

Fine-Tuning of the Unfolded Protein Response: Assembling the IRE1 α Interactome

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Endoplasmic reticulum (ER) stress is a hallmark feature of secretory cells and many diseases, including cancer, neurodegeneration, and diabetes. Adaptation to protein-folding stress is mediated by the activation of an integrated signal transduction pathway known as the unfolded protein response (UPR). The UPR signals through three distinct stress sensors located at the ER membrane—IRE1 α , ATF6, and PERK. Although PERK and IRE1 α share functionally similar ER-luminal sensing domains and both are simultaneously activated in cellular paradigms of ER stress *in vitro*, they are selectively engaged *in vivo* by the physiological stress of unfolded proteins. The differences in terms of tissue-specific regulation of the UPR may be explained by the formation of distinct regulatory protein complexes. This concept is supported by the recent identification of adaptor and modulator proteins that directly interact with IRE1 α . In this Review, we discuss recent evidence supporting a model where IRE1 α signaling emerges as a highly regulated process, controlled by the formation of a dynamic scaffold onto which many regulatory components assemble.

Introduction

A number of conditions interfere with oxidative protein-folding processes in the endoplasmic reticulum (ER) lumen (Ron and Walter, 2007), leading to accumulation of misfolded proteins, a cellular condition referred to as “ER stress.” Adaptation to ER stress is mediated by engagement of the unfolded protein response (UPR), an integrated signal transduction pathway that transmits information about protein-folding status in the ER lumen to the nucleus to increase protein folding capacity. Conversely, cells undergo apoptosis if these mechanisms of adaptation and survival are insufficient to handle the unfolded protein load.

The occurrence of ER stress is observed in many physiological processes, especially in highly secretory cells such as plasma B lymphocytes, salivary glands, and pancreatic β cells. The high demand for efficient protein folding and secretion processes in these cells constitutes a constant source of stress initiated by the presence of large amounts of misfolded proteins that are normally generated during the protein maturation process. These folding subproducts are eliminated through ER-associated degradation (ERAD), where misfolded proteins translocate to the cytosol and are degraded by the proteasome (reviewed in Vembar and Brodsky, 2008). ER stress is also triggered by conditions that alter proteostasis associated with perturbations in protein maturation, expression of certain mutant proteins, decreased chaperone function, abnormal ER calcium content redox metabolism, altered trafficking, and many others (Powers et al., 2009). As an initial response to ER stress, cells activate the UPR to decrease the unfolded protein load and recover homeostasis. In doing so, UPR signaling enforces global changes in expression of proteins related to nearly every aspect of the

secretory pathway. For example, gene-expression profiling has demonstrated that the UPR regulates genes involved in protein entry into the ER, folding, glycosylation, ERAD, protein quality control, redox metabolism, autophagy, lipid biogenesis, and vesicular trafficking (Figure 1A). Increasing attention has been given to the regulation of the UPR based on substantial evidence for the involvement of chronic ER stress in many diseases, including neurodegenerative conditions (Matus et al., 2008), cancer (Moenner et al., 2007), diabetes (Lipson et al., 2006), and inflammation (Todd et al., 2008), hence offering new therapeutic targets to treat these diseases.

Distinct UPR Signaling Branches

The UPR was first characterized in yeast where a single signaling pathway governs the response to ER stress mediated by a type I transmembrane ER protein known as IRE1p (*inositol-requiring transmembrane kinase/endonuclease*) (Ron and Walter, 2007). In higher eukaryotes, the UPR gained complexity as it is mediated by at least three distinct UPR signaling pathways initiated by the sensors IRE1 α and IRE1 β , PERK (*PKR-like ER kinase*), and ATF6 α and ATF6 β (*activating transcription factor 6*) (Figure 1A). Activated PERK inhibits protein translation into the ER through the inactivation of the initiation factor eIF2 α , alleviating ER stress by decreasing the overload of misfolded proteins. Phosphorylation of eIF2 α allows the expression of ATF4 (*activating transcription factor 4*), a transcription factor that upregulates UPR genes that function in amino acid and redox metabolism (Harding et al., 2003), including *chop/gadd153* and *gadd34* (Figure 1A). A second UPR pathway is initiated by ATF6 α and ATF6 β , type II ER transmembrane proteins whose cytosolic domain encodes a bZIP transcriptional factor

