Misfolding of mutant adenine nucleotide translocase in yeast supports a novel mechanism of Ant1-induced muscle diseases

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ABSTRACT
Approximately one-third of proteins in the cell reside in the membrane. Mutations in membrane proteins can induce conformational changes and expose nonnative polar domains/residues to the lipid environment. The molecular effect of the resulting membrane stress is poorly defined. Adenine nucleotide translocase 1 (Ant1) is a mitochondrial inner membrane protein involved in ATP/ADP exchange. Missense mutations in the Ant1 isoform cause autosomal dominant progressive external ophthalmoplegia (adPEO), cardiomyopathy, and myopathy. The mechanism of the Ant1-induced pathologies is highly debated. Here we show that equivalent mutations in the yeast Aac2 protein cause protein misfolding. Misfolded Aac2 drastically affects the assembly and stability of multiple protein complexes in the membrane, which ultimately inhibits cell growth. Despite causing similar proteostatic damages, the adPEO- but not the cardiomyopathy/myopathy-type Aac2 proteins form large aggregates. The data suggest that the Ant1-induced diseases belong to protein misfolding disorders. Protein homeostasis is subtly maintained on the mitochondrial inner membrane and can be derailed by the misfolding of one single protein with or without aggregate formation. This finding could have broad implications for understanding other dominant diseases (e.g., retinitis pigmentosa) caused by missense mutations in membrane proteins.

INTRODUCTION
Adenine nucleotide translocase (Ant) belongs to a large family of proteins known as mitochondrial carriers, which transport nucleotides, cofactors, signaling molecules, and metabolic intermediates across the mitochondrial inner membrane (Palmieri, 2004, 2014). Ant primarily catalyzes the export of ATP out of mitochondria by exchanging with cytosolic ADP (Klingenberg, 2008). Under low-membrane potential conditions, Ant reverses its transport directionality, allowing the import of cytosolic ATP into mitochondria against matrix ADP. In addition to ATP/ADP exchange, Ant also mediates proton leak across the membrane (Brand et al., 2005). This activity mildly uncouples the membrane, which prevents hyperpolarization and excessive production of superoxide.

Ant is one of the most conserved proteins through evolution and is present in multiple isoforms in different species. In humans, Ant1 is predominantly expressed in skeletal muscle and heart (Stepien et al., 1992). Yeast cells express three isoforms of Ant, with Aac2 being the major ADP/ATP carrier under respiring conditions (Lawson and Douglas, 1988; Lawson et al., 1990). Ant contains three repeats of a sequence module of \( \sim 100 \) amino acids, with each predicted to form two transmembrane \( \alpha \)-helices (Saraste and Walker, 1982; Walker and Runswick, 1993). The compact organization was confirmed by atomic structures established for the bovine Ant1 and yeast Aac2 and Aac3 isoforms in the presence of the competitive inhibitor carboxyatractyloside (Pebay-Peyroula et al., 2003; Ruprecht et al., 2014). These structures revealed that the six transmembrane helices in an Ant monomer form a central channel that is sufficiently large to accommodate the bulky ADP and ATP. This led to the alternating-access transport model, predicting that an
Ant monomer alternates between the cytosolic and matrix conformations, which allows the alternate access and then translocation of cytosolic ADP and matrix ATP during the nucleotide exchange cycle (Kunji and Crichton, 2010).

Several missense mutations in Ant1 (A90D, L98P, D104G, A114P, and V289M) cause autosomal dominant progressive external ophthalmoplegia (adPEO), characterized by late or adult-onset muscle weakness and exercise intolerance (Kaukonen et al., 2000; Napoli et al., 2001; Komaki et al., 2002; Siciliano et al., 2003; Deschauer et al., 2005). Another missense mutation, A123D, leads to hypertrophic cardiomyopathy and myopathy in homozygous patients (Palmieri et al., 2005). These diseases are commonly manifested by multiple mitochondrial DNA (mtDNA) deletions. Several models have been proposed to explain the Ant1-induced pathogenesis. Fontanesi et al. (2004) found that the adPEO-type mutants preferentially import ATP over ADP in vitro reconstituted proteoliposomes. This novel property was proposed to promote a futile ATP/ATP homoexchange and elevated mitochondrial ATP and dATP levels, which consequently affect the accuracy of mtDNA replication and stability. In mouse C2C12 myotube cells, Kawamata et al. (2011) found that mitochondria of myotubes replicating the ANT$^{A114P}$ allele are switched to the reversal ATP$_{cytosol}$/ADP$_{matrix}$ exchange mode at a higher membrane potential. This may also lead to increased ATP level and possibly, nucleotide imbalance in the mitochondrial matrix.

