

Aberrant substrate engagement of the ER translocon triggers degradation by the Hrd1 ubiquitin ligase

Eric M. Rubenstein,¹ Stefan G. Kreft,¹ Wesley Greenblatt,¹ Robert Swanson,² and Mark Hochstrasser¹

¹Department of Molecular Biophysics and Biochemistry, Yale University, New Haven, CT 06520

²Department of Biology, Valparaiso University, Valparaiso, IN 46383

Little is known about quality control of proteins that aberrantly or persistently engage the endoplasmic reticulum (ER)-localized translocon en route to membrane localization or the secretory pathway. Hrd1 and Doa10, the primary ubiquitin ligases that function in ER-associated degradation (ERAD) in yeast, target distinct subsets of misfolded or otherwise abnormal proteins based primarily on degradation signal (degron) location. We report the surprising observation that fusing *Deg1*, a cytoplasmic degron normally recognized by Doa10, to the Sec62 membrane protein rendered the

protein a Hrd1 substrate. Hrd1-dependent degradation occurred when *Deg1*-Sec62 aberrantly engaged the Sec61 translocon channel and underwent topological rearrangement. Mutations that prevent translocon engagement caused a reversion to Doa10-dependent degradation. Similarly, a variant of apolipoprotein B, a protein known to be cotranslocationally targeted for proteasomal degradation, was also a Hrd1 substrate. Hrd1 therefore likely plays a general role in targeting proteins that persistently associate with and potentially obstruct the translocon.

Introduction

The ubiquitin–proteasome system accounts for a major portion of eukaryotic protein degradation. Proteins destined for proteolysis by this pathway are covalently modified with chains of the protein ubiquitin through the sequential action of ubiquitin-activating enzymes (E1s), ubiquitin-conjugating enzymes (E2s), and ubiquitin ligases (E3s; Ciechanover et al., 1980; Scheffner et al., 1995). Ubiquitylated substrates are degraded by the 26S proteasome (Ravid and Hochstrasser, 2008). In the yeast *Saccharomyces cerevisiae*, the transmembrane (TM) RING domain proteins Hrd1 and Doa10 are the predominant ER-associated degradation (ERAD) E3s that target misfolded and regulatory proteins for proteolysis (Hampton et al., 1996; Plemper et al., 1999; Wilhovsky et al., 2000; Bays et al., 2001; Swanson et al., 2001). With a few exceptions, Hrd1 and Doa10 recognize unique substrate classes depending on degron disposition relative to the ER membrane (Vashist and Ng, 2004; Carvalho et al., 2006). Specifically, Hrd1 recognizes proteins

with ER luminal or intramembrane degrons, called ERAD-L (lumen) and ERAD-M (membrane) substrates, respectively (Carvalho et al., 2006; Gauss et al., 2006; Sato et al., 2009). In contrast, Doa10 promotes degradation of proteins with cytoplasmically disposed degrons, ERAD-C (cytoplasm) substrates (Huyer et al., 2004; Ravid et al., 2006; Metzger et al., 2008). Hrd1 and Doa10 exhibit overlapping E2 requirements. Hrd1 mediates ubiquitin transfer from Ubc7 and, to a lesser extent, Ubc1 (Plemper et al., 1999; Bays et al., 2001), whereas Doa10 functions with Ubc6 and Ubc7 (Swanson et al., 2001).

The prototypical Doa10 substrate is the transcriptional repressor MAT α 2, the proteolytic control of which is critical for regulating the yeast-mating phenotype (Swanson et al., 2001; Laney and Hochstrasser, 2003). The N-terminal 67 residues of MAT α 2, *Deg1*, contain a degron for Doa10-dependent degradation (Hochstrasser and Varshavsky, 1990; Johnson et al., 1998; Swanson et al., 2001). Importantly, *Deg1* confers Doa10-dependent instability when fused to the cytoplasmic N termini of heterologous soluble proteins (Hochstrasser and Varshavsky, 1990; Chen et al., 1993; Johnson et al., 1998;

Correspondence to Mark Hochstrasser: mark.hochstrasser@yale.edu

S.G. Kreft's present address is Dept. of Biology, University of Konstanz, 78457 Konstanz, Germany.

W. Greenblatt's present address is Harvard-MIT Division of Health Sciences and Technology, Harvard Medical School, Boston, MA 02115.

Abbreviations used in this paper: ERAD, ER-associated degradation; PTM, post-translational modification; PTT, posttranslational translocation; TM, transmembrane; WT, wild type.

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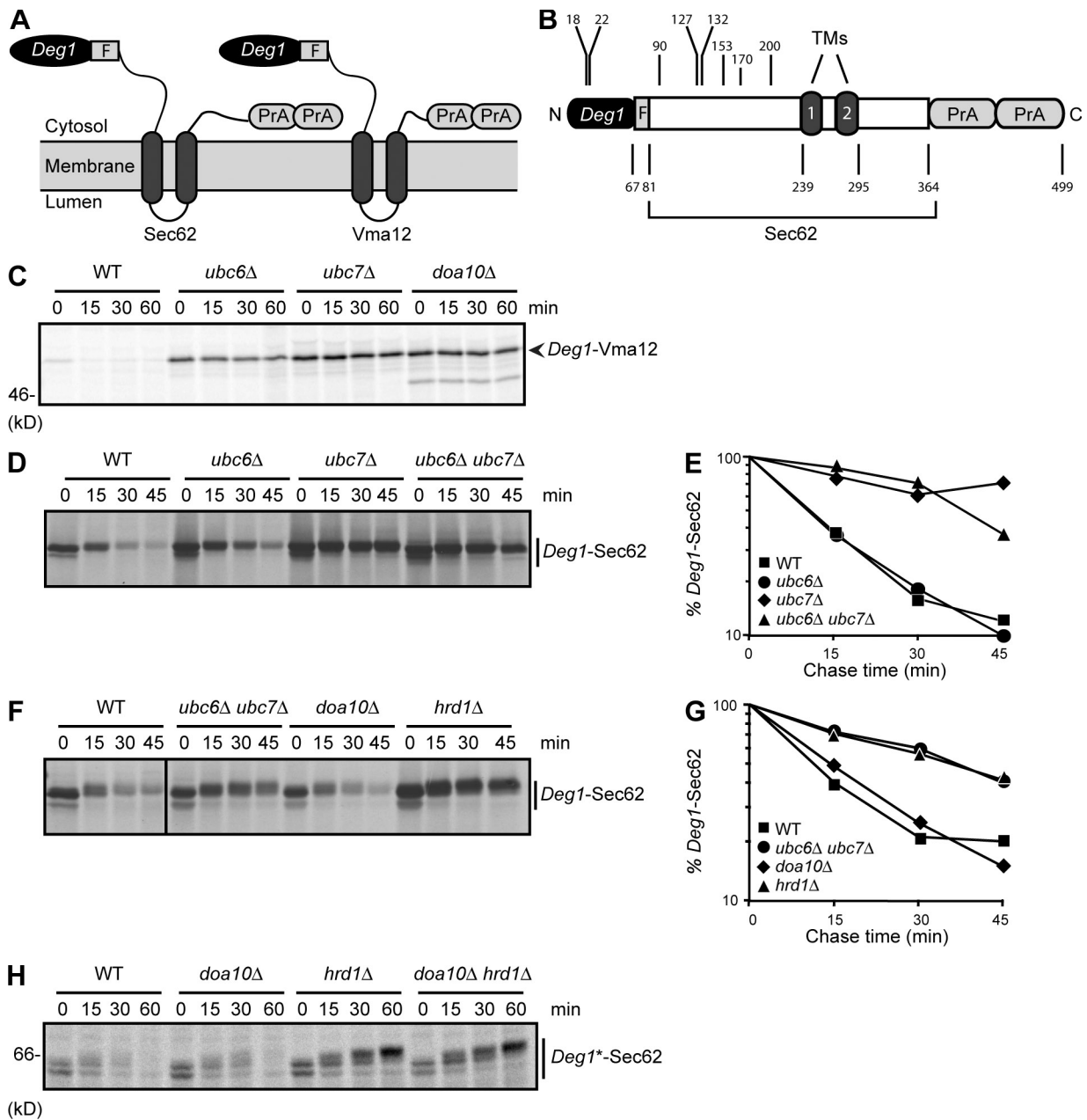


Figure 1. ***Deg1-Sec62* is a Hrd1 substrate.** (A) Schematic diagram of *Deg1* fusion proteins (with predicted topologies) used in this study. For each construct, the N-terminal *Deg1* is followed, in sequence, by the Flag (F) epitope, the 2-TM ER protein (Sec62 or Vma12), and two copies of the Protein A (PrA) tag. For clarity, the fusion proteins are referred to as *Deg1-Sec62* or *Deg1-Vma12*. Any additional alterations to protein sequence will be noted. (B) Linear representation of *Deg1-Sec62*, drawn to scale. Positions of boundaries between elements within the fusion protein and residue numbers mentioned in the text are highlighted. (C) Pulse-chase analysis of *Deg1-Vma12* in the indicated yeast strains. (D and F) Pulse-chase analysis of *Deg1-Sec62* in the indicated yeast strains. Molecular weight markers for these autoradiographs are not available; however, migration of the same protein can be seen in Fig. 4 B. (E and G) Quantification of autoradiographs in D and F. Data are representative of three experiments. The black line indicates that intervening lanes have been spliced out. (H) Pulse-chase analysis of *Deg1*-Sec62* in the indicated yeast strains. *Deg1* fusions in C, D, and F were precipitated with anti-*Deg1* antibodies. *Deg1*-Sec62* in H was precipitated with anti-Flag antibodies. Cycloheximide was included in the chase depicted in H. *Deg1**, F18S/I22T double mutant.