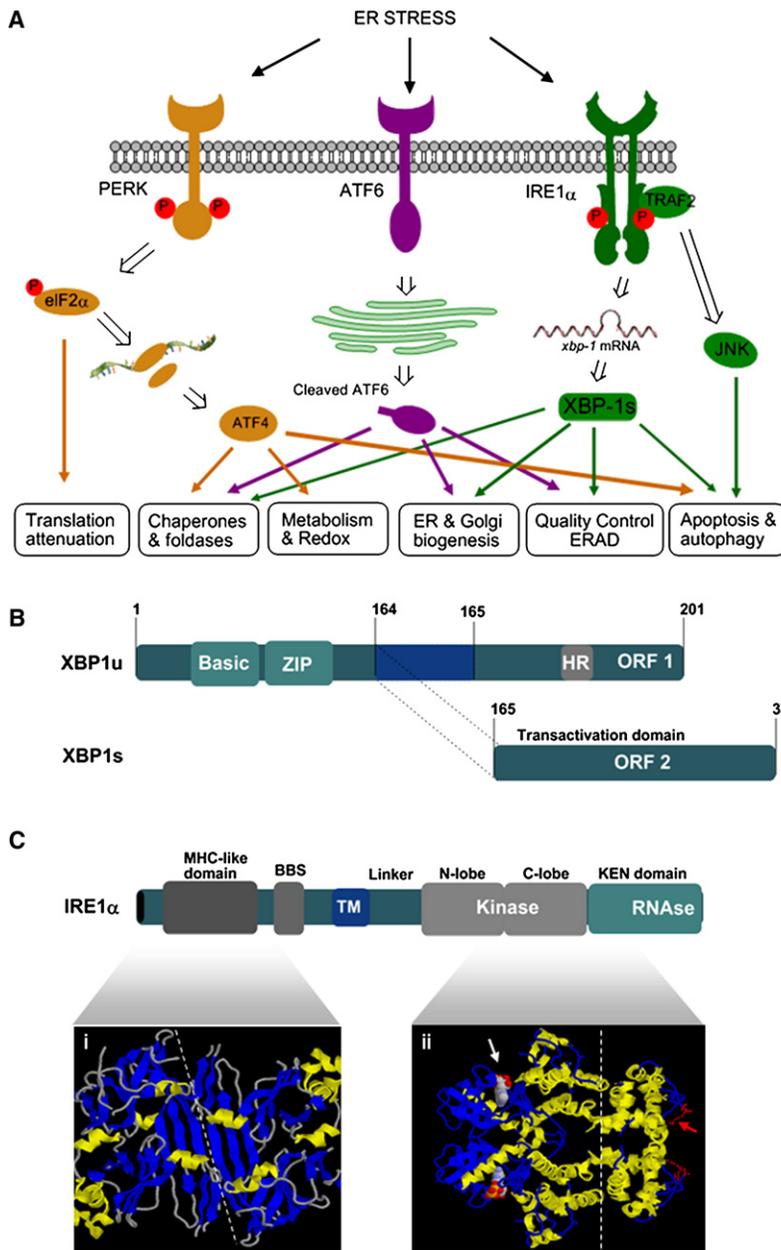


Figure 1. Essential Components of the Unfolded Protein Response

(A) UPR signaling. Accumulation of misfolded protein inside the endoplasmic reticulum (ER) lumen triggers a stress response known as UPR. There are at least three main stress sensors at the ER membrane: IRE1 α , PERK, and ATF6. In cells undergoing ER stress, IRE1 α autophosphorylates, leading to the activation of its endoribonuclease domain. This activity mediates the processing of the mRNA encoding XBP1, which is a transcriptional factor that upregulates many essential UPR genes involved in folding and protein quality control and regulates ER/Golgi biogenesis. Active IRE1 α binds the adaptor protein TRAF2, triggering JNK activation, which may participate in the regulation of autophagy and apoptosis. Alternatively, activated PERK phosphorylates and inhibits translation initiator factor eIF2 α , decreasing the synthesis of proteins and the overload of misfolded proteins at the ER. In addition, this event leads to the specific translation of ATF4, a transcription factor that induces the expression of genes that function in amino acid metabolism, the antioxidant response, and apoptosis regulators including CHOP. A third UPR pathway is initiated by ATF6, a type II ER transmembrane protein encoding a bZIP transcriptional factor in its cytosolic domain and localized in the ER in unstressed cells. Upon ER stress induction, ATF6 is processed, increasing the expression of some ER chaperones, and ERAD-related genes. At the bottom, the cellular functions affected by each UPR-signaling branch are indicated.

(B) XBP1 splicing. Schematic representation of the unspliced and spliced forms of XBP1 (XBP1u and XBP1s, respectively). Numbers indicate amino acid positions with the initiation methionine set at 1. ORF1 and ORF2 for the C-terminal domain as well as the basic and leucine zipper (ZIP) domains are indicated. Putative hydrophobic region (HR) of XBP-1u related to targeting is also indicated. (C) IRE1 structure. A schematic representation of the primary structure of IRE1p is presented indicating the kinase and RNase domains. BiP-binding site (BBS), the MHC-like domain, linker region, transmembrane region, and kinase and RNase domains are indicated. In the bottom panel, the crystal structure of the ER luminal domain groove (MHC-I like structure) of yeast IRE1p is shown (Credle et al., 2005). The dimer interface is indicated with a white line. This structural domain of IRE1p is proposed to bind misfolded proteins to stabilize the oligomeric conformation. In addition, the three-dimensional structure of the cytosolic domain of IRE1p is presented highlighting the two lobes of the kinase domain (Lee et al., 2008b). The ADP and the kinase domain are indicated with a white arrow. The KEN domain containing the RNase activity is also shown where a red arrow indicates the putative RNase active site (Lee et al., 2008b).

(Ron and Walter, 2007). Upon ER stress induction, ATF6 is processed at the Golgi, releasing its cytoplasmic domain, which acts as a transcriptional activator controlling many UPR genes related to ERAD and folding at the ER among others (Yamamoto et al., 2007) (Figure 1). This branch of the UPR is very complex and is formed by a series of newly identified ATF6 homologs that are modulated by ER stress in specific tissues, including OASIS, CREBH, LUMAN/CREB3, CREB4, and BBF2H7 (Ron and Walter, 2007). All of these ATF6-related bZip factors are processed at the Golgi in a similar way as ATF6, but their function in the UPR is poorly characterized. Mori's group recently generated ATF6 α - and ATF6 β -single knockout mice, which developed normally (Yamamoto et al., 2007). However, double-knockout mice are embryonic lethal, similar to the phenotype of X-Box-

binding protein 1 (XBP1) (Reimold et al., 2000) or IRE1 α (Urano et al., 2000) -deficient mice. In contrast, PERK-, ATF4-, and CHOP-deficient animals are viable and have varied defects in pancreatic function, metabolism, and skeletal development (Zhang et al., 2002; Harding et al., 2001; Tanaka et al., 1998; Zinszner et al., 1998).

IRE1 α is the most evolutionarily conserved branch of the UPR. Nevertheless, little is known about the regulation of IRE1 α activity. IRE1 α is a Ser/Thr protein kinase and endoribonuclease that, upon activation, initiates the unconventional splicing of the mRNA encoding the transcriptional factor XBP1 (Ron and Walter, 2007) (Figures 1B and 1C). In mammalian cells, a 26 nucleotide intron of *xbp1* mRNA is spliced out by activated IRE1 α , leading to a shift in the codon reading frame. Translation of the

new reading frame results in the conversion of XBP1 from an unspliced form of 267 amino acids to a spliced form of 371 amino acids that comprises the original N-terminal DNA-binding domain plus an additional, potent transactivation domain in the C terminus (Figure 1B).

Spliced XBP1 (XBP1s) controls the upregulation of a broad spectrum of UPR-related genes involved in protein folding, protein entry to the ER, ERAD, and protein quality control (Lee et al., 2003b; Shaffer et al., 2004). A regulatory circuitry governed by XBP1 was interrogated by a genome-wide profiling approach. In addition to classical UPR-related genes, unexpected cell-type-specific targets were identified that are linked to cell differentiation, signaling, and DNA-damage pathways (Acosta-Alvear et al., 2007). A recent genetic screen systematically characterized the functional interdependencies between UPR-target genes in yeast. These factors included chaperones, glycosylation enzymes, and ERAD components as well as trafficking pathways, transcriptional regulatory networks, modulators of lipid and ion composition, and vacuolar function (Jonikas et al., 2009). The complexity of activities/processes described in this work support the concept that proteostasis emerges from the dynamic interplay between synthesis/folding, degradation, and export processes. In addition, XBP1s regulates the expansion of the secretory pathway by controlling phospholipid biosynthesis and ER/Golgi biogenesis (Shaffer et al., 2004; Sriburi et al., 2004). Interestingly, XBP1 heterodimerizes with ATF6 α for the induction of ER-associated degradation components (Yamamoto et al., 2007; Wu et al., 2007) and ATF6 α may also modulate lipid biosynthesis and ER expansion under stress conditions (Bommiasamy et al., 2009).