More recently, Ravaud et al. (2012) showed that the missense pathogenic variants of Ant1 have altered nucleotide-binding affinity and translocation kinetics when expressed on the plasma membrane of Escherichia coli. This observation raises the possibility that loss of nucleotide transport function may be responsible for the pathogenesis, possibly by a haploinsufficient mechanism. However, this model is inconsistent with the findings that both humans and mice heterozygous for a complete loss-of-function ant1 allele are healthy (Graham et al., 1997; Palmieri et al., 2005; Echaniz-Laguna et al., 2012) and that homozygous ant1-null alleles do not cause ophthalmomilia or ocular motility defect (Echaniz-Laguna et al., 2012; Strauss et al., 2013). The latter findings tend to suggest that the pathogenesis induced by the missense mutations involves a novel mechanism that is different from the loss-of-transport-function mode.

Our previous studies showed that the adPEO-type Ant mutations in a yeast model severely affect cell viability in a dominant manner and that mtDNA instability in the mutant cells arises independent of adenine nucleotide transport (Chen, 2002; Liu and Chen, 2013; Wang et al., 2008a,b). We proposed that the mutant proteins may interfere with general mitochondrial biogenesis in addition to their effect on oxidative phosphorylation and mtDNA stability and that mtDNA instability likely arises as a secondary effect of mitochondrial damage. By using the filamentous fungus Podospora anserine as a model system, El-Khoury and Sainsard-Chanet (2009) also showed that expression of the adPEO-type Ant variants leads to severe mitochondrial damage, as manifested by low mitochondrial membrane potential. The mechanism by which the mutant Ant causes mitochondrial damage is unknown. In the present study, we provide evidence suggesting that pathogenic missense mutations cause the misfolding of Ant, which subsequently induces proteostatic stress on the membrane and loss of cell viability.

**RESULTS**

**Mutant Aac2 proteins affect protein quality control on the mitochondrial inner membrane**

We previously showed that the A106D, M114P, A128P, and A137D alleles of AAC2, equivalent to the pathogenic A90D, L98P, A114P, and A123D mutations in human ANT1, are synthetically lethal, with the disruption of YME1 even in cells expressing an endogenous wild-type copy of AAC2 (Wang et al., 2008a). YME1 encodes the i-AAA protease involved in the quality control of proteins on the mitochondrial inner membrane (Thorsness et al., 1993; Leonhard et al., 1999). This observation suggested that expression of the mutant Aac2 may dominantly affect protein homeostasis on the membrane, which cannot be tolerated in cells lacking Yme1. In support of this idea, we found that cells expressing a mutant allele of AAC2 are hypersensitive to diverse conditions that affect proteostasis and the biochemical properties and functionality of the membrane. First, meiotic spores cosegregating AAC2$^{A128P}$ with the disruption of OXA1 failed to form viable colonies on yeast extract/peptone/dextrose (YPD) medium (Figure 1A). OXA1 encodes the mitochondrial inner membrane insertase that mediates the insertion of mtDNA-encoded proteins from the matrix side (Bonnefoy et al., 1994; Herrmann et al., 1997; Stuart, 2002). Loss of Oxa1 directly affects the assembly of respiratory complexes on the inner membrane. Second, AAC2$^{A128P}$ was found to be synthetically lethal with ybr238cΔ (Supplemental Figure S1). Ybr238c is an inner membrane–associated protein that, together with its parologue Rmd9, controls mRNA processing/stability in mitochondria (Nouet et al., 2007; Williams et al., 2007). Given that the ybr238cΔ single mutant has barely detectable defect in respiratory growth, the synthetic lethality suggests that mitochondria expressing AAC2$^{A128P}$ are sensitive to subtle disturbance to the respiratory complexes on the membrane due to reduced mitochondrial gene expression. Third, the AAC2$^{A128P}$ allele was also synthetically lethal, with the disruption of PSD1 (Figure 1B), encoding the phosphatidylserine decarboxylase on the inner membrane involved in the synthesis of phosphatidylethanolamine (Figure 1C). Compositional alterations to the lipid environment on the membrane are therefore detrimental to mitochondria expressing AAC2$^{A128P}$. Finally, we found that haploid cells coexpressing the wild-type AAC2 and AAC2$^{A128P}$ are hypersensitive to antimycin on YPD, which inhibits the bc$_1$ complex in the electron transport chain (Supplemental Figure S2). This indicates that the defect in the electron transport chain synergizes with AAC2$^{A128P}$, which leads to membrane stress and the inhibition of cell growth.

