

# A Golgi rhomboid protease Rbd2 recruits Cdc48 to cleave yeast SREBP

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## Abstract

Hypoxic growth of fungi requires sterol regulatory element-binding protein (SREBP) transcription factors, and human opportunistic fungal pathogens require SREBP activation for virulence. Proteolytic release of fission yeast SREBPs from the membrane in response to low oxygen requires the Golgi membrane-anchored Dsc E3 ligase complex. Using genetic interaction arrays, we identified Rbd2 as a rhomboid family protease required for SREBP proteolytic processing. Rbd2 is an active, Golgi-localized protease that cleaves the transmembrane segment of the TatA rhomboid model substrate. Epistasis analysis revealed that the Dsc E3 ligase acts on SREBP prior to cleavage by Rbd2. Using APEX2 proximity biotinylation, we demonstrated that Rbd2 binds the AAA-ATPase Cdc48 through a C-terminal SHP box. Interestingly, SREBP cleavage required Rbd2 binding of Cdc48, consistent with Cdc48 acting to recruit ubiquitinated substrates. In support of this claim, overexpressing a Cdc48-binding mutant of Rbd2 bypassed the Cdc48 requirement for SREBP cleavage, demonstrating that Cdc48 likely plays a role in SREBP recognition. In the absence of functional Rbd2, SREBP precursor is degraded by the proteasome, indicating that Rbd2 activity controls the balance between SREBP activation and degradation.

**Keywords** Cdc48/p97; intramembrane proteolysis; rhomboid; SREBP; ubiquitin

**Subject Categories** Membrane & Intracellular Transport; Post-translational Modifications, Proteolysis & Proteomics

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## Introduction

Sterol regulatory element-binding proteins (SREBPs) play a key role in transcriptional regulation of lipid metabolism in response to intracellular levels of cholesterol and fatty acids in mammals (Espenshade & Hughes, 2007). SREBPs are synthesized as inactive

precursor proteins containing two hydrophobic transmembrane-spanning segments for membrane insertion into the endoplasmic reticulum (ER). In response to cholesterol depletion, SREBP cleavage-activating protein (SCAP) escorts SREBP via COPII vesicles to the Golgi apparatus where it is processed by two proteases: Site-1 protease (S1P; a membrane-bound subtilisin-related serine protease) cleaves in the lumen and Site-2 protease (S2P; a membrane-bound zinc metalloprotease) cleaves in the membrane. The proteolytically released N-terminal segment containing the transactivation domain and the basic helix-loop-helix leucine zipper DNA binding (bHLH-zip) domain travels to the nucleus where it activates transcription of genes encoding enzymes for the biosynthesis and uptake of cholesterol, fatty acids, and triglycerides, as well as activating its own transcription (Shao & Espenshade, 2012).

Sterol regulatory element-binding proteins are conserved across fungal phyla, including *Cryptococcus neoformans*, *Aspergillus fumigatus*, *Candida albicans* and *Schizosaccharomyces pombe* (Lane *et al*, 2001; Chang *et al*, 2007, 2009; Chun *et al*, 2007; Willger *et al*, 2008; Bien & Espenshade, 2010). In *S. pombe*, the SREBP called Sre1 controls the hypoxic response in addition to lipid homeostasis (Hughes *et al*, 2005; Todd *et al*, 2006; Bien & Espenshade, 2010). Under low oxygen conditions, Sre1 is proteolytically cleaved and the N-terminal transcription factor (Sre1N) is released from the membrane and enters the nucleus to activate hypoxic genes. Importantly, the SREBP oxygen-responsive pathway is conserved in human opportunistic pathogens, such as *C. neoformans* and *A. fumigatus* (Bien & Espenshade, 2010). SREBP is critical for pathogenic fungi to adapt and survive in hypoxic host conditions, which frequently occur at sites of fungal infections (Grahl *et al*, 2012). Consequently, SREBP is required for *C. neoformans* and *A. fumigatus* virulence, and SREBP pathway components are antifungal drug targets.

In addition to Sre1, fission yeast codes for a second SREBP, called Sre2. Sre1 and Sre2 have the same predicted topology, but Sre2 lacks the C-terminal SCAP binding domain (Hughes *et al*, 2005). Thus, Sre2 cleavage is not regulated by sterols or oxygen, but is constitutively cleaved (Stewart *et al*, 2011). Despite these differences in regulation, Sre1 and Sre2 cleavage require the same machinery (Stewart *et al*, 2011, 2012).