Mechanism of ER Stress Sensing by IRE1

Role of the IRE1 ER Luminal Domain in Its Activation

It was originally proposed that, under normal conditions, the ER chaperone BiP/Grp78 binds to IRE1 α or the yeast homolog IRE1p maintaining the protein in an inactive monomeric state (Bertolotti et al., 2000; Kimata et al., 2003). In ER-stressed cells, BiP is released, allowing IRE1 α to multimerize and autophosphorylate its cytosolic domain. This phosphorylation event triggers the activation of the RNase activity, initiating XBP1 mRNA splicing and UPR responses. The functional impact of BiP association to IRE1p was addressed by mutagenesis analysis, observing that disruption of this interaction does not drastically alter the ability of IRE1p to detect protein misfolding (Kimata et al., 2004). Recently, new insights into the mechanism of IRE1 α /IRE1p activation have emerged from two groups who independently solved the structure of the ER luminal domain of yeast and human IRE1 protein. Peter Walter's group speculated that misfolded proteins may directly bind to the N-terminal region of IRE1p, facilitating its oligomerization through a binding motif similar to an MHC-like groove (Figure 1C), and mutations in amino acids present in the groove or in the dimerization interface abrogated the ability of IRE1p to engage the UPR (see comparison with MHC structure in Credle et al. [2005]). Thus, misfolded proteins may be directly recognized by yeast IRE1p. The general structure of the ER stress-sensing domain of IRE1p is conserved in mammals (Zhou et al., 2006). Recent *in vitro* studies consolidated both models for yeast IRE1p activation, suggesting that

BiP first dissociates from IRE1p leading to its dimerization and cluster formation (Figure 2A, see below). In the second step, direct interaction of unfolded proteins with the stress-sensing domain may orient the protein into an active IRE1p signaling cluster with full ribonuclease activity (Kimata et al., 2007).

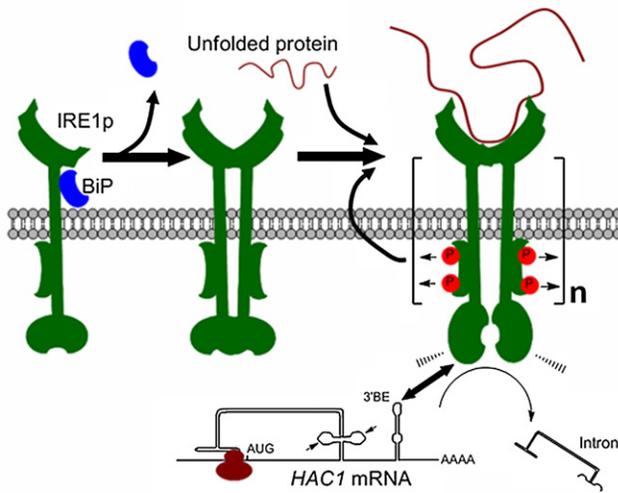
Interestingly, a recent study described the generation of luminal-domain mutants of mammalian IRE1 α that have low affinity for BiP, that gain significant activation even under unstressed conditions (Oikawa et al., 2009). Moreover, the luminal fragments of mammalian IRE1 α did not interact with unfolded proteins in an *in vitro* assay (Oikawa et al., 2009) as was described previously for yeast IRE1p (Kimata et al., 2007). These data suggested that, in contrast to yeast IRE1p, the regulation of mammalian IRE1 α may actually depend on the dissociation of BiP and may be independent of misfolded protein binding (Figure 2B) (Oikawa et al., 2009). This idea correlates well with the prediction that the MHC-like groove observed in the human IRE1 α ER luminal domain may not be able to accommodate an unfolded protein peptide as indicated in the crystal structure (Zhou et al., 2006). In this study it was proposed that IRE1 α undergoes different stages of phosphorylation, where dimerization of the ER luminal region is essential to get fully phosphorylated IRE1 α and subsequent RNase activation (Zhou et al., 2006). A similar dimer interface for the PERK-luminal region was also predicted.

The mechanisms involved in the activation of PERK and ATF6 and how these receptors sense the unfolded protein load, have not been directly investigated. Initial studies indicated that the luminal domains of both sensors bind BiP under resting conditions, and this association is lost under ER stress (Bertolotti et al., 2000; Shen et al., 2002; Chen et al., 2002). In addition, the primary sequence of the sensing domains of IRE1 α and PERK are similar, and a MHC-I-like groove is also predicted to be present in PERK (see comparison of primary sequences between PERK and IRE1 in Liu et al. [2000] and Credle et al. [2005]). In addition, the ER luminal domains of PERK and IRE1 α are interchangeable, without affecting the rate of activation of the proteins under ER stress conditions (Liu et al., 2000). More studies are required to define the mechanisms underlying PERK and ATF6 activation.

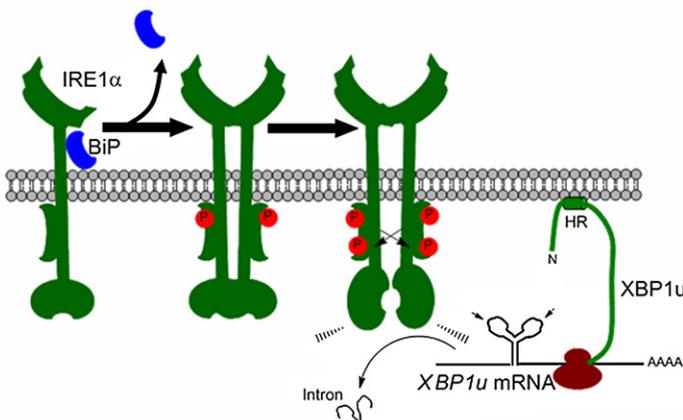
Cluster Formation by IRE1p

The crystal structure of the cytosolic domain of IRE1p was recently solved by two independent groups (Korennykh et al., 2009; Lee et al., 2008b) (Figure 1C). Korennykh and coworkers were able to visualize the architecture of IRE1p oligomers and depicted a high-order rod-shaped assembly of the cytosolic domain. This polymer-like organization was critical for IRE1p signaling as demonstrated through targeted mutagenesis of the interaction interfaces between dimers (Korennykh et al., 2009). The tridimensional structure revealed a possible mechanism where oligomerization of IRE1p dimers positions the kinase domain for transautophosphorylation, generates the RNase active site, and creates an additional interaction surface for binding of the mRNA substrate (Korennykh et al., 2009). Oligomerization of the unphosphorylated IRE1p opens the kinase domain and positions it for transautophosphorylation as a second step, leading then to activation of the RNase domain (Figure 2A). The authors also speculated that the association

A Mechanism of yeast IRE1p signaling



B Mechanism of IRE1 α signaling



between multiple IRE1p polymers may underlie visible foci formation in yeast cells undergoing ER stress.