Swanson et al., 2001). Besides soluble *Deg1* fusions, we have fused *Deg1* to the ER-resident TM protein Vma12 to study Doa10-dependent ERAD (Ravid et al., 2006). *Deg1* has also been fused to the topologically similar ER protein Sec62 (Mayer et al., 1998). Like Vma12, Sec62 has two TM segments, with both protein termini on the cytoplasmic side of the membrane (Deshaies and Schekman, 1990). Attachment of *Deg1* destabilizes both Sec62 and Vma12 (Mayer et al., 1998;

Ravid et al., 2006; Scott and Schekman, 2008). As expected for a Doa10 substrate, *Deg1-Sec62* is stabilized in cells lacking both Ubc6 and Ubc7 (Mayer et al., 1998). Because eliminating Ubc7 inactivates both Doa10 and Hrd1 pathways, however, it remained formally possible that *Deg1-Sec62* is a Hrd1 substrate. We show that this is indeed the case and that aberrant engagement of the translocon by *Deg1-Sec62* is responsible for this E3 switch. This engagement triggers a topological

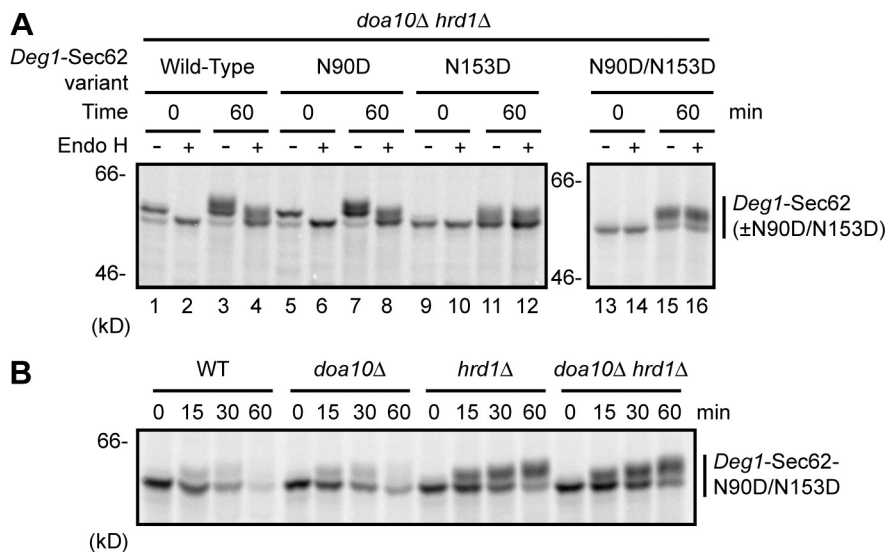


Figure 2. N-glycosylation of *Deg1-Sec62*. (A) *doa10Δ hrd1Δ* yeast cells expressing *Deg1-Sec62* or *Deg1-Sec62* with the indicated consensus N-glycosylation sites mutated were pulse labeled for 10 min and lysed immediately or after 60 min in the presence of excess nonradioactive amino acids and cycloheximide. *Deg1* fusion proteins were precipitated with anti-Flag antibodies and incubated in the presence or absence of Endo H. (B) Pulse-chase analysis of *Deg1-Sec62-N90D/N153D* in the indicated yeast strains. *Deg1-Sec62* was precipitated with anti-Flag antibodies. Cycloheximide was included in the chase.

change in *Deg1-Sec62* that is closely associated with Hrd1-dependent degradation.

We extended our investigation to a known cotranslationally degraded protein. Apolipoprotein B (apoB), the major protein component of low- and very-low-density lipoproteins, stalls in the translocon if it cannot associate with its lipid ligand and is subsequently degraded by the proteasome (Yeung et al., 1996; Fisher et al., 1997; Pariyarath et al., 2001). We found that, like *Deg1-Sec62*, an apoB variant expressed in yeast that is also associated with the translocon (Hrizo et al., 2007) is targeted by the Hrd1 ubiquitin ligase. These results suggest that Hrd1 may play a general role in the degradation of proteins that aberrantly or persistently engage the translocon.

Results

Deg1-Sec62 is a Hrd1 substrate

We used pulse-chase analysis to examine the metabolic stability of the ER-resident TM proteins Sec62 and Vma12 when the *Deg1* degron was appended to their N termini (depicted in Fig. 1, A and B). *Deg1* destabilized both proteins (Fig. 1, C and D). As previously demonstrated (Ravid et al., 2006), *Deg1-Vma12* was strongly stabilized in cells lacking Doa10. It was also stabilized when either of the E2s that function with Doa10 (Ubc6 or Ubc7) were absent (Fig. 1 C). Unexpectedly, *Deg1-Sec62* was only stabilized in the absence of Ubc7, which works with either Doa10 or Hrd1, but not in cells lacking Ubc6, which functions exclusively in Doa10-dependent ERAD (Fig. 1, D and E).

We directly tested which ERAD E3s were responsible for *Deg1-Sec62* degradation. Surprisingly, deletion of *DOA10* did not impair *Deg1-Sec62* degradation, whereas the *Deg1* fusion protein was strongly stabilized in the absence of Hrd1 (Fig. 1, F and G). The Hrd1 dependence for *Deg1-Sec62* degradation was unanticipated because *Deg1* typically targets proteins to which it is fused for Doa10-dependent proteolysis. Moreover, the predicted cytoplasmic disposition of *Deg1* represents an

apparent exception to the division of ERAD-L/M and ERAD-C substrates between Hrd1 and Doa10, respectively (Vashist and Ng, 2004; Carvalho et al., 2006).

Doa10-mediated degradation of *Deg1*-harboring proteins depends on the hydrophobic face of a predicted amphipathic helix within *Deg1* (Johnson et al., 1998). When mutations in this helix that stabilize *Deg1*-containing Doa10 substrates were introduced into *Deg1-Sec62*, the resultant protein (*Deg1**-Sec62) was still degraded in a Hrd1-dependent manner (Fig. 1 H), unlike *Deg1**-Vma12, which is strongly stabilized (Ravid et al., 2006). Thus, the details underlying molecular recognition of *Deg1*-containing substrates differ between the two ERAD E3s.

Posttranslational modification (PTM) of *Deg1-Sec62*

Deg1-Sec62 migrated as multiple species by SDS-PAGE and became increasingly modified with time after its synthesis (Fig. 1, D and F). It was previously noted that *Deg1-Sec62* is N-glycosylated, in contrast to wild-type (WT) Sec62, which is not typically posttranslationally modified (Kim et al., 2006; Scott and Schekman, 2008). Immediately after radiolabeling, *Deg1-Sec62* migrated as two distinct species (Fig. 2 A, lane 1). After treatment with Endoglycosidase H (Endo H), which removes all N-glycans from yeast proteins, the slower-migrating band collapsed to the faster migrating species (lane 2). Mutational analysis revealed that N153 was the major site of N-glycosylation (with a possible minor contribution by N90; lanes 5–12), which is consistent with an earlier study (Scott and Schekman, 2008). After 60 min, the higher mass species appeared as multiple diffuse bands (lane 3). Endo H treatment only partially increased the mobility of these poorly resolved bands (lane 4), which indicates additional PTM.