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Proteolytic activation of fission yeast SREBP (both Sre1 and Sre2) occurs through a mechanism different than mammalian SREBP. Proteolytic activation requires the Golgi Dsc (defective for SREBP cleavage) E3 ligase, a complex of five subunits, and the AAA-ATPase Cdc48, the yeast homolog of p97/VCP (Stewart *et al*, 2011, 2012; Lloyd *et al*, 2013). Cdc48 binds to the Dsc E3 ligase through the UBX domain of the Dsc5 subunit (Stewart *et al*, 2012). The Dsc E3 ligase complex possesses E3 ubiquitin ligase activity, and its complex architecture resembles the ER-associated degradation (ERAD) E3 ligase gp78 (Lloyd *et al*, 2013; Raychaudhuri & Espenshade, 2015). In ERAD, a membrane-associated E3 ligase core complex recognizes substrate proteins in the ER and mediates ubiquitylation (Olzmann *et al*, 2013). The Cdc48/p97 complex recognizes the poly-ubiquitin chain and extracts ubiquitylated substrates to the cytosol for proteasomal degradation. Despite their subunit architectural and functional conservation, the ERAD and Dsc E3 ligases have distinct functional outcomes. While ERAD E3 ligase machinery mediates substrate degradation, current data support a model in which the Dsc E3 ligase ubiquitylates yeast SREBP to direct its subsequent proteolytic activation.

Many unanswered questions stemming from major gaps in our knowledge of how yeast SREBP is activated need to be explored. First, which protease(s) cleaves SREBP? Subunits of the Dsc E3 ligase do not contain predicted protease domains, and S2P is absent from fission yeast (Bien & Espenshade, 2010). Given that activation of SREBP is required for fungal pathogen virulence, identification of an SREBP protease could provide a new drug target. Second, why is Cdc48 required for SREBP cleavage? Point mutants of *cdc48* fail to support yeast SREBP cleavage (Stewart *et al*, 2012). However, the Dsc5 UBX domain that recruits Cdc48 to the Dsc E3 ligase is not required for cleavage, suggesting an independent function for Cdc48 in SREBP cleavage. Finally, is yeast SREBP ubiquitylation required for cleavage? Our recent studies demonstrate that Dsc E3 ligase activity is required for Golgi localization of the Dsc E3 ligase complex (Raychaudhuri & Espenshade, 2015). Thus, mislocalization of the Dsc E3 ligase confounds the interpretation of *dsc* mutant data.

In this study, we provide answers to each of these questions. We demonstrate that *S. pombe* Rbd2 is a Golgi-resident rhomboid protease that is required for yeast SREBP cleavage activation. Kim and colleagues recently reported an initial genetic characterization of *rbd2* showing that Sre1 cleavage activation requires *rbd2* in *S. pombe*, and our results confirm these findings (Kim *et al*, 2015). Rhomboid proteases are a family of intramembrane serine proteases, whose catalytic site is buried within the membrane bilayer (Urban, 2006; Urban & Dickey, 2011). Conserved in prokaryotes and eukaryotes, rhomboid proteases regulate a wide range of divergent biological processes, such as growth factor signaling, mitochondrial membrane dynamics, protein quality control, and parasite invasion of host cells (Bergbold & Lemberg, 2013; Lemberg, 2013; Rather, 2013). Our epistasis analysis shows that the Dsc E3 ligase complex functions prior to Rbd2, suggesting that Dsc E3-mediated ubiquitylation is a prerequisite for Rbd2 function. The C-terminal, cytosolic tail of Rbd2 binds Cdc48, and disruption of Rbd2–Cdc48 binding blocks SREBP cleavage. Genetic experiments indicate that Cdc48 recruits SREBP to Rbd2 for cleavage, identifying a function for Cdc48 that is independent from its role in binding to the Dsc E3 ligase. The mammalian rhomboid RHDBL4 also binds