Cluster formation of IRE1 α has not yet been described and early studies from David Ron's laboratory suggested that IRE1 α forms mostly dimers, in contrast to PERK that multimerizes in high-molecular weight complexes upon activation (Bertolotti et al., 2000). Further studies are needed to resolve these issues regarding the similarities and differences in the mechanism of sensing ER stress in yeast and mammals.

mRNA Targeting to IRE1

Recent work has provided novel insights into how the HAC1/XBP1 mRNA is recognized by IRE1p/IRE1 α . Intriguing differences in the way that the XBP1 and HAC1 mRNAs are targeted to IRE1 α and IRE1p, respectively, were described. In unstressed yeast, most of the unspliced HAC1 mRNA is cytoplasmic and remains attached to the ribosomes in an untranslated form due to intrinsic properties of its secondary mRNA structure (Figure 2A). ER stress leads to colocalization of the HAC1 mRNA to the IRE1p clusters, and this process was shown to be

Figure 2. Mechanism of IRE1 Activation in Yeast and Mammals

(A) A direct recognition model proposes that unfolded proteins bind directly to the luminal domains of IRE1p, facilitating the assembly of highly ordered IRE1p clusters exemplified by the parenthesis and "n" IRE1p units). This oligomerization may orient the cytosolic region of the dimer to form the ribonuclease active site and generation of an mRNA docking region. BiP dissociation from IRE1p may play an indirect role in unfolded-peptide loading. Oligomerization of IRE1p is essential for its autophosphorylation between dimers (as indicated with arrows). IRE1p clusters recruit untranscribed HAC1 mRNA contained in ribosomes (inhibited by the secondary structure of the HAC1 intron), an association that depends on structural motifs in IRE1p and the HAC1 mRNA including a bipartite element at the 3' end (3'BE). (B) In mammalian cells, IRE1 α is maintained in a repressed state through an association with BiP. Upon ER stress BiP dissociates, leading to partial IRE1 α phosphorylation and IRE1 α dimerization mediated by the N-terminal ER luminal region. Dimerization triggers further phosphorylation events (autophosphorylation, indicated with arrows) and activation of the RNase domain of IRE1 α . The unspliced XBP1 mRNA is translated in mammals and a hydrophobic region (HR) on the nascent peptide targets the translated XBP1 mRNA to the ER membrane, enhancing its processing by active IRE1 α . XBP1 mRNA targeting to the ER membrane does not depend on the expression of IRE1 α . In (i) and (ii) splicing sites on the XBP1 and HAC1 mRNA are indicated with an arrowhead.

IRE1p-dependent (Aragon et al., 2009). HAC1 mRNA targeting to IRE1p also requires a bipartite stem loop structure in the nontranslated region of the mRNA in addition to translational repression through the intron sequence to be excised. In contrast, in mammals unspliced XBP1 (XBP1u) is normally translated, and a recent report indicated that the targeting of the XBP1 mRNA to the ER membrane is dependent on the expression of XBP1u (Figure 2B). Upon translation, it was shown that XBP1u associates with membranes and recruits the XBP1 mRNA to the ER membrane through a well-conserved hydrophobic region at its C terminus (Figure 2B) that is predicted to form an α helix. This mechanism may provide close proximity of the mRNA substrate, facilitating IRE1 α -mediated splicing (Yanagitani et al., 2009). IRE1 α was dispensable for ER association of XBP1 mRNA. In summary, it would appear that clear differences exist between IRE1 α and IRE1p signaling mechanisms.

**Diversity of IRE1 α Proximal Signaling
Control of Alarm Signaling Pathways**

In mammals, in addition to catalyzing XBP1 mRNA processing, IRE1 α has additional functions in cell signaling (Figure 3). The cytosolic domain of activated IRE1 α binds to the adaptor protein TNFR-associated factor 2 (TRAF2), triggering the activation of the apoptosis signal-regulating kinase 1 (ASK1) and cJun-N terminal kinase (JNK) pathway (Urano et al., 2000; Nishitoh et al., 2002). IRE1 α also modulates other "alarm genes" such as the activation of the p38, ERK (Nguyen et al., 2004), and NF- κ B pathways (Hu et al., 2006), possibly by the binding of the SH2/SH3 containing adaptor proteins Nck and a protein complex between inhibitor κ B kinase (IKK)/TRAF2, respectively.

