Published in final edited form as:

Nature. 2009 February 5; 457(7230): 687–693. doi:10.1038/nature07661.

The unfolded protein response signals through high-order assembly of Ire1

Alexei V. Korennykh^{1,3}, Pascal F. Egea¹, Andrei A. Korostelev⁴, Janet Finer-Moore¹, Chao Zhang^{2,3}, Kevan M. Shokat^{2,3}, Robert M. Stroud¹, and Peter Walter^{1,3}

¹Department of Biochemistry and Biophysics, University of California at San Francisco, San Francisco, California 94158, USA

²Department of Cellular and Molecular Pharmacology, University of California at San Francisco, San Francisco, California 94158, USA

³Howard Hughes Medical Institute, University of California at San Francisco, San Francisco, California 94158, USA

⁴Department of Molecular Cell and Developmental Biology at University of California, Santa Cruz, Santa Cruz, California 95064, USA

Abstract

Aberrant folding of proteins in the endoplasmic reticulum activates the bifunctional transmembrane kinase/endoribonuclease Ire1. Ire1 excises an intron from *HAC1* messenger RNA in yeasts and *Xbp1* messenger RNA in metazoans encoding homologous transcription factors. This non-conventional mRNA splicing event initiates the unfolded protein response, a transcriptional program that relieves the endoplasmic reticulum stress. Here we show that oligomerization is central to Ire1 function and is an intrinsic attribute of its cytosolic domains. We obtained the 3.2-Å crystal structure of the oligomer of the Ire1 cytosolic domains in complex with a kinase inhibitor that acts as a potent activator of the Ire1 RNase. The structure reveals a rod-shaped assembly that has no known precedence among kinases. This assembly positions the kinase domain for *trans*-autophosphorylation, orders the RNase domain, and creates an interaction surface for binding of the mRNA substrate. Activation of Ire1 through oligomerization expands the mechanistic repertoire of kinase-based signalling receptors.

Approximately one-third of all proteins in eukaryotes enter the endoplasmic reticulum (ER) for processing and folding. The quality of protein folding is monitored by the ER-membrane-resident kinase/RNase Ire1, which is activated by misfolded proteins. On activation, Ire1 initiates a non-spliceosomal mRNA splicing reaction. Translation of the spliced mRNA yields an unfolded protein response (UPR)-specific transcription factor, termed Hac1 (ref. 1) in yeasts and Xbp1 (ref. 2) in metazoans, that induces a comprehensive transcriptional program to adjust

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Correspondence and requests for materials should be addressed to A.V.K. (alexei.korennykh@ucsf.edu).

Author Contributions A.V.K. designed and prepared protein and RNA constructs and carried out kinetic and biophysical analyses. A.V.K. and P.F.E. carried out crystallization and data collection. A.A.K. performed structure determination. J.F.-M. and A.V.K. contributed to crystallographic data processing and model building. C.Z. and K.M.S. selected and provided the kinase inhibitors. P.W. and R.M.S. supervised the work. A.V.K. and P.W. wrote the manuscript.

Author Information Atomic coordinates and structure factors for the reported crystal structure have been deposited in the Protein Data Bank under accession number 3FBV. Reprints and permissions information is available at www.nature.com/reprints.

Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

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the protein-folding capacity of the ER according to need. The UPR is activated in cancers^{3, 4}, viral infections⁵, protein-folding diseases^{6, 7} and other cellular anomalies^{8, 9}.

Under ER stress conditions, the ER-luminal domain of Ire1 acts as a sensor of unfolded proteins¹⁰ (Fig. 1a). It crystallizes as a polymer that has two distinct crystallographic interfaces important for function¹⁰. This feature can explain an early observation of oligomerization of Ire1 during the UPR¹¹ and provide a structural rationalization of Ire1 organization into UPR-induced clusters (foci) that can be observed by live cell imaging^{12, 13}. Oligomerization of the ER-luminal domain of Ire1 has been proposed to promote dimerization and activation of the kinase/RNase domains^{10, 14} analogous to other cell-surface signalling receptors¹⁵. Additionally, *trans*-autophosphorylation and binding of ADP are thought to contribute to activation of the Ire1 RNase^{16, 17}.