To provide direct evidence for the proteostatic stress hypothesis, we examined whether general protein quality control is affected on the inner membrane in cells coexpressing the wild-type and a mutant variant of Aac2. The hemagglutinin (HA)-tagged NADH dehydrogenase Nde1 is unstable in vivo and is stabilized in cells defective in the Yme1 protease (Figure 1, D and E; Augustin et al., 2005). We found that the Yme1-dependent turnover of Nde1-3HA is marginally delayed by AAC2$^{A128P}$ and AAC2$^{A137D}$ (Figure 1, F and G) but significantly reduced by AAC2$^{A106D}$ and AAC2$^{M114P}$ (Figure 1, H and I). These data suggest that the mutant Aac2 proteins dominantly affect the turnover of misfolded proteins on the inner membrane, including Nde1-3HA.

**Aac2 variants with the adPEO-type mutations are prone to aggregation**

We hypothesized that the mutant Aac2 proteins may be misfolded, which causes general proteostatic stress on the inner membrane. Misfolded proteins often have increased propensity to form aggregates. To test whether the mutant Aac2 proteins form aggregates, mitochondria were isolated from cells cultured at 30°C. After detergent solubilization, the proteins were analyzed by Blue Native PAGE (BN-PAGE). As shown in Figure 2A, the monomeric wild-type Aac2 migrated as a band of ∼60 kDa, in contrast to an
In contrast to the adPEO-type mutant Aac2 proteins, no apparent aggregates were detected with the cardiomyopathy- and myopathy-type Aac2\(^{A137D}\) variant under the same conditions (Figure 2B). The monomeric form of Aac2\(^{A137D}\) was noticeably reduced in BN-PAGE. This was not caused by protein degradation, as SDS–PAGE analysis of \(n\)-dodecyl-\(β\)-maltoside (DDM)–extracted mitochondrial proteins showed that the level of Aac2\(^{A137D}\) was little changed after the 16-h incubation (Figure 2C). It is possible that Aac2\(^{A137D}\) has a poor reactivity with the antibody under the native conditions.

Further, in organello aggregation assays revealed that Aac2\(^{A106D}\), Aac2\(^{M114P}\), and Aac2\(^{A128P}\) formed aggregates 8–12 h after the incubation at 25°C (Figure 2E). Extended incubation eventually led to noticeable reduction of signals for the aggregates. This could be due to the formation of excessively large aggregates that become resistant to DDM extraction and separation by BN-PAGE. The time-course study also confirmed that Aac2\(^{A137D}\) does not form aggregates during the entire period of incubation at 25°C.
To evaluate the possible effect of Aac2 misfolding on membrane function, we first determined whether the assembly state of the oxidative phosphorylation pathway is dominantly affected in strains coexpressing the wild-type and mutant AAC2 alleles. In yeast mitochondria, the complexes III (coenzyme Q:cytochrome c oxidoreductase) and IV (cytochrome c oxidase) in the electron transport chain form the III$_2$IV$_2$ and III$_2$IV$_1$ supercomplexes to facilitate substrate channeling, whereas complex V (ATP synthase) is present in both monomeric and dimeric forms (Schagger and Pfeiffer, 2000).

The mutant Aac2 proteins affect the assembly and stability of multiple protein complexes on the inner membrane

The three adPEO-type mutations occur at Ala-106, Met-114, and Ala-128, which are located at the end of $\alpha$-helices 2 (H2) and 3 (H3) on the intermembrane space side, with their side chains facing the lipid phase in the carboxylatractyloside-inhibited conformation (Figure 3A). The cardiomyopathy- and myopathy-type mutation occurs at Ala-137, which is also located on H3 but is deeply embedded in the membrane with its side chain relatively facing inward.