Deg1-Sec62 is also modified by O-mannosylation, an ER-specific modification with a role in the ERAD of some Hrd1 substrates (Vashist et al., 2001; Hirayama et al., 2008). We performed pulse-chase analysis of *doa10Δ hrd1Δ* yeast expressing *Deg1-Sec62-N90D/N153D* cultured in the presence of a global

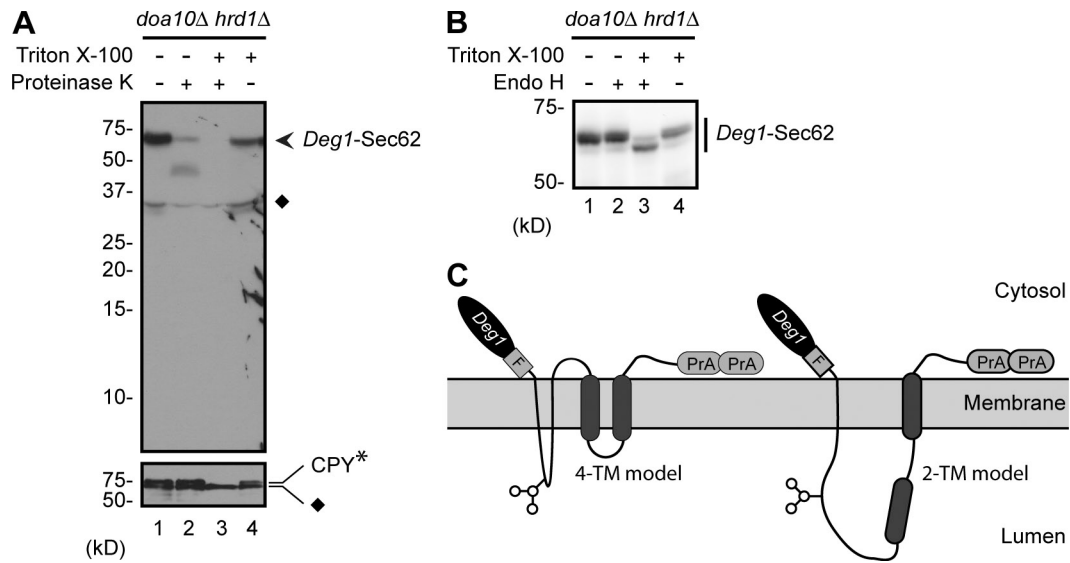


Figure 3. Membrane topology of *Deg1-Sec62*. (A) Intact microsomal membranes prepared from *doa10Δ hrd1Δ* cells expressing *Deg1-Sec62* and ER luminal control protein CPY* were treated with 5 μ g/ml Proteinase K (or mock-treated) in the presence or absence of 1% Triton X-100. Samples were separated by SDS-PAGE and detected by immunoblotting with antibodies against *Deg1* (top) or CPY (bottom). Diamonds denote nonspecific bands. Partially protease-resistant, anti-*Deg1*-reactive species are seen between 37 and 50 kD when detergent is excluded (lane 2). Species of similar size and intensity are observed when *Deg1-Vma12* is subjected to the same treatment (Fig. S3, lane 8). Thus, the *Deg1* moiety of *Deg1-Sec62* exhibits protease accessibility similar to that of *Deg1-Vma12*. (B) The same as in A, but microsomal preparations were incubated with Endo H instead of Proteinase K. *Deg1-Sec62* was detected by immunoblotting with peroxidase anti-peroxidase antibody, which recognizes the Protein A epitope. (C) Models for topological rearrangement of *Deg1-Sec62* with respect to the ER membrane. In each case, both N-terminal *Deg1* and C-terminal *Sec62* tail remain on the cytoplasmic face of the ER membrane. In the 4-TM model, the normally cytoplasmic sequence of *Deg1-Sec62* downstream of *Deg1* loops into the ER lumen, flanked by two novel membrane-spanning segments. In the 2-TM model, the first membrane-spanning segment of *Deg1-Sec62* is significantly upstream of the first normally used *Sec62* TM. The approximate position of the N-glycosylated N153 residue is indicated with a cartoon representation of the N-glycan.

inhibitor of O-mannosylation (Orchard et al., 2004). Under these conditions, the more slowly migrating species of *Deg1-Sec62*-N90D/N153D exhibited enhanced mobility but did not obtain the mobility of the fastest migrating, presumably unmodified species (Fig. S1 A). The full complement of PTMs decorating *Deg1-Sec62* and the substrate residues that they modify remain to be characterized.

Neither N-glycosylation nor O-mannosylation was required for Hrd1-mediated degradation of *Deg1-Sec62* (Fig. 2 B and Fig. S1 B). Preventing glycosylation also did not restore Doa10-mediated targeting of *Deg1-Sec62* (Fig. 2 B and Fig. S1 B). Conversely, N-glycosylation at engineered sites in *Deg1-Vma12* did not prevent its Doa10-dependent degradation (Fig. S2 A). Nonetheless, *Deg1-Sec62* PTMs serve as a convenient indicator of a topological transition that, as subsequent experiments will demonstrate, plays an important role in the switch from Doa10 to Hrd1 dependency.

Topology of *Deg1-Sec62*

Strikingly, N-glycosylation of *Deg1-Sec62* occurred on residues in the predicted cytoplasmic N-terminal domain (Fig. 1 B), whereas the glycosylation machinery resides in the ER lumen. Therefore, fusion of *Deg1* to *Sec62* must stimulate dislocation of at least a portion of the cytoplasmically disposed N-terminal domain into the ER lumen. As judged by immunoblot analysis, the majority of *Deg1-Sec62* at steady state was posttranslationally modified, and therefore topologically altered (Fig. S1 C). Previous work indicated that both the Flag epitope immediately downstream of *Deg1* and the C-terminal Protein A tag

(Fig. 1, A and B) of *Deg1-Sec62* remain on the cytoplasmic face of the ER membrane (Scott and Schekman, 2008). To confirm the cytoplasmic disposition of *Deg1*, ER-derived microsomal membranes were incubated with Proteinase K followed by anti-*Deg1* immunoblotting (Fig. 3 A). The *Deg1*-dependent topological rearrangement that allows glycosylation did not protect *Deg1* from protease exposure to the external (cytoplasmic) face of microsomal membranes in the absence of detergent (lane 2). Cytoplasmic exposure of *Deg1* is particularly intriguing given that Doa10 is incapable of targeting *Deg1-Sec62*.

The susceptibility of the *Deg1-Sec62* N terminus to proteolysis despite ER luminal access of N153 could be explained in two ways. *Deg1* fusion to *Sec62* might stimulate stable movement of a portion of the normally cytoplasmic N-terminal domain downstream of *Deg1* into the ER lumen. Alternatively, *Deg1* attachment may allow reversible movement of the entire N-terminal *Deg1-Sec62* domain into and out of the ER lumen. Return of the N terminus to the cytoplasmic face of the ER would explain protease accessibility. Consistent with the first alternative, we found that microsomal *Deg1-Sec62* was sensitive to Endo H only when detergent was added (Fig. 3 B, lane 3). Therefore, fusion of *Deg1* to *Sec62* triggers the stable movement of a portion of the N-terminal *Sec62* tail into the ER lumen while *Deg1* remains on the cytoplasmic side. These experiments are consistent with two models for the final topology of *Deg1-Sec62*, which differ in how much of the N-terminal domain is dislocated into the ER lumen (Fig. 3 C; see Discussion).

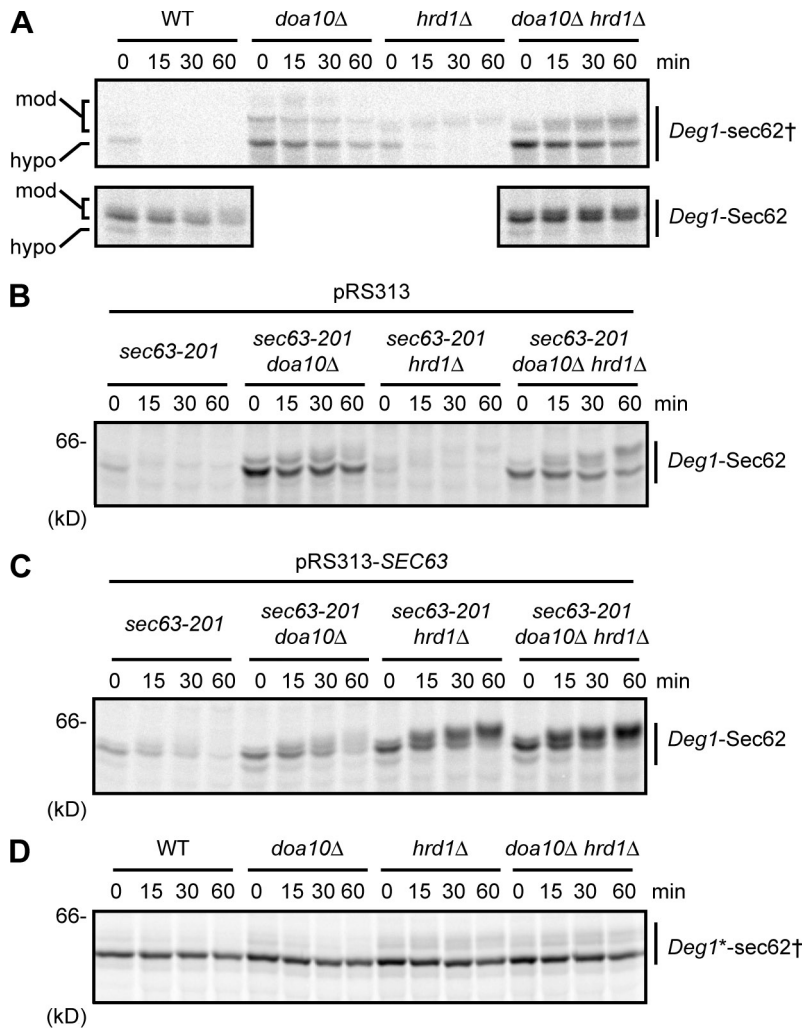


Figure 4. **Inhibiting *Deg1-Sec62* association with the translocon delays its PTM and restores *Doa10*-dependent degradation.** (A–D) Pulse-chase analysis of *Deg1-Sec62* variants in the indicated yeast strains. Where noted, yeast harbor a plasmid encoding WT *Sec63* or an empty vector. *Deg1* fusions in A and D were precipitated with anti-*Deg1* antibodies. *Deg1* fusions in B and C were chased in the presence of cycloheximide and precipitated with anti-Flag antibodies. Molecular weight markers for the autoradiograph in A are not available; however, migration of the same protein can be seen in Fig. S1 D. *sec62†*, G127D of *Deg1-Sec62*, equivalent to G37D of untagged *Sec62*. *sec63-201*, 27-residue C-terminal truncation of *Sec63*. *Deg1**, F18S/I22T double mutant.