A recent crystal structure of the kinase/RNase domain of Ire1 reveals a two-fold symmetric dimer with a back-to-back arrangement of the kinase domains, compactly attached to an RNase dimer proposed to have two independent active sites¹⁴. The back-to-back arrangement of the kinases in the dimer is unexpected because it positions the phosphorylation sites in the activation loops more than 40 Å away from the active site of the partnering molecule in the dimer. This arrangement does not seem to be productive for the *trans*-autophosphorylation of Ire1 observed *in vivo*¹¹. We propose that a different Ire1 dimer enables the *trans*-autophosphorylation reaction (below). Dimerization of the RNase domains has been proposed to match functionally the conserved pair of splice sites in *HAC1/Xbp1* mRNA¹⁴ (Fig. 1b). Our results indicate an alternative explanation because we observe fully reactive RNA substrates that contain only a single splice site, as well as poorly reactive RNA substrates that contain dual splice sites. In this work, we combine several approaches to show that the cytosolic region of Ire1 from *Saccharomyces cerevisiae* undergoes spontaneous oligomerization that activates Ire1 for signalling in the UPR.

Activation of the Ire1 RNase by oligomerization

We prepared variants of the cytosolic portion of Ire1 that contain the kinase and the RNase domains (Ire1KR), as well as the kinase and the RNase domains extended by 24 (Ire1KR24), 32 (Ire1KR32) or 120 (Ire1KR120, ref. 18) amino acids towards the amino terminus. These extensions are part of a ~120-amino-acid-long linker domain that tethers the kinase/RNase domains to the transmembrane region (Fig. 1a, c and Supplementary Fig. 1a). Ire1KR120 showed an RNase activity indistinguishable from that of Ire1KR32 but proved unsuitable for crystallization and was not pursued further. All Ire1 constructs site-specifically cleaved 5'-³²P-labelled stem-loop oligoribonucleotide¹⁷ (HP21) derived from the *Xbp1* mRNA (Fig. 1b and Supplementary Fig. 1b; Methods). The observed rate constant exhibited a non-Michaelis dependence on the enzyme concentration and increased cooperatively with a Hill coefficient $n = 2$ for Ire1KR and Ire1KR24 and, surprisingly, a Hill coefficient $n = 3.5-8$ for Ire1KR32 (Fig. 1d). This observation indicates that the RNase activity of Ire1 arises from self-association with the formation of predominantly dimers for Ire1KR and Ire1KR24, and oligomers for Ire1KR32.

At protein concentrations above 10 μM, reactions with Ire1KR32 appeared as a heterogeneous suspension, indicating self-association of Ire1KR32 (Fig. 2a). The presence of several oligomeric species was apparent on analytical ultracentrifugation of the sample (Fig. 2b). The oligomerization could be readily reversed and RNase activity suppressed by addition of salt to the solution (Fig. 2a, c). The visible aggregation seemed to be specific because it was strongly induced by cofactors. In contrast, solutions of Ire1KR and Ire1KR24 remained clear at all concentrations with no signs of protein oligomerization, consistent with the lower cooperativity of their activation profiles (Fig. 1d).

Ire1KR32 cleaved HP21 ~100-fold faster than did Ire1KR and Ire1KR24 (Fig. 2d, left panel). The observed rate constants were compared at 1 μ M concentration of the enzymes, at which reactions occur in the same kinetic regime characterized by a log-linear concentration response of the observed rate constant. The catalytic advantage and the highly cooperative activation of Ire1KR32 were even more apparent with the *Xbp1* 443-base polymer, which more closely mimics the natural mRNA substrate of Ire1 (Fig. 2d, right panel, and Supplementary Fig. 2). These observations demonstrate that the N-terminal linker domain, particularly the eight amino acids that constitute the difference between Ire1KR32 and Ire1KR24 (Supplementary Fig. 1a), defines the self-association properties and the RNase activity of the cytosolic domains of Ire1. Notably, point mutations within these eight amino acids abrogate Ire1 signalling *in vivo*¹⁹. This functionally important linker extension was absent in the Ire1 variant used previously to obtain the back-to-back dimer structure¹⁴. The use of Ire1 that includes the extended N terminus resulted in the crystal structure of the oligomeric state of Ire1.