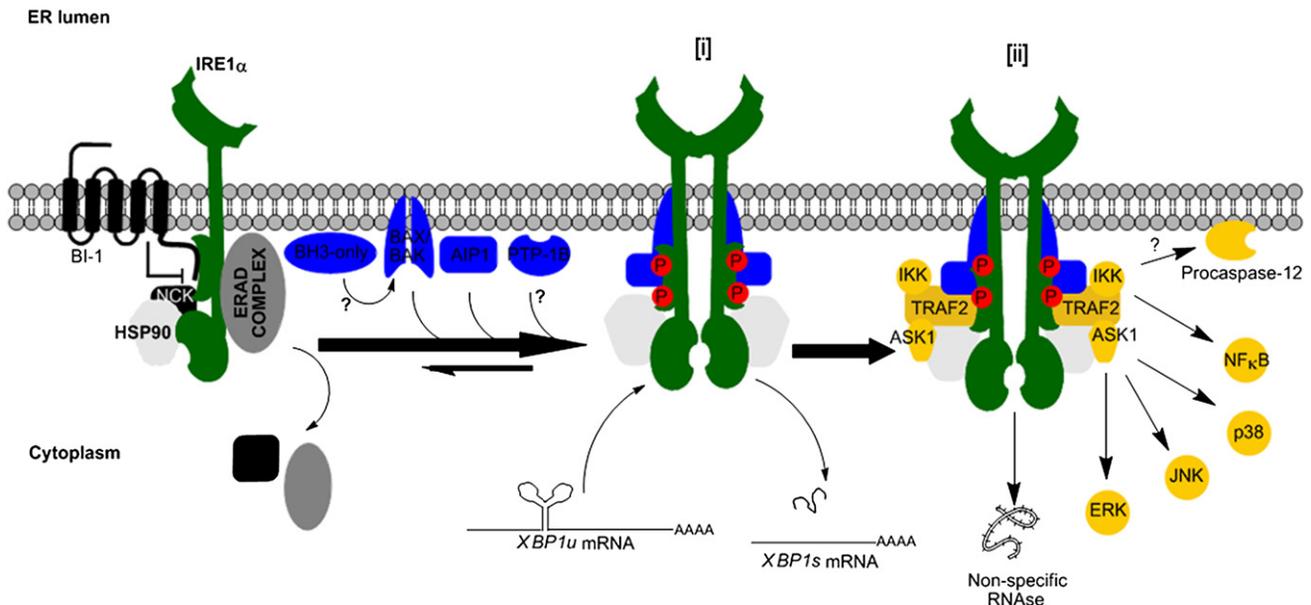


Figure 3. The IRE1 α Interactome

Mammalian IRE1 α signaling is initiated by the formation of a complex protein platform at the ER membrane here referred as the *UPRosome* where multiple factors assemble and modulate its activity. For example, activation of IRE1 α requires the binding of accessory proteins, such as BAX, BAK, AIP1, and possibly BH3-only proteins such as PUMA and BIM (upstream of BAX/BAK), in addition to the activity of the ER-located phosphatase PTP-1B. Under chronic or prolonged ER stress, IRE1 α signaling is downregulated and the ER located protein BI-1 is involved in the inactivation of IRE1 α , whereas HSP90 binding decreases its turnover. In addition, active IRE1 α initiates a variety of signaling responses through the binding of TRAF2 and possibly other adaptor proteins. These events trigger the activation of ASK1/JNK, ERK, and p38 kinases, which may regulate apoptosis and autophagy. Sequestration of IKK by IRE1 α induces NF- κ B signaling. In addition, a nonspecific RNase activity has been described for IRE1 α in flies to degrade the mRNA of proteins that have a high tendency to misfold under ER stress conditions. Inactive IRE1 α interacts with components of the ERAD machinery and may regulate this process and modulators of ERK such as NCK. For simplicity, the figure separates the components that control IRE1 α activation/inactivation in relation to XBP1 mRNA splicing activity (i), and the components related to the regulation of other signaling branches (ii), and this graphical separation of the complexes does not reflect a temporal dissociation between i and ii. Proteins that binds to inactive IRE1 α (resting condition) are shown in gray scale colors, that regulates its activation and XBP-mRNA splicing in blue and that controls other signaling pathways in yellow.

However, the function of these UPR-signaling branches in the context of protein misfolding is still not well understood. Activation of ASK1/JNK through the IRE1 α /TRAF2 complex has been proposed to mediate at least in part apoptosis under irreversible ER stress in an analogous fashion to TNF receptor signaling (Kanda and Miura, 2004; Mauro et al., 2006). In agreement with this idea, a recent high-throughput chemical screen for inhibitors of ER stress-induced cell death revealed a crucial role of ASK1 in the process (Kim et al., 2009). These data suggest that IRE1 α has a dual function in ER stress responses: (1) regulating adaptation to stress and cell survival through the control of XBP-1 s expression and (2) activation of apoptosis in cells irreversibly damaged by the activation of the JNK/ASK1 pathway.

Regulation of Autophagy

A new function for IRE1 α was recently proposed through activation of JNK whereby it controls levels of autophagy under ER stress (Ogata et al., 2006). Autophagy is a survival pathway classically linked to adaptation to nutrient starvation. Conversely, in cells undergoing ER stress, autophagy may serve as a mechanism to eliminate damaged organelles and aggregated proteins (Levine and Kroemer, 2008). Remarkably, an initial study showed that the upregulation of autophagy under ER stress conditions is initiated by the kinase domain of IRE1 α , independently of the RNase/XBP1 signaling branch (Ogata et al., 2006). How JNK regulates autophagy is not known, but a recent report suggested

that phosphorylation of the anti-apoptotic protein BCL-2 at the ER membrane by this kinase may directly affect the initiation of autophagy by modulating the activity of Beclin-1 (Pattinre et al., 2009), an essential component of the autophagy machinery (Levine and Kroemer, 2008). A genomic screen in fly cells demonstrated that knocking down UPR components, including XBP-1, increases basal autophagy levels (Arsham and Neufeld, 2009). Besides, autophagy-defective cells show upregulation of essential ER chaperones (Mathew et al., 2009), suggesting a close homeostatic balance between the autophagy and UPR pathways.

Interaction with the ER-Associated Degradation Machinery

A well defined subset of XBP1s target genes are related to ERAD and the ER translocon, including EDEM, HERP and Sec61. Using a two-hybrid screen a recent study identified the physical interaction between the ubiquitin specific protease (USP) 14 and IRE1 α . USP14 interacted with the cytoplasmic region of inactive IRE1 α , and their association was inhibited by ER stress and IRE1 α activation. Interestingly, a function for USP14 in ERAD was proposed, where inhibition occurs in an IRE1 α -dependent manner (Nagai et al., 2009). In addition, the authors reported the association of IRE1 α with essential ERAD components such as DERLIN-1, DERLIN-3, SEL1, and HRD1, suggesting that inactive IRE1 α may form a macromolecular

platform with the ERAD machinery (Nagai et al., 2009). The possible physiological relevance of these findings remains to be established.