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supramolecular structures in wild-type mitochondria can be readily resolved by BN-PAGE (Figure 3B). We found that expression of the four mutant Aac2 proteins drastically affected the formation of these structures. IIIIV₁ was reduced by AAC2\(^{A106D}\) and AAC2\(^{A128P}\) and became barely detectable in cells expressing AAC2\(^{A106D}\) and AAC2\(^{A137D}\). IIIIV₂ was severely reduced by all the four mutant alleles (Figure 3B). Immunoblotting of the native gels revealed that the assembly state of complex V using antisera against the α and β subunits.

Finally, we determined the effect of the misfolded Aac2 on the TIM23 complex. As shown in Figure 4B, TIM23 was barely detectable in cells coexpressing the wild-type Aac2 and AAC2\(^{A106D}\) or AAC2\(^{A114P}\). The level of TIM23 was strongly reduced by AAC2\(^{A128P}\) and AAC2\(^{A137D}\). The TIM23 protein level was only slightly reduced by the mutant Aac2 alleles, as shown by SDS–PAGE analysis (Figure 4C). These observations indicate that the assembly of TIM23 is also dominantly deterred by the mutant Aac2 proteins. The overall loss of Tim23 signal in the mutant mitochondria on the BN-PAGE gel may be caused by the aggregation of unassembled Tim23 or the poor solubility of structurally altered TIM23 complex, which may cause poor protein recovery before electrophoretic separation.

The i-AAA protease Yme1 plays a critical role in the quality control of misfolded Aac2
Misfolded membrane proteins are expected to be degraded by specific protein quality control machineries. To provide further support to the model that the mutant Aac2 proteins are misfolded, we first determined the in vivo stability of AAC2\(^{A128P}\) compared with the
induced mitochondrial damage, by screening for genes that suppress Aac2\textsuperscript{A128P}-induced growth defect (Figure 6A). Expression of AAC2\textsuperscript{A128P} from the galactose-inducible GAL10 promoter strongly inhibited cell growth on galactose medium. We found that overexpression of YME1, but not RCA1 and AFG3, suppressed growth inhibition by Aac2\textsuperscript{A128P}. In contrast to YME1, which encodes the i-AAA protease to preferentially degrade misfolded membrane proteins from the intermembrane space side, YTA12/RCA1 and YTA10/AFG3 encode the two subunits of the hetero-oligomeric m-AAA protease, which degrades misfolded and unassembled proteins on the inner membrane from the matrix side (Arlt et al., 1996). Overexpression of SOD2, CTA1 and CTT1, encoding the mitochondrial superoxide dismutase and catalases that reduce oxidative stress, failed to suppress aac2\textsuperscript{A128P}. These observations suggest that the Yme1 protease likely plays a major role in the quality control of Aac2.

To further support a role of Yme1 in the surveillance of Aac2 quality, we asked whether loss of Yme1 accelerates Aac2\textsuperscript{A128P} aggregation. The pathogenic Aac2 variants are synthetically lethal with yme1Δ (Wang et al., 2008a). We previously showed that the wild type. We observed that both Aac2\textsuperscript{A128P} and its wild-type control remain stable 24 h after cycloheximide chase (Figure 5A). We then determined the stability of Aac2\textsuperscript{A128P} by an in organello assay. Isolated mitochondria were incubated at different temperatures, and the levels of Aac2 were monitored by immunoblotting (Figure 5B). We found that the wild-type Aac2 remained stable 3.5 h after incubation at all three temperatures tested. Aac2\textsuperscript{A128P} was stable at 25°C but became unstable at 30°C. Aac2\textsuperscript{A128P} degradation was further accelerated at 37°C. The temperature-dependent turnover of Aac2\textsuperscript{A128P} correlated with the growth phenotype of the AAC2\textsuperscript{A128P} mutant (Figure 5C). Incubation at 25°C, but not 37°C, inhibited cell growth on complete medium containing glucose or glycerol plus ethanol as carbon sources. It remains unclear as to why AAC2\textsuperscript{A128P} turnover is detected in the in organello but not in vivo assays. We speculate that the in organello conditions may exacerbate the misfolding of the mutant protein, which accelerates its recognition and turnover by protein quality control systems.