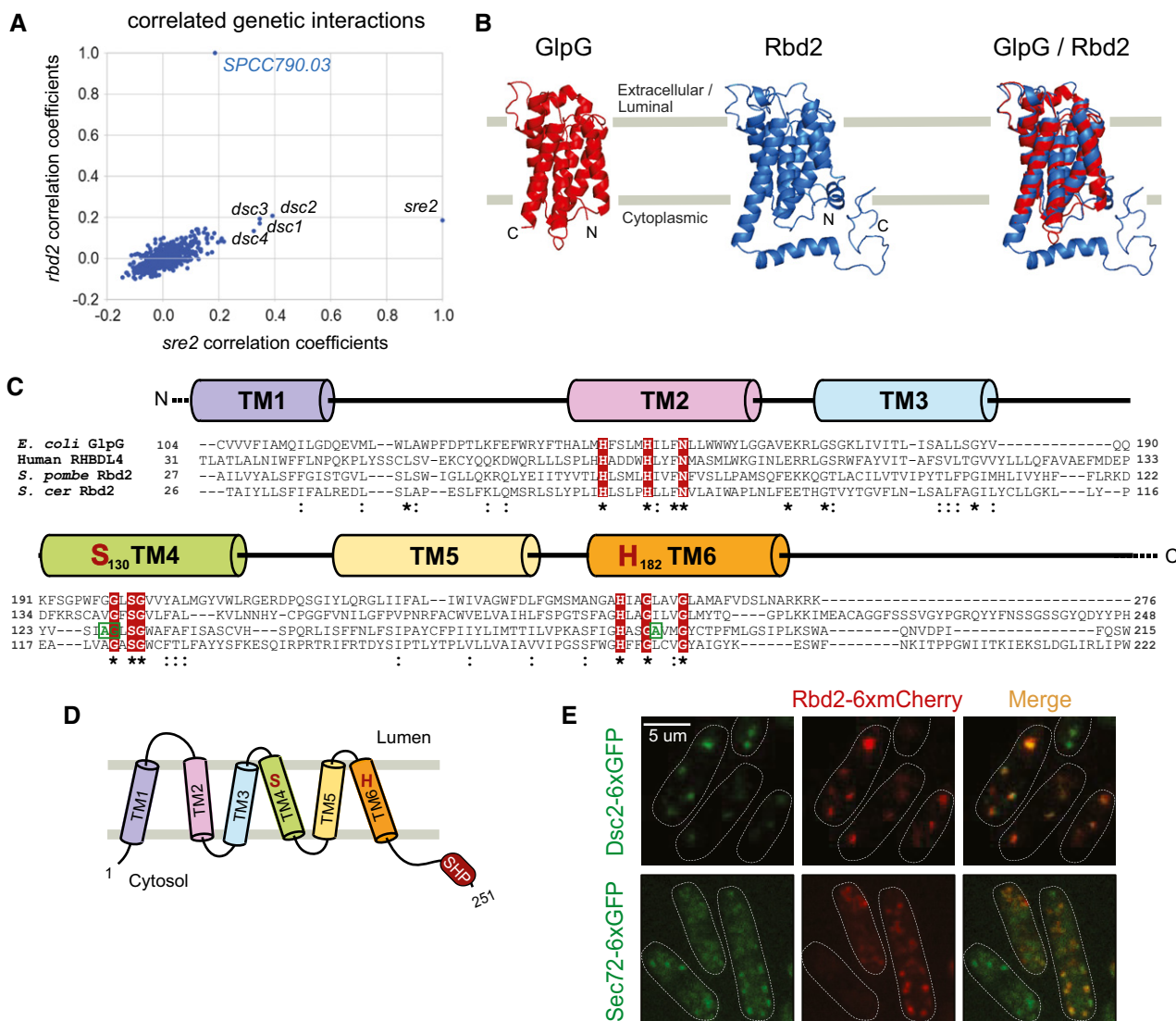
p97, but in this case, p97 functions after cleavage to promote degradation of cleaved products (Fleig *et al*, 2012). Interestingly, the absence of functional Rbd2 leads to proteasomal degradation of the uncleaved SREBP precursor form in a Dsc E3-dependent manner, supporting a model in which the Dsc E3 ligase ubiquitylates SREBP prior to cleavage by Rbd2. Thus, this study answers key questions regarding the mechanism of yeast SREBP cleavage activation.