Structure of the Ire1 oligomer

To establish the mechanism of Ire1 oligomerization and activation, we used X-ray crystallography. Because efforts to crystallize Ire1KR32 with ADP produced crystals unsuitable for X-ray data collection, we attempted to co-crystallize Ire1KR32 with structurally diverse protein kinase inhibitors. Remarkably, several kinase inhibitors activated the RNase function revealing synthetic activators of wild-type Ire1 (Fig. 3a, b and Supplementary Fig. 3). These results have profound implications for therapeutic uses of kinase inhibitors (see Conclusions).

Crystals obtained with the inhibitor APY29 allowed determination of the structure of the Ire1KR32•APY29 complex at 3.9 Å resolution. The resolution improved to 3.2 Å with a mutant version of Ire1KR32, Ire1KR32 Δ 28•APY29, in which we deleted the α F– α EF loop (28 amino acids, 865–892). The α F– α EF loop is not evolutionary conserved and was disordered in the 3.9-Å structure. Its deletion had no effect on the RNase activity of Ire1 (Supplementary Fig. 4). Electron density for the APY29 molecule was found in the ATP-binding pocket of the Ire1 kinase domain (Fig. 3c). The position of APY29 indicates that it could form three hydrogen bonds with residues Glu 746 and Cys 748 of the main chain and two additional hydrogen bonds or van der Waals contacts with the side chains Asn 751 and Asp 828 at the active site (Fig. 3d).

Although all tested compounds can potentially form hydrogen bonds with the protein backbone (Fig. 3b), the most potent activators, APY29 and APY24, also interact with the side chain Asn 751 and insert bulky aromatic rings in place of the ribose-phosphate moiety of ADP. Manual fitting of the FDA-approved anti-cancer drug Sunitinib guided by known structures of kinase•inhibitor complexes (Protein Data Bank (PDB) IDs 2G9X and 2F4J) predicts that the compound fills the adenine-binding site, but not the ribose and the phosphate subsites. Such partial occupancy could explain the fairly good binding of Sunitinib to Ire1 accompanied by partial activation of the enzyme (Fig. 3a). AIN54 could not be fit to the ATP pocket owing to steric clashes with the β 1 strand.

The interactions of APY29 with the nucleotide-binding pocket closely mimic those of ADP except that APY29 does not use a divalent metal ion for docking (Fig. 3d, e). Accordingly, addition of EDTA inhibits the RNase activity of Ire1 for reactions stimulated by ADP but not APY29 (Fig. 3f). These findings support a model first proposed based on Ire1 mutants¹⁶: that ADP and kinase inhibitors activate Ire1 RNase by filling the ATP pocket. For maximum activity the adenine and the ribose subsites should be occupied, apparently to stabilize the active open conformation of the kinase that favours self-association of Ire1. Electrostatic interactions

due to coordination of the magnesium ion and the phosphate groups of ADP do not have an indispensable role as the charged moieties can be replaced with neutral space-filling groups.

In contrast to Ire1, which lacks the oligomerization-inducing N-terminal segment and crystallizes as a back-to-back dimer¹⁴, Ire1KR32 and Ire1KR32Δ28 crystallize as a symmetric high-order assembly (Fig. 4a, b). Fourteen Ire1 molecules constitute the asymmetric unit in the crystal lattice. Formation of the oligomer can be described by incremental addition of symmetric back-to-back Ire1 dimers to an end of a growing filament, with a simultaneous clockwise turn of 51.4° per dimer, with a complete 360° turn every 14 molecules.