Impaired translocon binding to *Deg1-Sec62* allows *Doa10*-dependent degradation

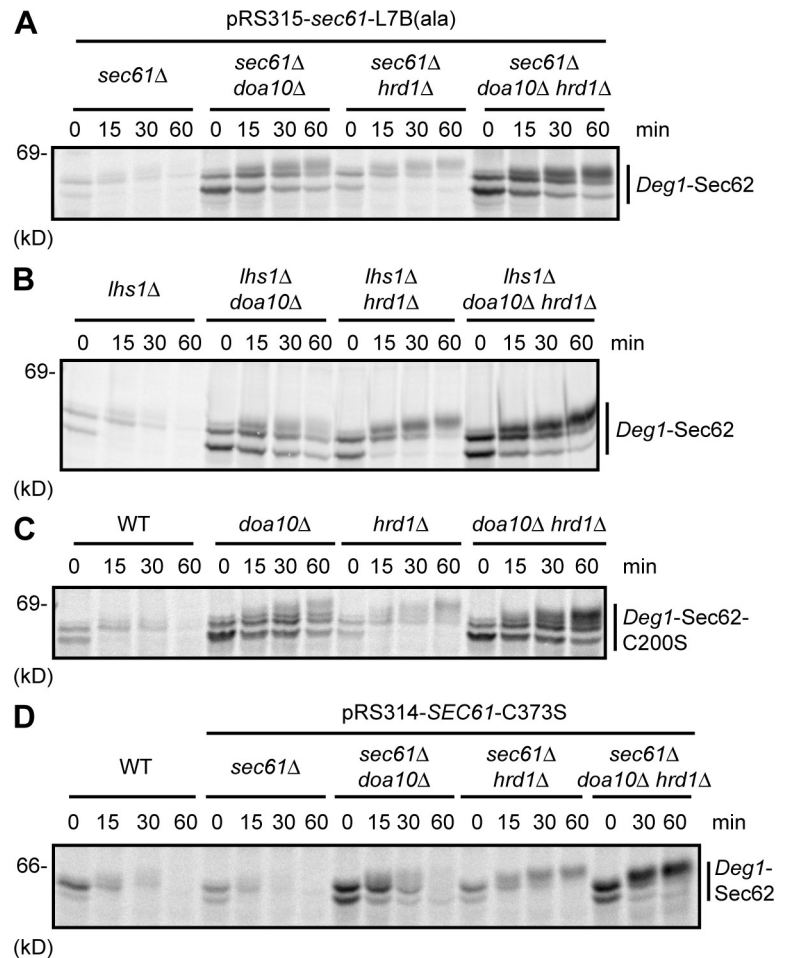
Based on genetic and physical interactions, *Sec61* has been proposed to function as a retrotranslocation channel for *Hrd1* substrates, although this remains controversial (de Virgilio et al., 1998; Willer et al., 2008; Schäfer and Wolf, 2009). *Sec62* is a component of the *Sec61* complex involved in posttranslational translocation (PTT; Deshaies et al., 1991; Ng et al., 1996). In a strain background that limits its degradation, *Deg1-Sec62* complements *sec62* mutants, which suggests that it associates with *Sec61* in a manner similar to WT *Sec62* (Mayer et al., 1998). We hypothesized that recognition of *Deg1-Sec62* as a *Hrd1* substrate is mediated by its incorporation into the translocon complex. To test this, we introduced a point mutation into *Deg1-Sec62* (G127D, *Deg1-sec62†*) that partially impairs *Sec62* association with *Sec63*, its interacting partner within the translocon complex (Deshaies and Schekman, 1990; Deshaies et al., 1991; Wittke et al., 2000), and assayed protein degradation.

Deg1-sec62† exhibited a shorter half-life than *Deg1-Sec62* (Fig. 4 A, compare WT cells, top and bottom). Strikingly, this single point mutation directed a switch in the E3 dependence of degradation (Fig. 4 A, top). Unlike *Deg1-Sec62*, *Deg1-sec62†* was strongly (though incompletely) stabilized

in *doa10Δ* cells and remained unstable in *hrd1Δ* cells. Full metabolic stabilization was not observed unless both E3s were absent, which is consistent with the incomplete ablation of *sec62†* interaction with *Sec63* (Wittke et al., 2000).

The dramatically reduced *Hrd1* dependence of *Deg1-sec62†* degradation coincided with significantly delayed glycosylation (which requires topological rearrangement of the *Sec62* N-terminal domain), which suggested a mechanistic link between the two (Fig. 4 A, compare *doa10Δ hrd1Δ* cells, top and bottom panels; and Fig. S1 D). The remaining trace of post-translationally modified *Deg1-sec62†* was stabilized in *hrd1Δ* cells (Fig. 4 A). As with the *Deg1-sec62†* mutant, the *sec63-201* mutant, which interacts poorly with WT *Sec62* (Ng and Walter, 1996; Wittke et al., 2000), caused an increase in the degradation rate, a switch in E3 specificity, and a delay in modification of *Deg1-Sec62* (Fig. 4 B). Complementation of *sec63-201* cells with WT *SEC63* caused all these changes to be reverted (Fig. 4 C). When the *sec62†* mutation was introduced in the context of *Deg1**, which is not recognized by *Doa10*, the protein was almost completely stabilized (Fig. 4 D). These data strongly suggest that persistent interaction of *Deg1-Sec62* with the translocon stimulates both *Deg1-Sec62* domain dislocation into the ER lumen and its switch to *Hrd1*-dependent degradation.

Figure 5. *Deg1*-*Sec62* topological rearrangement and *Hrd1* targeting depend on the PTT pathway. (A–D) Pulse-chase analysis of denoted *Deg1*-*Sec62* variants in the indicated yeast strains. Where noted, yeast harbor a plasmid encoding a variant of *Sec61*. Cycloheximide was included in the chase. *Deg1* fusion proteins were precipitated with anti-Flag antibodies.



Topological rearrangement of *Deg1*-*Sec62* via PTT

Most N-glycosylated proteins are modified while they are being translocated (Whitley et al., 1996; Popov et al., 1997; Scheper et al., 2003). Like WT *Sec62*, *Deg1*-*Sec62* is likely cotranslationally translocated into the ER membrane. The gradual acquisition of N-glycosylation, especially when translocon binding is partially impaired (Fig. 4 and Fig. S1 D), suggested that the topological transition allowing such modification occurs after the initial “normal” membrane insertion, possibly through the *Sec61* channel itself. We hypothesized that appending *Deg1* to *Sec62* triggers insertion of a portion of the cytoplasmic N-terminal domain through the *Sec61* translocon via a PTT mechanism after initial cotranslational translocation of *Deg1*-*Sec62* (Zimmermann et al., 2011). Moreover, we speculated that this insertion is critical for triggering *Hrd1*-dependent degradation and preventing *Doa10* recognition of *Deg1*-*Sec62*.

To test these ideas, we analyzed *Deg1*-*Sec62* E3 specificity and PTM kinetics in cells expressing a variant of *Sec61*—*sec61*-L7B(ala)—which harbors four mutations in a luminal loop and is predominantly defective for PTT (Trueman et al., 2011). Strikingly, in the context of *sec61*-L7B(ala), *Deg1*-*Sec62* degradation reverted to *Doa10* dependence (Fig. 5 A). A delay in PTM coincided with impaired degradation in *doa10*Δ cells expressing *sec61*-L7B(ala), likely signifying delayed movement of the *Sec62* N-terminal domain into the ER lumen.

The gradually appearing, posttranslationally modified species of *Deg1*-*Sec62* were cleared in a *Hrd1*-dependent manner. A similar delay in PTM and reversion to *Doa10*-dependent degradation were observed in cells lacking the BiP co-chaperone *Lhs1*, which is required for efficient PTT of many proteins (Fig. 5 B; Hamilton et al., 1999; Tyson and Stirling, 2000; Steel et al., 2004). Passage of a portion of *Deg1*-*Sec62* through the translocon is consistent with the reported formation of a transient disulfide linkage between a cysteine within the channel interior of *Sec61* and a cysteine in the N-terminal domain of *Deg1*-*Sec62* (but not WT *Sec62*; Scott and Schekman, 2008). Indeed, mutating either participating cysteine accelerated *Deg1*-*Sec62* degradation and partially switched E3 dependence from *Hrd1* to *Doa10* (Fig. 5, C and D).

In summary, these data strongly suggest that topological rearrangement of *Deg1*-*Sec62* takes place through the *Sec61* translocon by a PTT mechanism and that this is necessary for *Hrd1* recognition.