## Results

### Genetic interaction mapping identifies function for *rbd2* in SREBP pathway

To identify additional genes involved in SREBP cleavage in *S. pombe*, we previously carried out an unbiased, high-throughput genetic interaction screen (Frost *et al*, 2012). Hierarchical clustering of genetic interaction profile correlations between a set of 1,297 G418-marked array strains and 579 nourseothricin-marked query strains revealed functionally related genes, including those involved in *dsc*-dependent SREBP pathway (Frost *et al*, 2012). A gene encoding a putative rhomboid protease, *SPCC790.03*, was correlated with *dsc1-dsc4* and *sre2* (Fig 1A). The absence of *sre1* from the cluster was likely due to the fact that the study was performed under normoxic conditions when *sre1* is inactive, an observation made in a previous study (Eisenman *et al*, 2005; Stewart *et al*, 2011).

*SPCC790.03* codes for a putative 251 amino acid (aa) rhomboid protease that is homologous to *RBD2* in *Saccharomyces cerevisiae*. Bioinformatic analysis of *S. pombe* Rbd2 revealed highly significant shared homology between Rbd2 and GlpG, a well-studied bacterial member of the rhomboid family of intramembrane proteases (Fig 1B) (Kelley *et al*, 2015). Furthermore, a multiple sequence alignment revealed that Rbd2 shares significant sequence conservation with rhomboid proteases from other kingdoms (Fig 1C). Based on known rhomboid structures and topologies, Rbd2 has six predicted transmembrane domains (TM) with the N- and C-termini in the cytoplasm (Fig 1D), and its predicted catalytic Ser-His dyad is positioned in the fourth (S130) and sixth (H182) TM, respectively. In addition, Rbd2 also contains the conserved “GxSG” motif surrounding the catalytic Ser in TM4, the “GxxxG” helix dimerization motif in TM6, and the “HxxxxHxxxN” motif in TM2 characteristic of rhomboid proteases (Fig 1C) (Urban & Baker, 2008; Urban & Shi, 2008). Interestingly, a homology search program also identified a SHP box, a putative Cdc48 interaction domain, in the cytosolic C-terminus of Rbd2 that has not been described in rhomboid proteases (Fig 1D) (Sato & Hampton, 2006; Kelley *et al*, 2015). This SHP box is not present in *S. cerevisiae* Rbd2, suggesting that the two orthologs have distinct functions. The predicted subcellular localization of Rbd2 is the Golgi apparatus (Matsuyama *et al*, 2006). Consistent with this prediction, Rbd2 fused to six tandem copies of mCherry (Rbd2-6xmCherry) appeared as puncta in cells that colocalized with two Golgi proteins, Sec72-6xGFP and Dsc2-6xGFP (Fig 1E) (Stewart *et al*, 2011). Our combined structural modeling, bioinformatic analysis, and subcellular localization data suggest that Rbd2 is a *S. pombe* Golgi-localized, rhomboid intramembrane protease.



**Figure 1. Rbd2 is a Golgi-resident rhomboid protease.**

**A** Plot of correlation coefficients generated from comparison of the genetic profiles from *sre2Δ* and *SPCC790.03Δ* (*rbd2Δ*) to all other profiles from the genetic interaction array. Note the high correlations among *dsc1-4*, *sre2*, and *rbd2*.

**B** Protein structural predictions for *Schizosaccharomyces pombe* Rbd2 (blue) and *E. coli* GlpG (red). Protein structures were predicted using the Phyre2 server (<http://www.sbg.bio.ic.ac.uk/phyre2>). Rbd2 model structure was aligned to GlpG using PyMOL.

**C** Protein sequence alignment of *E. coli* GlpG, human RHD4, *S. pombe* Rbd2, and *S. cerevisiae* Rbd2. Alignments were generated using Clustal Omega server (<http://www.ebi.ac.uk/Tools/msa/clustalo>). Predicted transmembrane domains (TM) and shared sequence motifs were used for alignment. Positions of functionally important, conserved rhomboid amino acid residues (including predicted catalytic dyad serine and histidine) are shaded in red. Residues boxed in green were identified as required for Sre1 activation from a genetic selection and were analyzed in Fig 2E and F. The starting residue numbers of the indicated sequences within their full-length proteins are given. Asterisks mark residues identical in all sequences, and colons denote conservative substitutions.