The use of 14-fold non-crystallographic symmetry (NCS) improved the quality of averaged electron density maps and helped the modelling of all of the regions missing from the starting model (Supplementary Fig. 5). The structure of the kinase/RNase domain in the oligomer is similar to that in the Ire1•ADP dimer¹⁴. However, tight packing of Ire1 in the oligomer compared to the crystal packing of the Ire1•ADP dimers (Fig. 4a, inset) orders several fragments of Ire1 absent in the previous model (coloured green in Figs 4c–e). None of the new elements belong to the interface IF1^c defined previously in the back-to-back dimer¹⁴ (Fig. 4b, c). Two new interfaces, IF2^c and IF3^c, form in addition to the interface IF1^c in the oligomer. Interface IF2^c has a two-fold symmetry and forms by contacts between the RNase domains of monomers A–D, C–F, and so on (Fig. 4d). Interface IF3^c creates a linear side-to-side arrangement of monomers into filaments (B→D→F and, with opposite polarity, A←C←E). Interface IF3^c is formed by contacts between the kinase domains and involves two new elements, the αD' helix and the activation loop (Fig. 4e). The oligomerization-inducing N-terminal extension (residues 641–662) was disordered. Its structure and the mechanism of facilitating Ire1 oligomerization remain to be determined. It is possible that part of the N-tail contacts a dimerization interface, as proposed recently for the arginine-rich linker extension of epidermal growth factor receptor²⁰.

Architecturally, the oligomer resembles the double helix of DNA (Fig. 4b and Supplementary Fig. 6), where interface IF1^c parallels the interaction between nucleobases of opposing strands and interface IF3^c parallels phosphodiester linkages between nucleotides of the same strand.

The *trans*-autophosphorylation complex of Ire1

In the Ire1 oligomer, each kinase offers its activation loop to a new partner, thereby extending the filamentous oligomeric assembly. This interaction resembles the side-to-side arrangement of kinase dimers implicated in *trans*-autophosphorylation reactions^{21,22}. Ire1KR32 used in this work contained 17 phosphorylated residues (Supplementary Fig. 7). Phosphorylation was not observed on expression of Ire1 with a kinase-inactivating mutation (D828A), indicating that all Ire1 phosphates derive from its own kinase activity as the protein is expressed in *Escherichia coli*. Mass spectrometric analyses localized the phosphorylation sites to the activation loop and the αEF–αF loop (Supplementary Table 1), both of which face interface IF3^c. Together, the oligomer structure and mass spectrometry support a model wherein IF3^c serves for transfer of the phosphates *in trans*, resolving the difficulty in explaining *trans*-autophosphorylation of Ire1 in the back-to-back dimer¹⁴. The tightly packed oligomer makes it highly unlikely that kinases other than Ire1 have access to the phospho-acceptor sites. This feature can explain the specific phosphorylation of sites in Ire1 that are not part of any recognizable consensus motif and the apparent absence of other kinases known to phosphorylate Ire1.

Three phosphorylated residues important for Ire1 activation *in vivo*¹¹—Ser 840p, Ser 841p and Thr 844p—are resolved in the crystal structure (Fig. 4e). Thr 844p forms two intramolecular salt bridges positioned to stabilize the open state of the activation loop and conserved among kinases²³. Ser 840p and Ser 841p form two additional intramolecular salt bridges, and Ser 841p forms a unique intermolecular salt bridge with an adjacent Ire1 molecule at interface IF3^c. All

three phosphates are ideally placed to help Ire1 oligomerization by stabilizing the oligomerization-compatible open state of Ire1 kinase and positioning Ser 841p to stabilize interface IF3^c.

Three Ire1 interfaces control the RNase activity

The presence of the three distinct interfaces in the oligomer structure raised questions about their relative contribution to activation of the Ire1 RNase. Thus, we characterized Ire1 variants with each interface selectively impaired by mutations. For IF1^c, we prepared Ire1 with an E988Q mutation, which had the strongest deleterious effect on the RNase activity among the tested RNase IF1^c mutants¹⁴. For IF2^c and IF3^c, we identified previously uncharacterized contacts and mutated relevant residues to destabilize these contacts (Fig. 5a, b).

In the standard cleavage assay (3 μ M Ire1, HP21 substrate), mutations at all three interfaces had significant deleterious effects (Fig. 5c). Mutations mapping to IF2^c and IF3^c exhibited as strong or stronger effects on the RNase activity as did the mutation mapping to IF1^c. Sedimentation profiles and activation profiles show that mutations at each of the interfaces weaken the self-association properties of Ire1KR32 (Supplementary Fig. 8). The functional importance of residues at all three interfaces indicates a conjoint effort from IF1^c, IF2^c and IF3^c in activation of the RNase.