We next attempted to identify potential protein quality control mechanisms and other cellular pathways that might reduce Aac2\textsuperscript{A128P}-induced mitochondrial damage, by screening for genes that suppress Aac2\textsuperscript{A128P}-induced growth defect (Figure 6A). Expression of AAC2\textsuperscript{A128P} from the galactose-inducible GAL10 promoter strongly inhibited cell growth on galactose medium. We found that overexpression of YME1, but not RCA1 and AFG3, suppressed growth inhibition by Aac2\textsuperscript{A128P}. In contrast to YME1, which encodes the i-AAA protease to preferentially degrade misfolded membrane proteins from the intermembrane space side, YTA12/RCA1 and YTA10/AFG3 encode the two subunits of the hetero-oligomeric m-AAA protease, which degrades misfolded and unassembled proteins on the inner membrane from the matrix side (Arlt et al., 1996). Overexpression of SOD2, CTA1 and CTT1, encoding the mitochondrial superoxide dismutase and catalases that reduce oxidative stress, failed to suppress aac2\textsuperscript{A128P}. These observations suggest that the Yme1 protease likely plays a major role in the quality control of Aac2.

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of the Aac2 proteins may have been degraded by other proteases before maturing on the inner membrane. The remaining mutant Aac2 that is successfully matured on the inner membrane is sufficient to affect cell survival because of severe misfolding in the absence of Yme1.

**DISCUSSION**

We found that the adPEO-type Aac2A128P, Aac2M114P, and Aac2A128P share similar biochemical properties. Of note, they all have increased propensity to form large aggregates and are capable of inducing proteostatic stress and severe defects in the assembly and structural maintenance of multiple protein complexes on the mitochondrial inner membrane. The data strongly support the view that Ant1-induced adPEO is a protein misfolding disorder. In addition to damages to the oxidative phosphorylation pathway, the mutant Ant1 likely reduces cell viability by affecting the function of multiple membrane protein complexes, including TIM22 and TIM23, which directly participate in mitochondrial biogenesis. The mutant Aac2 proteins therefore gain a toxic function, which is consistent with the dominant nature of the disease.

**FIGURE 6:** Yme1 protease plays a key role in the quality control of Aac2. (A) Suppression of Aac2A128P-induced cell death by YME1 overexpression. Aac2A128P was expressed from the GAL10 promoter on complete medium containing galactose and raffinose as carbon sources at 30°C. All the genes were cloned into a multicopy vector and introduced into the PG1 strain by transformation. (B) BN-PAGE coupled with immunoblotting showing the aggregation of Aac2A128P in fresh mitochondria prepared from a strain with the yme1Δ background. The strains CS1153/1 (rpl6Δ, AAC2), CS1227-4A (rpl6Δ, Aac2, yme1Δ), CS1374-2D (rpl6Δ, AAC2A128P), and CS1699/1 (rpl6Δ, AAC2A128P, yme1Δ) were grown at 30°C in YPD. Mitochondria were isolated and solubilized by 2% DDM. Approximately 40 μg of solubilized proteins was applied to BN-PAGE, followed by immunoblotting using anti-Aac2 antibody. (C) Steady-state levels of Aac2A128P compared with wild-type Aac2 in the presence or absence of Yme1. Mitochondria were prepared as in B, and 30 μg of proteins was analyzed by SDS–PAGE and Western blotting. The matrix Ilv5 protein was used as a loading control. (D) Relative steady-state levels of Aac2 in C. Error bars indicate the SDs of three independent experiments. *p < 0.05 (unpaired Student’s t test). (E) A model for the membrane damage caused by misfolded Aac2. See Discussion for details. The red star denotes the pathogenic missense mutations. IM, inner membrane; IMS, intermembrane space.
We found that Aac2\textsuperscript{A137D}, mimicking Ant1\textsuperscript{A123D}, which causes cardiomyopathy and myopathy (Palmieri et al., 2005), does not form visible aggregates. However, cells expressing Aac2\textsuperscript{A137D} share many common phenotypes with those expressing the adPEO-type alleles (Wang et al., 2008a). These include uncoupled respiration, cold sensitivity, \(p\)-lei-lathyl, mtDNA instability, and synthetic lethality with \(yme1\Delta\). Furthermore, as shown in the present study, Aac2\textsuperscript{A137D} also interferes with the assembly and stability of multiple protein complexes on the membrane. These findings argue that Aac2\textsuperscript{A137D} may also be misfolded and affect mitochondrial function by the same mechanism as the adPEO-type Aac2 variants. Although Aac2\textsuperscript{A137D} does not aggregate in in organello assay, it may have altered conformation that causes proteostatic stress on the membrane. Definitive proof for Aac2\textsuperscript{A137D} misfolding is lacking. If Aac2\textsuperscript{A137D} were misfolded, an extended speculation would be that protein misfolding, rather than aggregation per se, may be the primary cause of membrane damage (Figure 6E). Because the A137D mutation (or A123D in human Ant1) completely abolishes nucleotide transport activity (Palmieri et al., 2005), the cardiomyopathy and myopathy phenotypes observed in the reported homozygous patient may result from a double hit. The pathogenesis may involve both Ant1 misfolding and defective ATP/ADP exchange. In the future, it would be interesting to see whether ANT1\textsuperscript{A123D} can cause adPEO in a dominant manner in heterozygous individuals if such clinical cases become available.