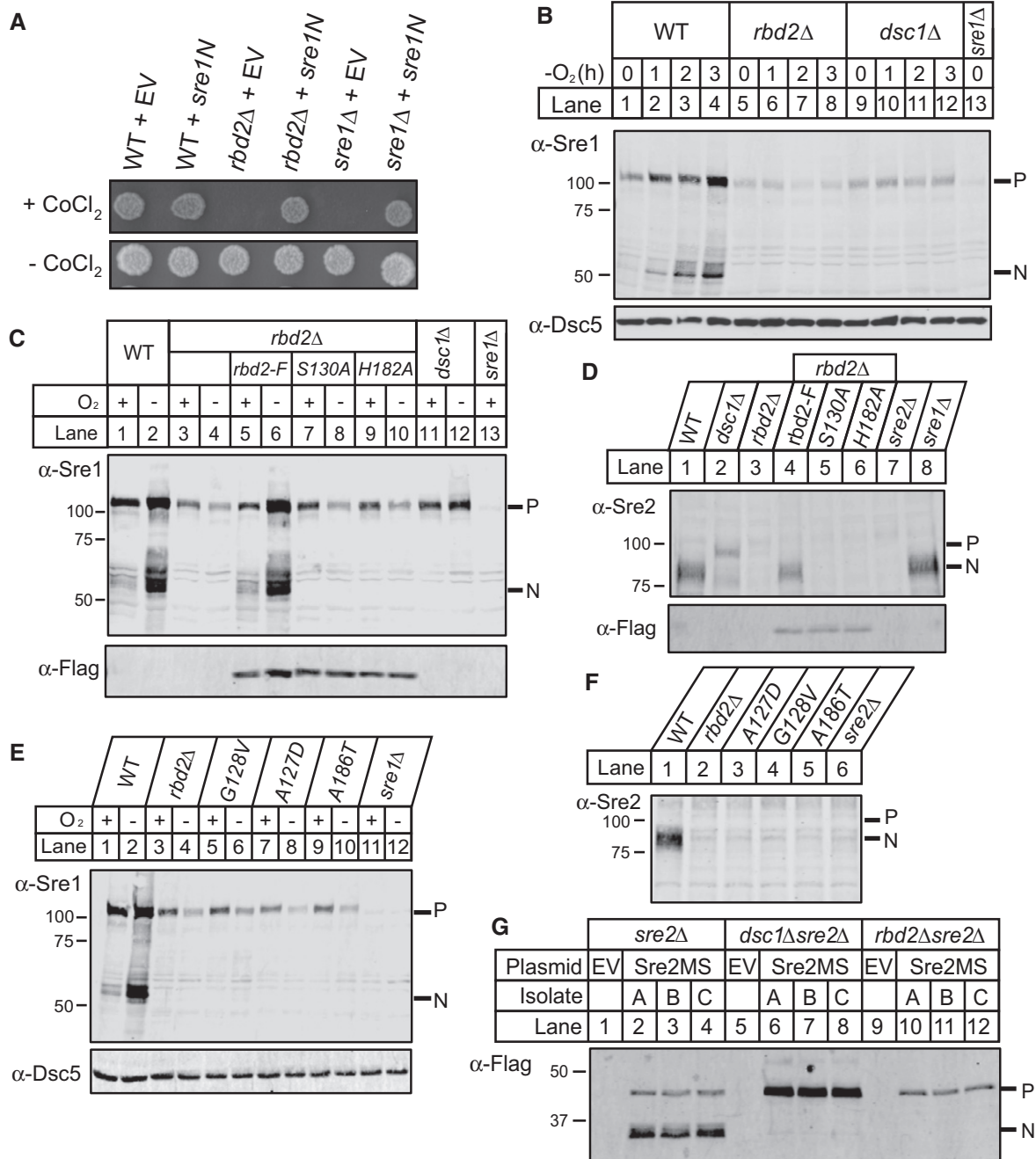
**D** Topology model for *S. pombe* Rbd2 with the conserved six-pass TM. Both the N- and C-termini face the cytoplasm. The protease active site serine and histidine form a catalytic dyad between TM4 and TM6. A SHP box, predicted to bind Cdc48, is positioned at the Rbd2 C-terminus.

**E** Cells carrying a plasmid expressing *rbd2-6xmCherry* and either of two Golgi proteins *dsc2-6xGFP* (*dsc2-6xGFP rbd2-6xmCherry*) or *Sec72-6xGFP* (*Sec72-6xGFP rbd2-6xmCherry*) were imaged by confocal fluorescence microscopy.