The activation mechanism of the Ire1 RNase

It has been suggested that dimerization activates the RNase of Ire1 by complementing the pair of splice sites in the *HAC1/Xbp1* mRNA¹⁴. In principle, this mechanism could explain why the stem-loop HP21 is cleaved slowly compared to *HAC1/Xbp1* mRNA (Fig. 2d). However, several pieces of evidence do not support the proposed model. Only Ire1KR32 shows a large preference for cleavage of *HAC1/Xbp1* mRNA over stem-loop HP21, whereas Ire1KR and Ire1KR24 show no discrimination (Fig. 2d). Furthermore, we prepared a 354-nucleotide RNA substrate that contains only a single stem-loop but reacts with Ire1KR32 at the rate of *HAC1* and *Xbp1* mRNA (Fig. 6a, b). We also prepared a 58-nucleotide RNA substrate that contains two stem-loops but reacts with Ire1KR32 at a rate of HP21 (Fig. 6c). These findings show that substrates with dual stem-loops do not have catalytic advantage compared to substrates with a single stem-loop and that self-association activates the Ire1 RNase by a mechanism different from steric complementation of the dual splice sites. In the oligomer structure, the RNase domains are linked into a continuous ribbon by two interfaces, IF1^c and IF2^c (Supplementary Fig. 9). IF2^c places the α 3' helices from the adjacent RNase monomers in reciprocal contact (Fig. 6d). The α 3' helix as well as the adjacent α 3'– α 4 loop (coloured green in Fig. 6d, e and Supplementary Fig. 9) are disordered in the back-to-back dimer structure¹⁴, indicating that IF2^c stabilizes this helix-loop element (designated HLE). Location of HLE near a dimerization interface may provide a dynamic switch that controls the RNase activity of Ire1. Indeed, HLE completes the proposed RNase active site and creates a cavity characteristic for substrate-binding pockets of enzymes (Fig. 6e), and point mutations in the HLE inactivate the RNase¹⁴.

An important additional contribution of oligomerization to the RNase activation may result from the extensive molecular surface of the oligomer, which would provide interactions with the substrate mRNA not possible with a monomer or a dimer. *Xbp1* mRNA binds to Ire1KR32 several orders of magnitude better than HP21 (Supplementary Fig. 10), indicating that protein–mRNA binding interactions spread beyond a stem-loop binding site of Ire1. These interactions could explain why Ire1KR and Ire1KR24, which do not form oligomers, do not discriminate between large and small RNA substrates (Fig. 2d). We conclude that yet to be characterized extended contacts between the Ire1 oligomer and mRNA feature prominently in Ire1 function.

Mechanistic implications

Key attributes of Ire1 activation emerged soon after its discovery and include Ire1 self-association, *trans*-autophosphorylation and the binding of ADP as a cofactor (Fig. 1a). Our present functional and structural data rationalize each of these events. In particular, we show that the primary step activating the Ire1 RNase is the self-assembly of the cytosolic region into a helical rod structure (Fig. 4a). The self-association equilibrium built into the cytosolic kinase/RNase module must be subservient to the ligand-controlled oligomerization of the ER-luminal domain of Ire1 to establish the flow of the UPR signal from the ER lumen towards the cytosol. Therefore, aggregation of the ER-luminal domain of Ire1 by unfolded proteins would serve to increase the local concentrations of the kinase/RNase domains on the cytosolic side of the ER membrane, passing the threshold for oligomerization and, consequently, RNase activation.