Protein Expression Control by Direct Regulation of mRNA Decay

A genome-wide search for substrates of the mRNA-splicing activity of IRE1 α revealed only HAC1 mRNA as a hit within the limits of detection, and no additional substrates were identified highlighting the specificity of the pathway (Niwa et al., 2005). Similar findings were described in a mammalian system (Nekrukenko and He, 2006). However, in insect cells, active IRE1 α was proposed to control the degradation of mRNAs encoding certain ER proteins that were predicted to be difficult to fold (Hollien and Weissman, 2006). Through gene-profiling analysis, Weissman's group demonstrated that a subset of RNAs is selectively downregulated during ER stress in an IRE1 α -dependent and XBP1-independent manner. These effects were proposed to be a direct consequence of mRNA degradation by IRE1 α ribonuclease activity (Hollien and Weissman, 2006). The authors proposed a model where selective mRNA targeting was related to misfolding propensity of certain proteins encoded by the degraded mRNA. Thus, in a dynamic way, the misfolding of a nascent protein during its translation may directly and locally activate IRE1 α and its RNase domain to produce the degradation of mRNA being translated by the local ribosome. Two recent studies described the occurrence of IRE1 α -dependent mRNA decay in mammalian cells (Hollien et al., 2009; Han et al., 2009). Surprisingly, although XBP1 splicing can be induced by artificial dimerization of IRE1 α in the absence of ER stress, IRE1 α -dependent mRNA decay was shown to require both ER stress and IRE1 α activity, suggesting that these two functions of IRE1 α are distinct (Hollien et al., 2009). While the precise mechanism involved in mRNA decay is not clear, Hollien et al. proposed that this novel function of IRE1 α is well suited to selectively decrease the production of proteins that challenge the ER at the folding level and alleviate stress. In contrast, Han and coworkers presented evidence suggesting that endonucleolytic mediated mRNA decay of ER-localized mRNAs, including those encoding chaperones, may also culminate in apoptosis (Han et al., 2009). These complex data reinforce the concept that IRE1 α controls cell fate, wherein a dual activity modulates cell survival and apoptosis in cells irreversibly damaged.

Differential Regulation of UPR Stress Sensors: IRE1 α

Recent evidence indicates that IRE1 α activation is specifically regulated by a set of different proteins (cofactors and inhibitors) (Figure 3). For example, the levels of IRE1 α signaling were shown to be controlled by the expression of the ER-located protein-tyrosine phosphatase 1B (PTP-1B) (Gu et al., 2004). The absence of PTP-1B caused impaired XBP1 splicing, JNK phosphorylation, and attenuated upregulation of XBP1 target genes such as EDEM (Gu et al., 2004). Remarkably, PTP-1B deficiency did not affect PERK signaling, suggesting a specific regulation of IRE1 α and ruling out possible general effects on protein folding at the ER lumen. However, a physical association between PTP-1B and IRE1 α was not evaluated, which could provide mechanistic insights about this regulation. A recent report pre-

sented evidence indicating that PTP-1B also regulates UPR signaling *in vivo* in diabetes models (Delibegovic et al., 2009).

IRE1 α signaling is instigated by the expression of proapoptotic BCL-2 family members (Hetz et al., 2006). The BCL-2 family is a group of evolutionarily conserved regulators of cell death composed of both anti- and pro-apoptotic members that operate at the mitochondrial membrane to control caspase activation (Danial and Korsmeyer, 2004). Accumulating evidence indicates that, in addition to the mitochondria, members of the BCL-2 family of proteins are located at the ER membrane (reviewed in Hetz, 2007). We have described a new function for the proapoptotic BCL-2 family members BAX and BAK at the ER where they positively regulate the amplitude of IRE1 α signaling by promoting its activation. These effects were specific for this branch of the UPR since PERK signaling was not altered by BAX and BAK deficiency (Hetz et al., 2006). This regulation was mediated by a physical association between the cytosolic domain of IRE1 α and BAX/BAK and depended on the BCL-2 homology (BH) domains 1 and 3, essential motifs in the regulation of apoptosis (Figure 3). This regulation was recapitulated *in vivo* after challenging BAX and BAK conditional DKO mice with the ER stress agent tunicamycin (inhibitor of N-linked glycosylation), since decreased XBP1s expression and JNK phosphorylation were observed (Hetz et al., 2006).

In addition to the "multidomain" members BAX and BAK, there is another subtype of pro-apoptotic BCL-2 family members known as the "BH3-only" proteins (i.e., BIM, PUMA, and NOXA) which contain a single α -helical domain critical for apoptosis activation. In the control of apoptosis, BH3-only proteins act as upstream activators of BAX and BAK (reviewed in Youle and Strasser, 2008). A recent report indicated that the specific expression of the BH3-only proteins BIM and PUMA at the ER leads to the activation of the IRE1 α /JNK pathway in a BAK-dependent manner (Klee et al., 2009). Notably, these findings were obtained in the absence of any ER stressor, suggesting that these BH3-only proteins are potent activators of the IRE1 α -JNK branch acting upstream of BAX and BAK. These two BH3-only proteins have been shown to be induced by ER stress (Hetz, 2007), suggesting an interesting regulatory feedback loop. In murine cells, the proteolytic processing of the ER-resident caspase-12 has been indirectly associated to the UPR pathway by an interaction with TRAF2 and possibly with active IRE1 α (Yoneda et al., 2001), but a complex between pro-caspase-12/TRAF2/IRE1 α has not been described. Translocation of BIM from microtubules to the ER membrane has been shown to activate pro-caspase-12 processing (Morishima et al., 2004), and BAX- and BAK-deficient cells are resistant to pro-caspase-12 processing (Zong et al., 2003). However, the possible role of IRE1 α in this process was not assessed in these two studies.

Similarly, the pro-apoptotic protein ASK1-interacting protein 1 (AIP1) was recently shown to specifically regulate and enhance IRE1 α signaling (Luo et al., 2008). AIP1-deficient mice and cells derived from this mouse model displayed impaired IRE1 α signaling after exposure to ER stress agents. Similar to the phenotype of PTP-1B or BAX-/BAK-deficient cells, the lack of AIP1 expression did not affect the PERK axis of the UPR. Structural and biochemical analyses suggested that AIP1 directly

interacts with IRE1 α through a domain homologous to the pleckstrin (PH domain), facilitating IRE1 α dimerization and activation (Luo et al., 2008). Interestingly, the association of AIP1, and also BAX/BAK (Hetz et al., 2006), was shown to be induced by ER stress conditions. In summary, these findings suggest a novel and specific role for pro-apoptotic genes as accessory factors for the instigation of certain early UPR-signaling events related to adaptation and survival (Figure 3). The possible allosteric site of IRE1 α that modulates its signaling remains to be determined.