Degradation of *Deg1*-*Sec62* differs from that of previously characterized ERAD-L and ERAD-M substrates

Deg1-*Sec62* domain dislocation moves normally cytoplasmic residues of *Sec62* to the ER lumen and membrane. We therefore determined if *Deg1**-*Sec62* degradation requires *Hrd1* cofactors required for ERAD-L and ERAD-M substrate degradation.

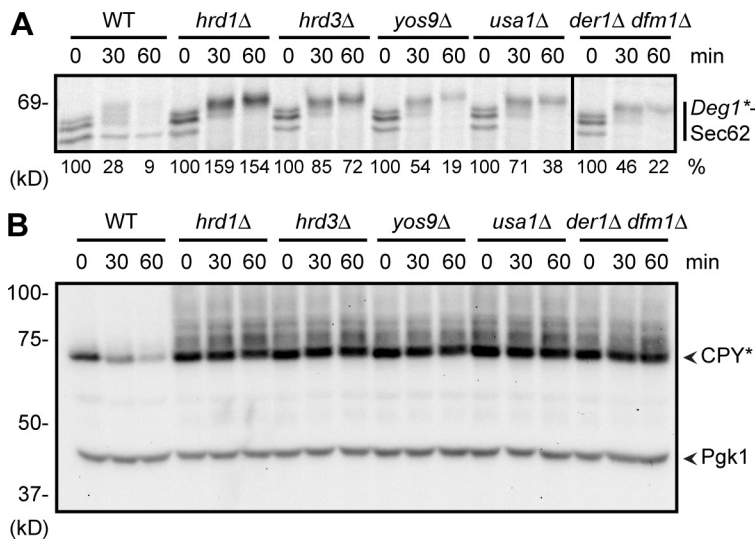


Figure 6. Hrd1 cofactor requirements for *Deg1*-*Sec62* degradation. (A) Pulse-chase analysis of *Deg1**-*Sec62* in the indicated yeast strains. The percentage of input protein remaining at each time-point is indicated below the autoradiograph. Cycloheximide was included in the chase. *Deg1**-*Sec62* was precipitated with anti-Flag antibodies. *Deg1**, F18S/I22T double mutant. The black line indicates that intervening lanes have been spliced out. (B) Cycloheximide chase analysis of CPY*-HA in the indicated yeast strains. CPY*-HA was detected by anti-HA immunoblotting. Pgk1 serves as a loading control and was detected by anti-Pgk1 immunoblotting.

*Deg1**-*Sec62* was efficiently degraded in the absence of Yos9 and the Derlins (Der1 and Dfm1), proteins with roles in ERAD-L, and only marginally stabilized in the absence of Usa1, which functions in both ERAD-L and ERAD-M (Fig. 6 A; Knop et al., 1996; Buschhorn et al., 2004; Bhamidipati et al., 2005; Kim et al., 2005; Szathmary et al., 2005; Carvalho et al., 2006; Goder et al., 2008; Horn et al., 2009; Carroll and Hampton, 2010; Carvalho et al., 2010; Kanehara et al., 2010). *Deg1**-*Sec62* was significantly, but incompletely, stabilized in the absence of Hrd3, which is required for degradation of both ERAD-L and ERAD-M substrates (Hampton et al., 1996; Carvalho et al., 2006; Gauss et al., 2006). However, this effect may be indirect, as Hrd1 levels are markedly reduced in the absence of Hrd3 (Plempner et al., 1999). A control ERAD-L substrate, CPY*-HA, was significantly stabilized in these mutant strains (Fig. 6 B). Degradation requirements for *Deg1**-*Sec62* bear some resemblance to those for the ERAD-M substrate Hmg2. However, Hmg2 degradation displays a stronger dependence on Hrd3 (Wilhovskiy et al., 2000) and potentially Usa1 (Horn et al., 2009; Carroll and Hampton, 2010) for its degradation. Together, these results suggest that recognition of *Deg1*-*Sec62* occurs through a novel mechanism requiring the minimal ERAD machinery consisting of Hrd1, Ubc7, presumably Cue1 (which anchors Ubc7 to the ER membrane), and possibly Hrd3. Novel cofactors might also be involved.

Degradation of an apolipoprotein B derivative requires Hrd1

We hypothesized that the mechanism used by Hrd1 to target *Deg1*-*Sec62* at the translocon may be related to that of a known cotranslocationally degraded protein with intriguing links to human health. Under conditions in which lipid binding of apoB (a major component of low-density and very-low-density lipoproteins) is impaired, apoB translocation is arrested and the protein remains associated with the translocon (Fisher et al., 1997; Pariyarath et al., 2001). Incompletely translocated apoB is redirected to proteasomal degradation in the cytoplasm (Yeung et al., 1996), perhaps via a mechanism similar to that of *Deg1*-*Sec62* degradation. ApoB copurifies with several

mammalian ERAD components, including HRD1, Derlin-1, and p97 (Fisher et al., 2008; Rutledge et al., 2009). Moreover, overexpression of the mammalian E3 gp78 (a HRD1 paralogue) stimulates degradation of apoB (Liang et al., 2003). Knock-down of gp78 expression decreases apoB ubiquitylation and enhances apoB secretion (Fisher et al., 2011). A model apoB derivative (apoB29) expressed in yeast copurifies with the translocon and is partially stabilized in *ubc7Δ* or *doa10Δ hrd1Δ* cells but not *ubc6Δ* cells, which is consistent with a wider role for Hrd1 in degradation of proteins persistently associated with the translocon (Hrizo et al., 2007). We therefore performed cycloheximide-chase/immunoblot analysis to test directly whether apoB29 degradation requires this E3. ApoB29 was in fact significantly stabilized in *hrd1Δ* cells (Fig. 7 A).

ApoB29 appeared as multiple bands when separated by SDS-PAGE, with a gradual conversion of lower to higher molecular weight species over time. This is most apparent under stabilizing conditions (i.e., in *hrd1Δ* cells). ApoB29 is a translocon-engaged protein with its N terminus exposed to the ER lumen. These slower migrating bands of apoB29 were sensitive to Endo H, which indicates N-glycosylation of apoB29 (Fig. 7 B). A complete collapse to the fastest migrating species was not observed, however, which suggests that apoB29, like *Deg1*-*Sec62*, acquires additional PTMs distinct from N-glycosylation. These additional PTMs became more prominent after inhibition of protein synthesis, as has previously been observed for other ERAD substrates (Wilhovskiy et al., 2000; Sato and Hampton, 2006).

Finally, we analyzed the cofactor requirements for apoB29 degradation. In contrast to *Deg1*-*Sec62*, apoB29 was markedly stabilized in cells lacking Hrd3, Yos9, Usa1, or the Derlins (Fig. 7 C). Thus, although both proteins may be targeted by Hrd1 by virtue of their aberrant or prolonged translocon engagement, they differ with respect to the exact cofactors required for their degradation.

Discussion

Many proteins pass through the Sec61 translocon en route to their ultimate subcellular or extracellular destinations.