The quality control mechanism of plasma membrane proteins has been extensively studied. It involves the recognition of misfolded domains by E3 ubiquitin ligases on the endoplasmic reticulum (ER) membranes, followed by ER-associated degradation (ERAD; Hirsch et al., 2009; Houck and Cyr, 2012). The removal of the misfolded proteins by ERAD decreases the delivery of functional proteins on the cell surface, leading to recessive pathologies such as cystic fibrosis. However, in other clinical cases, destabilizing mutations in transmembrane domains cause dominant diseases (e.g., retinitis pigmentosa). These diseases often involve the toxic accumulation of misfolded proteins (Sanders and Myers, 2004; Mendes et al., 2005). Not much is known about the extent to which the delivery of a misfolded protein might affect membrane functionality and cell viability. We provide evidence that the misfolded Aac2 proteins affect the assembly and stability of multiple protein complexes on the mitochondrial inner membrane. Among the adPEO-type mutations, M114P and A128P are expected to introduce a proline kink that would prematurely terminate \(\alpha\)-helices 2 and 3 (Figure 3A). A106D introduces a negative charge in the proximity of the membrane surface, which could stretch the helix 2 toward the positively charged intermembrane space. The cardiomyopathy- and myopathy-type A137D mutation introduces a negative charge at a deeper position within the membrane. Possibly, a common consequence of these structural and conformational changes could be the exposure of nonnative, especially polar residues to the membrane environment. This may induce structural perturbations (e.g., helix misalignment) in order to solvate the unfavorable energy potential within the membrane. In this regard, aggregation of Aac2\textsuperscript{A106D}, Aac2\textsuperscript{M114P}, and Aac2\textsuperscript{A128P} may be a compensatory molecular strategy that helps to shield the exposed nonnative patches to defuse the unfavorable energy potential. Although we detected visible aggregates of mutant Aac2 in freshly prepared mitochondria in cells lacking the Yme1 protease, Aac2 aggregates were not observed in fresh mitochondria isolated from YME1 cells, even when cells were grown at 25°C. Aggregates were readily detected in isolated mitochondria after extended incubation at 25°C. It remains unknown what exactly triggers the aggregation of the mutant Aac2 during the in organello incubation. Nonetheless, the data clearly indicate that the mutant proteins are biophysically distinguishable from the wild-type Aac2. Compared with Ala-106, Met-114, and Ala-128, which have side chains facing directly to the lipid phase, the side chain in Ala-137 faces relatively inwardly. It is possible that Aac2\textsuperscript{A137D} adopts a specific conformation that is unfavorable for oligomerization and aggregation.

In considering possible mechanisms of mitochondrial damage (Figure 6E), one can speculate that the misfolded Aac2 alters the surface property of the protein and exposes nonnative residues. This may in turn promote ectopic interactions with other membrane proteins. Interactions with unassembled proteins may prevent their incorporation into protein complexes. Previous studies suggested that Aac2 physically associates with multiple protein complexes in the membrane, including the III-IV supercomplexes and TIM23 (Cloppool et al., 2008; Dienhart and Stuart, 2008; Mehnert et al., 2014). The intrinsic affinity to Aac2 may explain the severe defects in the assembly of these complexes in cells expressing a misfolded Aac2. On the other hand, as Aac2 is inserted into the membrane through the TIM22 complex, the interaction with the misfolded Aac2 appears to directly destabilize the TIM22 structure, as manifested by the breakdown of the complex. As alternative mechanisms, it is also possible that the misfolded Aac2 proteins deplete the molecular chaperones and proteases required for the biogenesis and quality control of other protein complexes. The damage to TIM22 and defective membrane protein import may also contribute to the defect in the biogenesis of respiratory complexes and the TIM23 preprotein translocase.

