstrated.4,5 In PrD familial forms, PrP misfolding occurs due to direct mutations in the PRNP gene. The factors involved in the misfolding of PrP in sporadic cases are still unknown. It is possible that environmental factors, together with specific genetic backgrounds (mutation of genes different of PRNP), or undetected somatic mutations may increase the susceptibility to generate de novo PrPRES in sporadic CJD.

PrP is a membrane attached protein that undergoes post-translational processing in the endoplasmic reticulum (ER) and Golgi, including the addition of a glycosyl phosphatidyl inositol (GPI) anchor, C- and N-terminal proteolytic processing, formation of one disulphide bond, the addition of N-linked glycosylations.5 After trafficking through the secretory pathway, fully matured PrPc localizes in the outer surface of the plasma membrane in cholesterol-rich lipid domains (lipids rafts), and cycle through the endocytic pathway with a half live of ~6 hours.5 During the folding process at the ER, around 10% of PrPc is naturally misfolded and eliminated through the ER-associated degradation (ERAD) pathway by proteasomes.6 In the case of the familial forms of PrDs, some PrP mutants are retained and aggregated in the ER and Golgi, where they may exert their pathological effects.7 However, the molecular mechanisms explaining the neurotoxicity of PrP mutants and the factors regulating this process are still elusive. In contrast to PrP mutants linked to familial PrDs, the generation of misfolded PrP in infectious forms of the disease is proposed to occur at the plasma membrane and during its cycling through the endocytic pathway.7,8 However, many studies in infectious PrDs models also have shown the trafficking...
and accumulation of PrPRES at the cytosol and ER of the infected cells.9,12

**Altered ER Homeostasis in Prion-related Disorders**

Different stress condition can affect the normal ER homeostasis, interfering with the correct folding of proteins in the ER lumen. This condition, termed “ER stress,” engages a complex integrated signaling cascade known as the unfolded protein response (UPR).13 The UPR aims reestablishing homeostasis by recovering the capacity of the cell to synthesize properly folded proteins at the ER.

Several groups have observed the occurrence of ER stress responses in PrDs including the activation of the UPR transcription factor XBP-1 splicing14 and the action of JNK and ERK.14,15 In human CJD patients and mouse models, the upregulation of several chaperons and foldases such as such as Grp78/BiP, Grp94 and Grp58/ERp57 proteins is observed, suggesting abnormal ER homeostasis.14,20 The levels of ERp57 increase even during the asymptomatic stage of the disease, and it is detected in high levels in brain regions that are more resistant to undergo degeneration in scrapie prion-infected mice.18 Additionally, in studies using Neuro2a neuroblastoma cells we demonstrated that ERp57 operates as a neuroprotective factor against infectious PrPRES neurotoxicity.18 Interestingly, ERp57 co-immunoprecipitates with PrP.18 In addition to these results, a proteomic analysis of post-mortem brain samples from patients affected with sporadic CJD demonstrated that ERp57 is highly expressed in the pathology.20 A recent report suggests that the expression of a GSS-linked PrP mutant triggers ER stress in a cellular model.21

Perturbations of ER homeostasis leads to generation of intermediary misfolded forms of PrPC, increasing its susceptibility to be converted into PrPRES in vitro.22,24 This partial misfolding of PrPC, may be reverted by the overexpression of UPR components such as XBP-1, ATF4 and ATF6,23 suggesting that the UPR has an active role in preventing neurodegeneration. In cellular models, ER stress conditions reduce PrP co-translational translocation, favoring accumulation of aggregation-prone cytosolic species, which retain the signal sequence but lack N-glycans and disulfides.24 Inhibition of proteasomes further increases the levels of cytosolic PrP.24 Overexpression of UPR transcription factor XBP1 facilitated ER translocation, suggesting possible neuroprotective effects of this pathway.24 Similarly, other groups have shown that cytosolic accumulation of PrP lead to accumulation

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**Figure 1.** ER stress and altered calcium homeostasis in Prion-related Disorders. Misfolded mutant PrP associated with inherited forms of PrDs accumulate in the ER and may affect the activity of proteins controlling the influx/efflux of calcium (i.e., SERCA, IP3Rs and/or RYRs), decreasing in the long term ER calcium content. This event may alter the activity of several ER chaperones involved in protein folding and quality control mechanisms, generating ER stress. Chronic ER stress will induce sustained activation of the unfolded protein response (UPR), leading to neuronal apoptosis. In contrast, a fraction of the infectious PrP form may be transported from the plasma membrane/endosomes to ER, where it may interact with yet unknown receptors inducing an early and sustained calcium release to the cytoplasm sensitizing cells to mitochondrial-mediated apoptosis.
Abnormal Calcium Homeostasis: A Common Factor in the Pathogenesis of Prions

There are many different cellular perturbations that can generate ER stress including expression of mutant proteins, altered protein maturation, abnormal redox status, chaperone inactivation and inhibition of the proteasome or the ERAD pathway. In addition, decreased ER calcium content is an important factor leading to ER stress. In neurons, calcium signals play a relevant role as a second messenger, controlling synaptic functions and cell viability. At basal conditions, ER calcium concentrations are close to 300 μM, whereas cytosolic calcium concentrations usually are 5–50 nM. Several key ER chaperones require optimal calcium concentrations for their protein folding activity. Thus, ER-calcium depletion may inhibit the folding and maturation proteins. Besides, sustained increase of cytosolic calcium may also induce mitochondrial-mediated apoptosis via the activation of proteins like calcineurin or by overload of mitochondrial calcium (Fig. 1). For this reason, the regulation of specific concentration of calcium in the cytosol and ER lumen are critical for maintain normal cellular functions and protein homeostasis.

It has been suggested that synthetic peptides derived from PrP primary sequence may affect calcium homeostasis. However, these peptides as such have not been observed in PrDs in vivo. Recently, we described the contribution of calcium to the pathogenesis of infectious and familial PrPd forms. Using purified PrPRES from the brain of scrapie-infected mice we evaluated its impact on ER stress responses and calcium homeostasis. We found that acute exposition of PrPRES of Neuro2a cells induces release of ER-calcium and ER stress, associated with the upregulation of several chaperones and foldases including ERP57, BiP and Grp94. Consistent with these results, cells chronically infected with scrapie prions presented altered ER calcium content. Scrapie infected cells were more susceptible to undergo ER stress-mediated cell death, associated with a stronger UPR activation after exposition of these cells to ER stress-inducing agents. We then examined the possible contribution of calcium abnormalities in models of familial PrDs. We generated neuronal cells expressing PrP mutants related to FFI and familial CJD, in addition to a mutant form that is suggested to operate as a misfolded intermediate. Cells expressing mutant PrP were more susceptible to ER stress-inducing agents and presented abnormal ER calcium content.

It remains to be determined what is the exact mechanism underlying disturbed ER calcium homeostasis and ER stress in PrDs. Interestingly, similar to PrP mutants we observed that a fraction of infectious PrPRES was also locate to the ER. One interesting model to test is the possibility that PrP, may interact with proteins that control the levels of calcium like the calcium ATPase SERCA, or the calcium channels IP3 receptors and/or ryanodine receptors as suggested for example in Huntington’s disease models. Under chronic conditions, this release of calcium may occur slowly, generating in the long term a decrease in the ER steady state calcium levels, generating ER stress. Both, reduction of calcium inside of ER and increase in the cytosol could lead to drastic alterations in protein homeostasis, sensitizing cells to cell death (Fig. 1). In this scenario, the design of small molecules to target calcium handling proteins may represent a novel therapeutic strategy against PrDs and other protein mis-folding disorders.

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