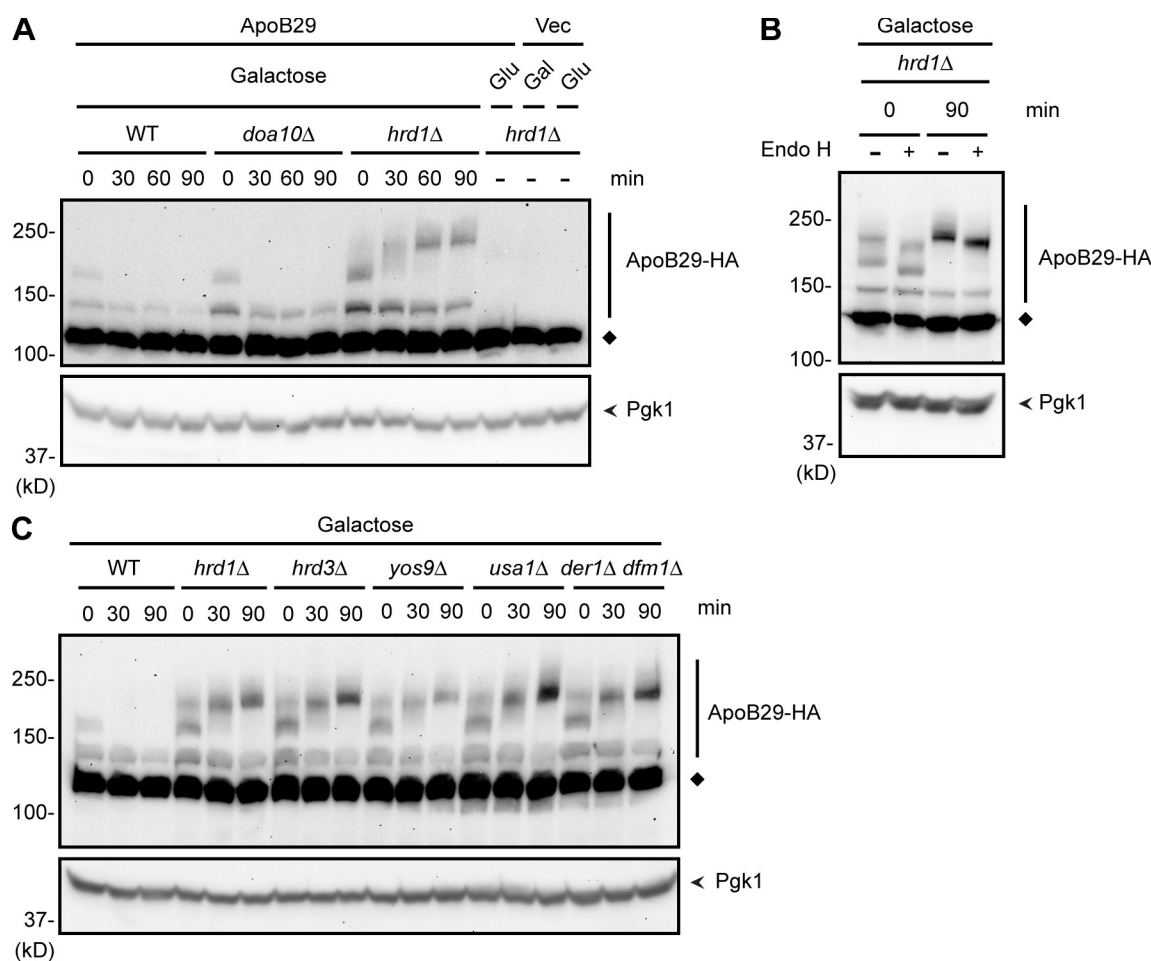


Figure 7. **ApoB29 is a Hrd1 substrate.** (A–C) Cycloheximide chase analysis of ApoB29-3HA, expressed under the control of the *GAL1* promoter in the indicated yeast strains, which were grown for 5 h in SD medium containing 2% galactose. ApoB29-3HA was detected by anti-HA immunoblotting. Pgk1 serves as a loading control and was detected by anti-Pgk1 immunoblotting. The diamonds denote a nonspecific band. (A) As controls, *hrd1Δ* yeast harboring a plasmid with *GAL1*-driven ApoB29-3HA were grown in 2% glucose (repressing) medium, and *hrd1Δ* yeast harboring an empty vector were grown in 2% glucose (repressing) or 2% galactose (inducing) medium and processed similarly. (B) Lysates prepared immediately or 90 min after cycloheximide addition were incubated in the presence or absence of Endo H.

Potentially, ERAD substrates may also pass through this channel in a retrograde fashion en route to proteasomal degradation. Aberrant, translocationally stalled proteins might limit flux through the channel in either direction, and eukaryotes may therefore have evolved mechanisms for eliminating such obstructions. Although *Deg1*-Sec62 (like many heavily studied model ERAD substrates) is an artificial protein, it may illuminate a previously unappreciated protein quality-control pathway mediated by the Hrd1 ubiquitin ligase and serve as a prototype for proteins that aberrantly engage or occupy the translocon. Indeed, we find that clearance of apoB, a protein previously reported to be cotranslocationally degraded, also depends on Hrd1.

Deg1-Sec62 inserts into the ER membrane in two distinct steps. Initially, the two TM segments of *Deg1*-Sec62 are probably cotranslationally translocated to yield the same topology as WT Sec62. In a second, *Deg1*-dependent step, a loop within the normally cytoplasmic N-terminal domain of *Deg1*-Sec62 penetrates the ER membrane. This rearrangement allows access of previously cytoplasmic residues to the N- and O-glycosylation machinery of the ER lumen.

Although glycosylation is not required for Hrd1-mediated degradation, the switch in degradation dependence from Doa10 to Hrd1 tracks very closely with acquisition of these PTMs. Our data strongly suggest that the topological rearrangement across the ER membrane is the crucial event dictating E3 specificity (Fig. 8).

Other work supports the 4-TM model of rearranged topology (Fig. 3 C). Potential O-mannosylation sites (serines) were systematically introduced throughout *Deg1*-Sec62 (Scott and Schekman, 2008). We inferred that residues that became O-mannosylated reached the ER lumen; nonmodified serine substitutions were judged to be in the membrane-spanning or cytoplasmic segments. O-mannosylation of serines inserted at positions 132 and 170 confirmed that this span is present within the ER lumen (see Fig. 1 B). Substitutions at other positions, such as immediately upstream of the first known TM, did not result in O-mannosylation. The most straightforward interpretation of these data is the 4-TM model of *Deg1*-stimulated topological rearrangement (Fig. 3 C), although the failure to observe O-mannosylation of a particular serine does not exclude its luminal residence (Lommel and Strahl, 2009).

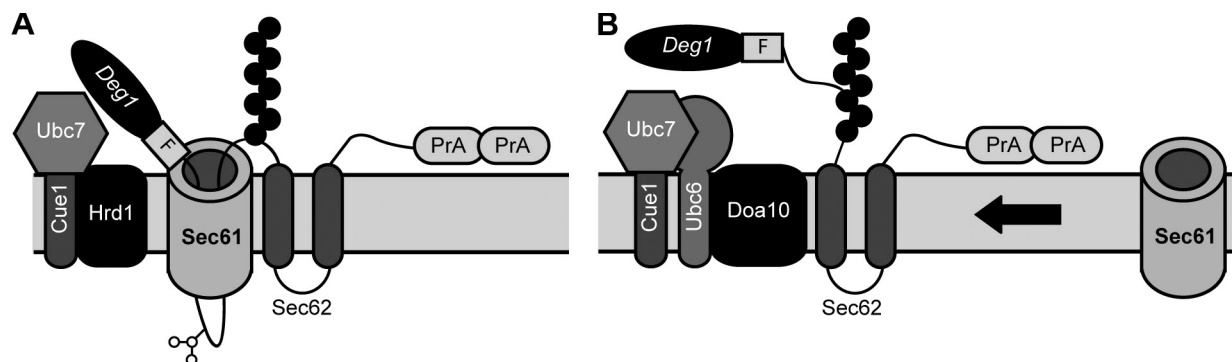


Figure 8. **Model for Hrd1-dependent ERAD of *Deg1-Sec62*.** (A) After its initial cotranslational translocation, *Deg1-Sec62* aberrantly engages the Sec61 translocon, resulting in PTT of the normally cytoplasmic N-terminal Sec62 tail. After membrane penetration, *Deg1-Sec62* becomes N-glycosylated. In this rearranged conformation, Doa10 does not recognize *Deg1-Sec62* as an ERAD-C substrate. *Deg1-Sec62* remains trapped in this orientation unless Hrd1 targets it for degradation, thereby restoring functionality to the Sec61 translocon. With the exception of the E2 Ubc7 (and presumably its membrane-anchoring binding partner Cue1; Biederer et al., 1997), and potentially Hrd3 (not depicted), well-characterized Hrd1 cofactors that function in ERAD-L and ERAD-M are dispensable for ERAD of *Deg1-Sec62*. The approximate position of N-glycosylated N153 residue is indicated with a cartoon representation of the N-glycan. (B) When translocon engagement is prevented by disrupting *Deg1-Sec62* interaction with the translocon or impairing PTT, *Deg1-Sec62* retains its original topology and is targeted by Doa10 as an ERAD-C substrate. Black circles, ubiquitin.

Topological rearrangement of *Deg1-Sec62* likely occurs via Sec61-mediated PTT. Disrupting contacts of *Deg1-Sec62* with the translocon or impairing PTT by Sec61 mutation prevents domain dislocation and significantly reverts degradation dependence to Doa10 (Figs. 4 and 5). Additionally, a cysteine found within the initially cytoplasmic N-terminal domain of *Deg1-Sec62* forms a transient disulfide linkage with a cysteine on the interior of the Sec61 channel (Scott and Schekman, 2008). This bond appears to facilitate the altered topology of the Sec62 N-terminal domain, as mutation of either participating cysteine delays *Deg1-Sec62* PTM and causes a partial switch to Doa10-dependent degradation (Fig. 5, C and D).

Fusion of *Deg1* to Sec62 was not anticipated to trigger PTT of the normally cytoplasmic Sec62 N-terminal domain. MAT α 2, from which *Deg1* is derived, is a soluble nuclear protein, and *Deg1* fused to the isolated soluble Sec62 N-domain (residues 1–149) does not trigger membrane translocation (Scott and Schekman, 2008). Rather, *Deg1-Sec62*_{1–149} is degraded in a Doa10-dependent manner, behaving like other characterized soluble *Deg1* fusions (Fig. S2 B). Potentially, fusing any protein sequence to the Sec62 N terminus could promote the observed topological rearrangement. This is not the case, however, as *Deg1* bearing a 20-residue internal deletion does not promote PTM (or degradation) when fused to Sec62 (Scott and Schekman, 2008). *Deg1*, by virtue of novel protein–protein and/or protein–membrane interactions stimulated by enforced proximity to the translocon, may conformationally alter the N-terminal portion of Sec62 or cause it to linger persistently near the opening of the Sec61 translocon channel. The translocon could respond by attempting to conduct the Sec62 N-terminal tail into the ER lumen. The disulfide linkage between *Deg1-Sec62* and the Sec61 channel may strengthen the interaction of the Sec62 N-domain with a signal sequence-binding site within the channel. Consistent with this, the cysteine cross-link is not strictly necessary for topological rearrangement but significantly accelerates it (Fig. 5, C and D; and Fig. S1 D).