The roles and the temporal separation of *trans*-autophosphorylation and ADP binding are now clear. Oligomerization of the unphosphorylated Ire1 opens the kinase domain and positions it for *trans*-autophosphorylation (Supplementary Fig. 11a). ATP enters the opened kinase and phosphorylates the activation loop *in trans* to lock it in the oligomerization-compatible open state and to introduce a phosphate-mediated salt bridge at the interface IF3^c. These events provide positive feedback for oligomer assembly (Fig. 6f). Binding of a cofactor occurs in the open state of Ire1 kinase, shifts the equilibrium from monomers towards multimers and provides an additional, phosphorylation-independent level of positive modulation for the activating transition (Fig. 6f). At increased concentrations, Ire1 self-associates and becomes activated independent of phosphorylation (Supplementary Fig. 11b) and cofactor binding (Fig. 1d, open circles), directly supporting the model wherein oligomerization is the earliest and centremost step of Ire1 activation. Cofactor binding and phosphorylation enhance the self-association properties of Ire1 but neither is strictly required.

A model for the structure of the UPR-induced Ire1 foci^{12,13} emerges from our work. Oligomers formed by the ER-luminal domain of Ire1 and the cytoplasmic domains can be arranged to give similar periodicity of monomers on both sides of the ER membrane (Supplementary Fig. 12a). The resulting mesh could provide a platform for the formation and growth of supramolecular Ire1 foci in two dimensions. The length of the linkers connecting the functional domains of Ire1 to the transmembrane region permits this arrangement (Supplementary Fig. 12b). Such an assembly would allow a cooperative response to unfolded proteins and a prolonged time to mount and extinguish the UPR.

Conclusions

Our knowledge of the multi-domain signalling proteins built around protein kinases is fairly immature. This work shows how one of these molecules, Ire1, operates by forming a supramolecular structure not observed previously. We found that kinase inhibitors—including Sunitinib—act as potent activators of the Ire1 RNase. Because Ire1 provides cytoprotective function^{24,25} from which cancer cells may benefit, it may be of therapeutic value to separate the intended function of kinase inhibitors towards the targets for which they were designed from activation of Ire1. Conversely, Ire1 activation might contribute to the beneficial effects of kinase inhibitors, including Sunitinib, in mouse models of type 1 diabetes²⁶, and the cytoprotective effect of the Ire1 activators may be harnessed to combat protein-folding diseases.

METHODS SUMMARY

Proteins were expressed in *E. coli* and purified using glutathione *S*-transferase (GST)-affinity purification and size-exclusion chromatography. The DNA oligonucleotides were made by

PCR or purchased from IDT. RNA oligonucleotides were purchased from Dharmacon Inc. or prepared by *in vitro* transcription with T7 RNA polymerase. All kinetic assays were done at 30 °C and neutral pH. Diffraction data were collected from cryo-preserved crystals at a beamline 8.3.1 (Advanced Light Source, Berkeley National Laboratories). The structure was solved at 3.2 Å resolution by molecular replacement followed by refinement in CNS27 and PHENIX28. The 2.4-Å structure of Ire1 dimer (PDB ID 2RIO14) was used as a molecular replacement search model in PHASER29. The final model containing amino acids 663–864 and 893–1115 of Ire1 has R/R_{free} of 0.235/0.283 and excellent stereochemistry (PDB ID 3FBV; Supplementary Fig. 13 and Supplementary Tables 2–5). A part of the N-tail (residues 641–662) is disordered.

Supplementary Material

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Acknowledgments

We thank A. Krutchinsky for the help with MALDI instruments and for the tryptic digest analysis of Ire1KR32, F. Gruswitz for useful discussions, C. Waddling for managing the protein crystallization facility of the molecular structure group (MSG) at UCSF, and to the staff of the beamline 8.3.1 at the Advanced Light Source (Berkeley). We thank members of the Walter laboratory for critical review of the manuscript. A.V.K. is a recipient of Jane Coffin Childs fellowship. C.Z. was supported by a grant from the National Parkinson Foundation. R.M.S., J.F.-M. and P.F.E. were supported by an NIH grant RO1 GM60641. P.W. and K.M.S. are Investigators of the Howard Hughes Medical Institute.

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Figure 1. Activation of Ire1 by self-association

a, A general scheme of Ire1 activation during the UPR summarizing the key events. The kinase domain of Ire1 is coloured light brown; the RNase domain is coloured purple. TM, transmembrane. **b**, Schematic representation of RNA substrates used in this work. Triangles mark sites of specific cleavage by Ire1. **c**, Ire1 constructs used for cleavage assays and structure determination. **d**, Cooperative activation profiles for Ire1KR, Ire1KR24 and Ire1KR32 obtained using 5'-³²P-HP21, with (filled circles) and without (open circles) cofactor. E, enzyme. Asterisks are used for reference in Fig. 2. Assay details are provided in Methods.