Two AAA proteases play key roles in the quality control of proteins on the mitochondrial inner membrane (Gerdes et al., 2012). The i-AAA protease (or Yme1) has its catalytic domain facing the intermembrane space, whereas the m-AAA protease, composed of the Yta12 (or Rca1) and Yta10 (or Afg3) subunits, has the catalytic domain facing the matrix. These proteases sense the folding state of solvent-exposed domains on either side of the membrane and specifically degrade unfolded membrane proteins (Arlt et al., 1996; Leonhard et al., 1996, 1999, 2000). The suppression of Aac2\textsuperscript{A128P}-induced cell lethality by YME1 overexpression and accelerated accumulation of Aac2\textsuperscript{A128P} aggregates in cells lacking Yme1 suggest that the i-AAA protease plays a key role in the quality control of Aac2\textsuperscript{A128P}. The premature termination of \(\alpha\)-helix 2 likely distorts the conformation of the solvent-exposed loop between helices 2 and 3, which is recognized and attacked by Yme1. Whether other mutants are also recognized by Yme1 for degradation has yet to be determined.

In summary, our data strongly suggest that Ant1-induced adPEO is a protein misfolding disease, which distinguishes it from clinical conditions caused by null mutations in ANT1. Of greater importance, our work provides evidence that the misfolding of one single protein could have detrimental effects on the biogenesis and stability of other membrane complexes on the membrane and that proteostatic stress could be a significant source of damage to the oxidative phosphorylation apparatus under pathophysiological conditions. Our finding could have general implications for understanding the physiologic consequences of other destabilizing mutations in transmembrane proteins that are associated with an increasing number of dominant diseases, including retinitis pigmentosa (Sanders and Myers, 2004; Mendes et al., 2005; Hirsch et al., 2009; Houck and Cyr, 2012).

MATERIALS AND METHODS

Cell growth conditions and yeast strains

Standard media were used to cultivate yeast cells. The genotypes and sources of Saccharomyces cerevisiae strains used in this study
are listed in Supplemental Table S1. To generate the NDE1-3HA allele, the HA epitope was added to the C-terminus of NDE1 by the integration of a PCR product amplifying from the 3HA-KanMX6 cassette. The strain PG1 (MATα, his3Δ1, leu2Δ0, lys2Δ0, ura3Δ0, trp1::GAL10-AAC2::128F, His3) was derived from BY4741.

Determination of protein stability
Yeasts were grown in YPD to late exponential phase and shifted to 37°C after addition of cycloheximide (1 mg/ml). At indicated time points, three OD600 equivalents of cells were harvested. The cell extract was prepared as previously described (Chen, 2001). Approximately 5 μl of cell extracts was analyzed by SDS–PAGE, followed by Western blot using antisera against Aac2, Cox2, or HA.

Mitochondrial preparation and BN-PAGE analysis
Yeasts were grown at 25 or 30°C in YPD. Crude mitochondria were isolated as described (Boldogh and Pon, 2007), except that the cell wall was removed by enzymatic digestion at 5 mg of Zymolyase 20T/g of wet cells at 25 or 37°C. Isolated mitochondria were resuspended in SEH buffer (0.6 M sorbitol, 20 mM 4-(2-hydroxyethyl)-1-piperazinethanesulfonic acid [HEPES]-KOH, pH 7.4, 2 mM MgCl2, 1 mM ethylene glycol tetraacetic acid, 1 mM phenylmethylsulfonyl fluoride). For BN-PAGE analysis, crude mitochondria were solubilized in a 2% DDM or digitonin solution containing 50 mM NaCl, 2.5 mM MgCl2, 20 mM HEPES-KOH (pH 7.4), 10% (vol/vol) glycerol, and 1 mM EDTA (pH 7.4) on ice for 30 min. Insoluble material was removed by centrifugation of 10,000 x g at 4°C for 15 min. Approximately 30–40 μg of mitochondrial proteins were mixed with approximately 5 μl of cell extracts was analyzed by SDS–PAGE, followed by Western blot using antisera against Aac2, Cox2, or HA.

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