Lateral release of true TM segments from the translocon into the plane of the lipid bilayer depends strongly on the hydrophobic character of the TM helices (Shao and Hegde, 2011; Zimmermann et al., 2011). The N-terminal loop of *Deg1-Sec62* does not include bona fide TM domains; thus, we speculate that the membrane-spanning polypeptide segments flanking the N-glycosylated luminal element resist lateral release from the translocon and persistently occupy the Sec61 channel. We propose that Hrd1 recognizes persistent translocon engagement by *Deg1-Sec62* after the aberrant translocation event. In this respect, *Deg1-Sec62* may resemble a translocationally stalled or arrested protein. Hrd1, by virtue of its reported association with the translocon (Schäfer and Wolf, 2009), may be ideally poised to target such incompletely translocated proteins for ubiquitin-mediated degradation. It is also possible that the Sec61-dependent topological change in *Deg1-Sec62* generates a permanently misfolded TM protein or misassembled translocon complex that is recognized by Hrd1. Nevertheless, the cofactor requirements for *Deg1-Sec62* differ from other well-characterized Hrd1 substrates (Fig. 8), which is consistent with a unique mechanism for recognition and degradation.

Deg1 has been found to destabilize other TM proteins in a partially Hrd1-dependent manner. Like *Deg1-Sec62*, both *Deg1-Hmg1* and *Deg1-Hmg2* fusion proteins are strongly stabilized in the absence of Ubc7. Deletion of *HRD1* partially impairs turnover of these proteasome substrates (Wilhovskiy et al., 2000). Whether these proteins are targeted in a mechanism similar to that targeting *Deg1-Sec62* remains to be determined.

The factors preventing Doa10 from recognizing *Deg1-Sec62* remain enigmatic. After topological rearrangement of *Deg1-Sec62*, *Deg1* remains on the cytoplasmic face of the ER membrane (Fig. 3). However, this posttranslationally translocated form of the fusion protein is resistant to Doa10-mediated proteolysis. The translocon-dependent rearrangement might alter the position or conformation of *Deg1* such that it is inaccessible to the Doa10 complex. Preventing interaction of *Deg1-Sec62* with the translocon causes a strong reversion to

Doa10-dependent degradation (Fig. 4). Therefore, *Deg1*-Sec62 is not inherently unrecognizable by Doa10. Interfering with events downstream of translocon binding (e.g., inhibiting PTT or preventing *Deg1*-Sec62–Sec61 disulfide formation) also causes a significant switch to Doa10-dependent degradation (Fig. 5). Thus, Doa10 is capable of recognizing *Deg1*-Sec62 even when conditions allow initial interaction with the translocon. In these cases, Doa10 might target *Deg1*-Sec62 that is in complex with the translocon but has not yet undergone PTT or might recognize *Deg1*-Sec62 that has transiently dissociated from Sec61.

The discovery that abnormal translocon engagement precedes *Deg1*-Sec62 degradation led us to hypothesize that cotranslocational proteolysis of apoB might occur by a similar mechanism. ApoB uses the translocon as a platform for progressive lipid binding. When lipid binding is inhibited, translocation into the ER lumen is slowed, and the protein is destroyed by the cytoplasmic proteasome (Fisher et al., 1997; Pariyarath et al., 2001). Earlier studies implied that degradation of a model variant of apoB occurred via the ERAD system in yeast (Hrizo et al., 2007). Our results directly demonstrate Hrd1 dependence for apoB29 degradation in yeast (Fig. 7 A). The cofactor requirements for *Deg1*-Sec62 and apoB Hrd1-dependent degradation, however, differed, which suggests that the details of E3-substrate recognition diverge for these substrates.

A previous study had led to the suggestion that *Deg1*-Sec62 glycosylation and disulfide formation with Sec61 occur during Sec61-dependent retrotranslocation of a Doa10 substrate (Scott and Schekman, 2008). Our results, however, strongly suggest that these events occur in the process of Hrd1 substrate generation. Given the general dependence of substrate retrotranslocation upon functional ubiquitin conjugation machinery (Biederer et al., 1997; deVirgilio et al., 1998), N-glycosylation of *Deg1*-Sec62 in *doa10Δ hrd1Δ* cells (Fig. 2 A) indicates that such modification happens upstream of substrate selection. The disulfide cross-link between *Deg1*-Sec62 and Sec61 was interpreted as representing a briefly stalled retrotranslocation intermediate, as preventing this disulfide from forming accelerated degradation. However, we found that the increased degradation rate of *Deg1*-Sec62 in the absence of the disulfide linkage is caused by a switch from Hrd1-dependent degradation to comparatively rapid Doa10-mediated proteolysis (Fig. 5, C and D). Therefore, although our data establish a role for Sec61 in the biogenesis of a Hrd1 substrate, they do not directly implicate Sec61 as a retrotranslocon for *Deg1*-Sec62, although this remains possible. It is tempting to speculate that, in a manner mirroring its membrane insertion, retrotranslocation of *Deg1*-Sec62 occurs in two steps: extraction of the aberrantly rearranged portion from the Sec61 channel followed by removal of the normal TM segments, potentially also through Sec61. Similarly, the precise mechanism by which apoB is extracted from the ER membrane remains to be elucidated. Translocationally stalled apoB might simply be extruded back into the cytoplasm via retrograde transport through the translocon in which it stalls, or it might be transferred to another retrotranslocating complex.

We speculate that Hrd1 may play an important role in removing proteins that persistently engage the translocon in a

process provisionally called ERAD-T (for “translocon associated”). ERAD-T substrates may include proteins that aberrantly stall in the translocon either because of an abnormality in the translocating protein (such as may be the case for *Deg1*-Sec62) or because of malfunction in the translocation process itself. Such substrates may be difficult to identify experimentally, as they likely represent a small subpopulation of any given translocated gene product. Artificial translocon-occupying substrates, such as *Deg1*-Sec62, may serve as a model for understanding such mechanisms. Other cellular pathways may have co-opted this quality-control pathway to mediate the regulated destruction of otherwise normal proteins. The low-density and very-low-density lipoprotein biosynthesis pathway provides a likely example of a protein—apoB—whose regulated degradation utilizes a basic cellular quality control mechanism. Future studies will seek to understand the mechanisms by which Hrd1 recognizes this potentially distinct class of substrates that persistently engage the translocon.

Materials and methods

Yeast and bacterial methods

With the exception of cells prepared for cycloheximide chase/immunoblot analysis of apoB29, yeast were grown in synthetic defined (SD) medium that was prepared as described previously (Guthrie and Fink, 1991). For experiments involving apoB29, yeast cells were cultured in modified SD medium with 5 mg/ml of acid-hydrolyzed casein and 0.08 mg/ml tryptophan as the amino acid source. Standard techniques were used for genetic manipulation of yeast strains (Guthrie and Fink, 1991). To introduce gene deletions, selection markers were PCR-amplified from donor strains with flanking sequences containing homology upstream and downstream of target gene start and stop codons. PCR products were transformed into yeast to replace genes by homologous recombination. Gene disruptions were confirmed by PCR. Standard techniques were used for plasmid manipulation in *Escherichia coli* (Ausubel et al., 1989). Site-directed mutagenesis was used to introduce point mutations (Zheng et al., 2004). Yeast strains and plasmids used in this study are presented in [Tables S1 and S2](#), respectively. Yeast strains and plasmids not constructed in our laboratory were provided by C. Andreasson (Stockholm University, Stockholm, Sweden), J. Brodsky (University of Pittsburgh, Pittsburgh, PA), E. Craig (University of Wisconsin, Madison, Madison, WI), R. Gilmore (University of Massachusetts Medical School, Boston, MA), Y. Jigami (National Institute for Advanced Industrial Science and Technology, Tsukuba, Japan), N. Johnsson (Ulm University, Ulm, Germany), D. Ng (National University of Singapore, Singapore), R. Schekman (University of California, Berkeley, Berkeley, CA), D. Scott (St. Jude Children’s Research Hospital, Memphis, TN), and T. Yoko-o (National Institute for Advanced Industrial Science and Technology, Tsukuba, Japan).

Pulse-chase analysis

Pulse-chase analysis was performed essentially as described previously (Chen et al., 1993). Yeast cells were labeled with ~20 μ Ci TRAN³⁵S-LABEL (MP Biomedicals) per 1 OD₆₀₀ unit of cells at 30°C for 10 min in SD medium lacking methionine and cysteine. Chases with excess unlabeled Met and Cys were performed in the absence of cycloheximide, except where noted in the figure legends. Inclusion of 500 μ g/ml cycloheximide had no detectable impact on degradation kinetics, modification kinetics, or E3 specificity. Immunoprecipitation of *Deg1* fusion proteins was performed with anti-FLAG M2 affinity resin (Sigma-Aldrich) or sequential incubation with anti-*Deg1* antibody (Hochstrasser and Varshavsky, 1990) and agarose cross-linked to recombinant Protein A (Repligen), as indicated in the figure legends. Immunoprecipitated proteins were separated by SDS-PAGE and analyzed by autoradiography, using a Storm 860 Phosphorimager system and ImageQuant 5.2 software (Molecular Dynamics).