Figure 2. Linker controls the oligomerization and activation of Ire1

a, Observation of visible self-association of Ire1KR32 that can be reversed by salt (NaCl). **b**, Analytical ultracentrifugation reveals monomers and dimers for Ire1KR as well as dimers and higher-order assemblies for Ire1KR32. Conditions were as in Fig. 1d; open symbols, 13.5 μ M Ire1 (20 $^{\circ}$ C). **c**, Salt inhibits the RNase activity of Ire1KR32. **d**, Ire1KR32 has higher RNase activity against HP21 and *Xbp1* RNA compared to Ire1KR and Ire1KR24 (Supplementary Fig. 1b; 2). Error bars show variability between single-exponential fits from two to five independent measurements. Conditions similar to those used in Fig. 1d are marked * and **.

Figure 3. Kinase inhibitors activate the RNase of wild-type Ire1

a, Activation of Ire1KR32 (3 μM) in the presence of different kinase inhibitors. **b**, Inhibitor structures, with probable hydrogen bonds shown by dashed lines. **c**, σ_A -weighted $3F_{obs} - 2F_{calc}$ map for APY29 bound to Ire1KR32Δ28 contoured at 1.5σ . **d**, The network of probable hydrogen bonds between APY29 and Ire1. **e**, The network of interactions between ADP•Mg and Ire1 (PDB ID 2RIO). **f**, Chelation of magnesium inhibits Ire1 RNase in the presence of ADP, but not of APY29. Error bars show variability between single-exponential fits from two independent measurements. Reactions contained 2 mM ADP or 100 μM APY29.

Figure 4. Structure of the Ire1 oligomer

a, Assembly of Ire1KR32Δ28•APY29. A parallel filament in the crystal packing is shown above the main filament. Domains are coloured as in Fig. 1a. The inset shows crystal packing of Ire1 dimers (PDB ID 2RIO). Letters A–N below the structure refer to individual monomers in the asymmetric unit. K, kinase; R, ribonuclease. **b**, Three intermolecular interfaces of Ire1KR32Δ28 in the oligomer. **c**, Dimer formed via interface IF1^c. **d**, Dimer formed via interface IF2^c. **e**, Dimer formed via interface IF3^c (left). Close view of the activation loop (right). Phosphates are shown in ball representation. Arrows in **b** and **e** show the direction of the activation loop donation. Regions with previously unknown structures are coloured green.

Figure 5. Three interfaces of Ire1 contribute to the RNase activity

a, Schematic representation of the Ire1 oligomer packing. Stars mark the ATP-binding pocket of the Ire1 kinase. Lower panel shows a single monomer; upper panel shows packing of four monomers. **b**, Contacts at the intermolecular interfaces of the oligomer. **c**, Mutations of the predicted interface residues designed to weaken IF1^c, IF2^c and IF3^c inhibit the RNase activity of Ire1KR32. Reactions contained 5'-³²P-HP21, 3 μM Ire1KR32 and 2 mM ADP. WT, wild type. Error bars show variability between single-exponential fits from two independent measurements. The colour of the bars matches that in **a** and **b**.

Figure 6. The mechanism of Ire1 activation

a, Time courses for cleavage of *Xbp1* (lanes 1–6) and of *Xbp1*/PstI (lanes 7–12) with 5 nM Ire1KR32. **b**, Quantification of the gel in **a**. **c**, Cleavage of RNA with one and two splice sites by Ire1KR32. Error bars show variability between single-exponential fits from two to five independent measurements. nt, nucleotide. **d**, Position of the HLE at the RNase–RNase interface IF2^c. **e**, Molecular surface representation of the RNase domain in the oligomer and in the dimer structures. Putative catalytic residues in **d** and **e** are coloured orange; HLE is coloured green. **f**, Revised model of Ire1 activation during the UPR.