Temporal Regulation of UPR Signaling: Turning Off IRE1 α

Peter Walter's group reported that XBP1 mRNA splicing levels decline after prolonged ER stress, whereas PERK signaling is sustained over time (Lin et al., 2007). The authors suggested that this may serve to sensitize cells to cell death after chronic or irreversible ER stress, shutting down the prosurvival effects of IRE1 α /XBP1 signaling, while enhancing the expression of the PERK-dependent pro-apoptotic factor CHOP/GADD153 and downstream expression of BIM (Puthalakath et al., 2007; Lin et al., 2009). However, the mechanism involved in the inactivation of IRE1 α was not addressed. A different study provided indirect evidence suggesting that a negative regulator of IRE1 α may exist. Along this line, proteasome inhibition was shown to selectively block IRE1 α but not PERK activation, perhaps by the accumulation of an unknown inhibitor (Lee et al., 2003a).

Reed and colleagues initially suggested that the IRE1 α pathway (and possibly ATF6) may be negatively modulated by the ER-located protein BAX inhibitor-1 (BI-1) in vivo (Bailly-Maitre et al., 2006). Under ischemic conditions, BI-1-deficient mice displayed increased expression of XBP1s and hyperactivated JNK in the liver and kidney, without affecting eIF2 α phosphorylation (Bailly-Maitre et al., 2006). BI-1 is a six transmembrane containing protein functionally related to the BCL-2 family of proteins (Xu and Reed, 1998). BI-1 has no obvious homology with BCL-2-related proteins, yet it physically interacts with some members of this family (Xu and Reed, 1998; Chae et al., 2004). Further studies revealed that BI-1 homologs are present in yeast, plants, viruses, and many other organisms (Chae et al., 2003; Huckelhoven, 2004), where their function remains poorly explored. Another study also suggested that BI-1 overexpression decreases activation of the UPR in vitro in classical paradigms of ER stress (Lee et al., 2007), but the mechanism underlying these observations was not directly defined.

We recently reported a direct role of BI-1 in the attenuation of IRE1 α signaling. BI-1-deficient cells showed hyperactivation of IRE1 α associated with increased XBP1 mRNA splicing and upregulation of XBP1s-dependent responses (Lisbona et al., 2009). Notably, attenuation of IRE1 α signaling over time was markedly delayed in BI-1-deficient cells, suggesting that BI-1 has a role in the inactivation of IRE1 α . The inhibition of IRE1 α by BI-1 was recapitulated in vivo in BI-1-deficient mice and flies overexpressing *Drosophila melanogaster* BI-1 (Lisbona et al., 2009), indicating that this regulation is conserved across species. However, yeast cells deficient for the putative BI-1 homolog, the Ynl305c protein (Chae et al., 2003), did not show deficiencies in HAC1 expression, suggesting that this regulation emerged in multicellular organisms (Lisbona et al., 2009).

The formation of a protein complex between the cytosolic domain of IRE1 α and BI-1 was observed. More importantly, this association was reconstituted in vitro with purified components, and BI-1 was shown to inhibit the endoribonuclease activity of IRE1 α in a cell-free system (Lisbona et al., 2009). The regulation of XBP1 splicing by BI-1 was mediated by its C-terminal cytosolic region, a domain previously linked with BI-1's anti-apoptotic activity (Chae et al., 2003) and to act as a pH sensor in the modulation of calcium release from the ER (Kim et al., 2008).

BI-1 and BAX/BAK's regulatory effects on XBP1 mRNA splicing were evident when moderate to low doses of ER stressors were employed, conditions which more closely resemble a physiological state where cells are equipped to cope with injury (adaptive conditions). In agreement with this idea, BI-1 was shown to modulate immunoglobulin secretion of primary B cells (Lisbona et al., 2009), a phenomenon strictly dependent on XBP1s activity (Iwakoshi et al., 2003; Reimold et al., 2001). Randal Kaufman's group also reported that mild to low ER stress conditions evoke distinct signaling processes, where apoptosis-related events are not observed under mild ER stress conditions (Rutkowski et al., 2006).

In addition to the regulation of IRE1 α activity, there is another checkpoint that controls the stability of IRE1 α , possibly affecting the amplitude of UPR responses. HSP90 was shown to interact with the cytosolic domain of IRE1 α and targeting this chaperone with the inhibitors geldanamycin or 514 disrupted this complex, leading to IRE1 α turnover by the proteasome (Marcu et al., 2002). HSP90 is known to regulate the stability of many protein complexes, raising the possibility that its association/dissociation with IRE1 α may control the composition of the IRE1 α interactome and its effects on cell signaling. Finally, Mori's group discovered that XBP1u markedly accumulates at the recovery phase of ER stress (Yoshida et al., 2006) to form a complex with XBP1s that is rapidly degraded by the proteasome. The authors proposed that this negative feedback loop helps to turn off the transcription of target genes during the recovery phase of ER stress.

Distinct Roles of UPR Components in Organ Physiology: Lessons from Genetic Mouse Models

XBP1 is essential for the differentiation of hepatocytes, as XBP1-deficient embryos die in utero from severe liver hypoplasia and a resulting fatal anemia (Reimold et al., 2000). The first insights about the function of XBP1 in adult animals came from studies in the immune system. XBP1 was originally identified in multiple myeloma cells as a gene that is induced by Interleukin-6 treatment (Todd et al., 2008). XBP1-deficient B cells are markedly defective in antibody secretion in vivo in response to antigenic challenge, an activity later shown to be directly dependent on XBP1 splicing in stimulated B cells (Reimold et al., 2001; Iwakoshi et al., 2003). These findings provided the first link between the UPR and secretory cell function, and soon after the role of IRE1 α was identified using similar experimental systems in plasma B cells (Zhang et al., 2005).

To circumvent the lethal liver phenotype of XBP-1^{-/-} mice, we targeted an XBP1 transgene back to liver using a liver-specific promoter (Lee et al., 2005). XBP1^{-/-};Liv^{XBP1} mice lacking

XBP1 in all organs except the liver died shortly after birth from a severe impairment in the production of pancreatic digestive enzymes leading to hypoglycemia and death. At the cellular level, expansion of the ER was drastically impaired in pancreatic exocrine cells, resulting in a complete disorganization of the ER network and decreased efficiency in zymogen granules/enzymes synthesis in the liver and salivary glands (Lee et al., 2005). These observations are consistent with a critical role of XBP1 in ER/Golgi biogenesis and phospholipid synthesis in secretory cells (Shaffer et al., 2004; Sriburi et al., 2004). Taken together with the requirement for XBP1 in plasma cell differentiation, these findings suggested that XBP1 is essential for the development of highly secretory exocrine cells.