Cycloheximide chase and standard immunoblot analyses

Cycloheximide was added to a final concentration of 250 μ g/ml to logarithmically growing cells. For analysis of CPY*-HA, cells were lysed essentially as described by Kushnirov (2000). 2.5 OD₆₀₀ units of cells were

collected at each time point, pelleted by centrifugation, resuspended in 400 μ l of 0.2 N NaOH, and incubated for 5 min at room temperature. Cells were pelleted by centrifugation and resuspended in sample loading buffer (Laemmli, 1970). Samples were boiled for 5 min before separation by SDS-PAGE and immunoblotting.

For analysis of apoB29, cells were lysed essentially as described by Loayza and Michaelis (1998). 2.5 OD₆₀₀ units of cells were collected at each time point, pelleted by centrifugation, and resuspended in 1 ml of ice-cold water. Cells were lysed by the addition of NaOH (to 0.26 N final concentration) and β -mercaptoethanol (to 0.13 M final concentration) and allowed to incubate on ice for 15 min. TCA was added to 5% final concentration to precipitate proteins. Proteins were pelleted by centrifugation at 4°C. Pellets were washed with 2% TCA and resuspended in 50 μ l TCA sample buffer (3.5% SDS, 0.5 M DTT, 80 mM Tris [not pH adjusted], 8 mM EDTA, 15% glycerol, and 0.1 mg/ml of bromophenol blue) before separation by SDS-PAGE and immunoblotting.

For analysis of Deg1-Sec62₁₋₁₄₉, 2.5 OD₆₀₀ units of cells were collected at each time point, pelleted by centrifugation, resuspended in 100 μ l of 0.2 N NaOH, and incubated for 5 min at room temperature. Cells were pelleted by centrifugation and resuspended in 30 μ l of urea lysis buffer (1% NP-40, 1% sodium deoxycholate, 1% SDS, and 8 M urea). Laemmli sample buffer was added and samples were boiled for 5 min before separation by SDS-PAGE and immunoblotting.

Proteins were immunoblotted to polyvinylidene fluoride membranes (Millipore), which were blocked with 5% skim milk in TBS/T (20 ml Tris, pH 7.6, 136 mM NaCl, and 0.1% Tween 20) for 60 min at room temperature or overnight at 4°C. All antibody incubations were conducted for 45–90 min at room temperature in 1% skim milk in TBS/T followed by three 5-min washes in TBS/T. The following mouse monoclonal antibodies were used at the indicated dilutions: anti-FLAG M2 (Sigma-Aldrich) at 1:10,000, anti-HA 16B12 (Covance) at 1:2,000, anti-yeast Pgk1 (Molecular Probes) between 1:20,000 and 1:40,000, and anti-yeast CPY at 1:5,000. Anti-Deg1 (MAT α 2) rabbit polyclonal antibodies were used at a dilution of 1:5,000. The anti-yeast CPY antibody is a monoclonal mouse antibody raised against yeast CPY and was a gift of N. Segev (University of Illinois at Chicago, Chicago, IL). Anti-Deg1 antibodies are polyclonal rabbit anti- α 2 antiserum raised against a full-length α 2- β -galactosidase fusion protein (Hochstrasser and Varshavsky, 1990). Mouse primary antibody incubations were followed by incubation with peroxidase-coupled sheep anti-mouse IgG (GE Healthcare) at a dilution of 1:5,000; anti-Deg1 antibody incubation was followed by incubation with peroxidase-coupled donkey anti-rabbit IgG (GE Healthcare). The peroxidase anti-peroxidase (PAP) antibody (Sigma-Aldrich) was used to detect the Protein A epitope at a dilution of 1:8,000 and required no secondary antibody. Bound antibody was visualized using Enhanced Chemiluminescence (GE Healthcare).

Endo H treatment of cellular proteins

After pulse chase, Deg1-Sec62 variants were immunoprecipitated. After five washes with wash buffer (150 mM NaCl, 50 mM Hepes, pH 7.5, 5 mM EDTA, 1% Triton X-100, and 0.1% SDS), two additional equilibrating washes were performed with 1 \times NEBuffer 4 (New England Biolabs). The resin was resuspended in 50 μ l of 1 \times NEBuffer 4 that had been supplemented with potassium acetate, pH 5.6, to an 80-mM final concentration. After adding 0.005–0.0075 U of Endo H (Roche) to the resin suspension, samples were incubated at 37°C for 2 h with gentle mixing every 20–30 min. Laemmli sample buffer was added, and samples were boiled for 8 min.

For treatment of whole cell lysates, lysates were prepared by glass-bead vortexing 3 \times 5.0 m/s for 20 s (with 5 min on ice between pulses) using a FastPrep-24 bead beating machine (MP Biomedicals) in a buffer containing 110 mM Hepes, pH 7.4, (adjusted with KOH), 2 mM MgCl₂, 0.1% Tween 20, and 1 \times Complete Protease Inhibitor Cocktail (Roche). Lysate volumes corresponding to 0.075 OD₆₀₀ units of cells were incubated with 0.005 U of Endo H (or mock-treated with water) at 37°C for 2 h. Laemmli sample buffer was added, and samples were boiled for 8 min.

Proteinase K and Endo H protection of microsomal proteins

Yeast microsomal membranes were prepared essentially as described previously (Kreft et al., 2006; Scott and Schekman, 2008). 15 OD₆₀₀ units of cells were harvested by centrifugation, resuspended in 1 ml resuspension buffer (10 mM Tris, pH 9.4, and 10 mM DTT), and incubated at room temperature for 10 min. Cell pellets were harvested by centrifugation and washed with spheroplast buffer (1 M sorbitol, 20 mM sodium

phosphate, pH 7.5, 150 mM NaCl, and 2 mM dithiothreitol). Cells were digested in 140 μ g zymolyase 100T (MP Biomedicals)/10 OD₆₀₀ units of cells in spheroplast buffer for 20 min at 30°C. Spheroplasts were harvested by centrifugation (5 min at 600 g at 4°C) and washed in spheroplast buffer containing 20 μ g/ml pepstatin A and 1 mM EDTA (for samples destined for protease protection analysis) or spheroplast buffer containing 1 \times EDTA-free Complete Protease Inhibitor Cocktail (Roche) and 1 mM EDTA (for samples destined for Endo H protection analysis). Spheroplasts were centrifuged again (5 min at 600 g at 4°C) and resuspended in fractionation buffer (200 mM D-mannitol, 20 mM sodium phosphate, pH 7.5, and 150 mM NaCl) containing appropriate protease inhibitors and lysed by vortexing in the presence of glass beads for three 30-s pulses with 1 min on ice between pulses. Unbroken cells and cellular debris were centrifuged (5 min at 600 g at 4°C), and the supernatant was used as the microsomal preparation. For Proteinase K protection experiments, microsome preparations corresponding to 3.75 OD₆₀₀ units were incubated with or without 1% Triton X-100 and treated with Proteinase K (Roche) in storage buffer (20 mM Tris, pH 7.5, 1 mM CaCl₂, and 50% glycerol) at a final concentration of 5 μ g/ml or mock-treated with storage buffer alone. Reactions were performed on ice for 30 min. Proteinase K activity was quenched with 10 mM PMSF. For Endo H protection experiments, microsome preparations corresponding to 3.75 OD₆₀₀ units were supplemented with potassium acetate, pH 5.6, to a final concentration of 80 mM. Samples were incubated with or without 1% Triton X-100 and treated with 0.004 U Endo H or mock-treated with water. Reactions were performed at 37°C for 2 h with gentle mixing every 20–30 min. Reactions were terminated with ice-cold TCA added to a final concentration of 10%.

For both Proteinase K and Endo H protection experiments, proteins were precipitated with 10% ice-cold TCA. Proteins were pelleted by centrifugation at 4°C. Pellets were washed with 2% TCA and resuspended in 50 μ l TCA sample buffer (Loayza and Michaelis, 1998). Acidic samples (which caused the bromophenol blue indicator to turn yellow) were neutralized, and samples were boiled for 8 min.

Inhibition of O-mannosylation

Yeast cells were cultured for 3 h at 30°C in the presence of 1 μ M 5-[[3-(1-phenylethoxy)-4-(2-phenylethoxy)phenyl]methylene]-4-oxo-2-thioxo-3-thiazolidineacetic acid [“Compound 5a”; Biotrend Chemicals; Orchard et al., 2004], provided by S. Strahl (University of Heidelberg, Heidelberg, Germany), a global inhibitor of O-mannosylation in *S. cerevisiae* (Strahl, S., personal communication; Arroyo et al., 2011).

Online supplemental material

Fig. S1 shows PTM of Deg1-Sec62. Fig. S2 shows Doa10-dependent degradation of Deg1 fusion proteins. Fig. S3 shows membrane topology of Deg1-Sec62, Deg1-sec62 \dagger , and Deg1-Vma12. Table S1 shows yeast strains used in this study. Table S2 shows plasmids used in this study. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.201203061/DC1>.

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