An XBP1 conditional knockout mouse was recently generated, revealing new physiological functions of the transcription factor in diverse organs. For example, XBP1 expression in the liver is required for normal fatty acid and cholesterol synthesis (Lee et al., 2008a). In addition, PERK has been implicated in the regulation of lipogenic pathway in the mammary gland (Bobrovnikova-Marjon et al., 2008). Along this line, a recent study suggested that all three ER stress-sensing pathways share in protecting the organism against the deregulation of lipid metabolism upon experimental ER stress in vivo by regulating a subset of metabolic transcription factors related to lipid homeostasis (Rutkowski et al., 2008). In other studies, XBP1 deletion in intestinal epithelial cells triggered spontaneous enteritis secondary to Paneth cell dysfunction and increased susceptibility to induced colitis (Kaser et al., 2008). XBP1 polymorphisms were identified as risk factors for the human inflammatory bowel diseases Crohn's disease and ulcerative colitis (Kaser et al., 2008). ER stress has been also suggested as relevant in the occurrence of diabetes. Obesity induces ER stress in the liver, playing a central role in the development of insulin resistance and diabetes by triggering JNK activity via IRE1 α and inhibition of insulin receptor signaling (Ozcan et al., 2004). Mice haploinsufficient for XBP1 showed an increased susceptibility to develop insulin resistance (Ozcan et al., 2004).

Correlative evidence suggests that activation of the UPR/IRE1 α pathway may be a primary response against neurodegeneration (Matus et al., 2008). The first insights about the function of XBP1 in the nervous system came from genetic studies of human patients with bipolar disorders (Kakiuchi et al., 2003; Kato et al., 2005) and a polymorphism in the XBP1 promoter was shown to be a risk factor for this illness in the Japanese population (Kakiuchi et al., 2003). However, these findings are still under debate (Hou et al., 2004). In contrast to the drastic phenotypes described in secretory organs, the specific deletion of XBP1 in the brain was shown to have no effect on the development of the central nervous system and did not trigger any spontaneous illness or enhance the progression of a prion disease model (Hetz et al., 2008).

The phenotypes caused by defects in the PERK/eIF2 α and IRE1 α /XBP-1 pathways are disparate, indicating some divergence in their functions in vivo. For example, UPR-mediated translational control through eIF2 α phosphorylation is not required for B lymphocyte maturation and/or plasma cell differentiation (Zhang et al., 2005). Similarly, PERK knockout mice do not show any deficiency in B cell function (Gass et al.,

2007). Interestingly, PERK deficiency leads to abnormalities in the exocrine pancreas with decreased secretion of digestive enzymes, distended ER, and increased apoptosis of acinar cells (Harding et al., 2001; Zhang et al., 2002), although the phenotype is modest as compared to XBP1^{-/-};LivXBP1 mice. Instead, PERK deficiency or lack of eIF2 α phosphorylation causes progressive loss of pancreatic islet β cells and impaired bone formation, indicating a function for the PERK pathway in β cells and osteoblasts rather than other secretory cells (Zhang et al., 2002; Harding et al., 2001; Scheuner et al., 2005). In contrast, XBP1^{-/-};LivXBP1 mice do not show drastic changes at birth in the endocrine pancreas, reflected in normal levels of insulin and insulin-containing granules in β cells. The PERK/eIF2 α pathway activates a broad range of target genes, which is not surprising given that various cellular stresses converge upon the regulation of eIF2 α activity. In contrast, as mentioned above, XBP1 target genes largely increase the folding capacity of the ER, trigger ER/Golgi biogenesis, and improve the quality control system through ERAD regulation.

The Emergence of the UPRosome Concept

The differences in cell type-specific requirements for PERK and IRE1 α pathways could be attributed to distinct modes of activation and/or downstream target genes regulated by each UPR branch. PERK and IRE1 α share functionally similar luminal sensing domains, and both are activated in cells treated in vitro with ER stress inducers. The ER stress sensing domains are, as mentioned, interchangeable between the two proteins without affecting cytosolic signaling (Liu et al., 2000). Based on the observation that PERK and IRE1 α are selectively activated in vivo, it is feasible that additional mechanisms differentially regulate their activity. Based on the compelling data discussed in this review, we speculate that the specific activation of ER stress sensors in different tissue contexts may be related to the engagement of particular regulatory complexes through the association of adaptors and direct binding of modulators.

We envision a model where a complex signaling platform is assembled at the level of IRE1 α to modulate its activation status in terms of signaling intensity and kinetics of activation/inactivation (Figure 3). In the context of cell fate and proteostasis, the fine tuning of UPR signaling responses is particularly relevant in life-to-death transitions by controlling transcriptional programs that regulate adaptation to stress or by initiating apoptosis of irreversibly damaged cells. To refer to the existence of an IRE1 α signaling macromolecular complex, we have previously used the term "UPRosome" (Hetz and Glimcher, 2008). This platform initiates multiple signaling responses in a highly regulated manner, providing a mechanism for selectivity and specificity in the signaling of IRE1 α (Figure 3). It remains to be determined if PERK and ATF6 are regulated in a similar manner by specific factors (i.e., UPRosome-2, UPRosome-3). It is interesting to note that a set of apoptosis-related proteins (i.e., BAX, BAK, AIP1, BI-1, and maybe PTP-1B and BH3-only proteins) interacts with IRE1 α , regulating its activation status. These findings suggest a model wherein the expression of anti- and pro-apoptotic proteins at the ER membrane may determine the amplitude of UPR responses and the ability of a cell to adapt to ER injuries or to initiate apoptosis (Figure 3). It remains to be

determined if distinct or dynamic IRE1 α -containing complexes exist in a tissue-specific context to regulate its signaling. This model may also be useful in addressing how the transition between the catalysis of XBP-1 mRNA processing and mRNA decay occurs. Based on the fact that active IRE1 α clusters in a highly organized manner, extensive biochemical characterization of the IRE1 α interactome is required to determine the stoichiometry of this complex and how its protein composition evolves under conditions of mild and chronic ER stress. This characterization will be particularly relevant due to the divergent and diverse effects of the UPR in cell physiology related to the control of essential processes such as secretion, folding, autophagy, calcium signaling, organelle biogenesis, apoptosis, inflammation, and cellular differentiation. Exploration of the molecular control of fine tuning of the UPR may provide new therapeutic targets to modulate ER stress responses in the setting of diverse diseases conditions such as cancer, diabetes, autoimmunity, and neurodegeneration.

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