**SREBP activation requires Rbd2**

To test whether Rbd2 is involved in SREBP proteolytic processing, we generated an *rbd2Δ* strain. Cells lacking Sre1 activity fail to grow on medium containing cobalt chloride (CoCl<sub>2</sub>), which mimics hypoxia (Hughes *et al*, 2005). *rbd2Δ* cells failed to grow on CoCl<sub>2</sub>

medium, and growth was rescued by the expression of the cleaved, soluble N-terminus of Sre1 (Sre1N) (Fig 2A), suggesting that the *rbd2Δ* growth defect was due to a failure to produce Sre1N. Consistent with this result, *rbd2Δ* cells failed to accumulate cleaved Sre1N when cells were grown in the absence of oxygen, a condition where Sre1 cleavage was robust in wild-type cells (Fig 2B, lanes 1–8). As



**Figure 2. SREBP activation requires *rbd2*.**

A Wild-type (WT), *rbd2Δ*, or *sre1Δ* yeast (200 cells) containing either empty vector (EV) or a plasmid expressing *sre1N* (Sre1 amino acids 1–440) were grown on rich medium in the presence or absence of cobalt chloride (CoCl<sub>2</sub>).

B Western blot was probed with anti-Sre1 IgG of phosphatase-treated, whole-cell lysates from WT and the indicated mutants grown for the indicated time in the absence of oxygen. Dsc5 serves as a loading control and was detected by chemiluminescence. P and N denote Sre1 precursor and cleaved nuclear forms, respectively.

C Indicated yeast strains expressing *rbd2* or chromosomal Flag-tagged *rbd2*, *rbd2-S130A*, or *rbd2-H182A* were analyzed for Sre1 cleavage. Western blot was performed using anti-Sre1 or anti-Flag IgG of phosphatase-treated, whole-cell lysates from cells grown for 3 h in the presence or absence of oxygen.

D Indicated yeast strains expressing *rbd2* or chromosomal Flag-tagged *rbd2*, *rbd2-S130A*, or *rbd2-H182A* were analyzed for Sre2 cleavage by Western blot probed with anti-Sre2 serum or anti-Flag IgG of phosphatase-treated, whole-cell lysates from cells grown in the presence of oxygen.

E Sre1 cleavage assay was performed as in (C). *G128V*, *A127D*, and *A186T* mutant strains were obtained from our previous mutagenesis screening (Stewart et al, 2012). Dsc5 serves as a loading control.

F Sre2 cleavage assay for indicated yeast strains was performed as in (D).

G A model substrate of Sre2 containing amino acids 423–793 (Sre2MS) with an N-terminal 3xFlag epitope tag was expressed on a plasmid under the control of a CaMV promoter in *sre2Δ*, *dsc1Δsre2Δ*, or *rbd2Δsre2Δ* yeast strains. Whole-cell lysates from indicated yeast strains were analyzed by Western blot with anti-Flag antibody. Three independent isolates (A–C) are shown for each strain. Empty vector (EV) transformation serves as a negative control.

expected, *dsc1Δ* cells also failed to produce Sre1N (Stewart *et al*, 2011). Interestingly, unlike *dsc1Δ* cells, the full-length Sre1 precursor form decreased in *rbd2Δ* cells (Fig 2B, lanes 5–13). Taken together, these data suggest that deletion of *rbd2* impairs Sre1 activation under hypoxic conditions.

To test whether the catalytic dyad residues of Rbd2 (Ser130 and His182) are required for SREBP activation, we assayed Sre1 cleavage in *rbd2Δ* cells expressing chromosomally Flag-tagged *rbd2*, *rbd2-S130A*, or *rbd2-H182A*. Wild-type and Rbd2 mutants were expressed equivalently (Fig 2C, bottom panel). While wild-type Rbd2 supported Sre1 cleavage, catalytically dead Rbd2 mutants did not (Fig 2C, lanes 5–10), indicating Sre1 activation requires Rbd2 protease activity.

Next, we examined effects on activation of the second fission yeast SREBP, Sre2. In wild-type cells, Sre2 is constitutively cleaved in the presence of oxygen, and the majority of Sre2 exists as Sre2N (Fig 2D, lane 1) (Stewart *et al*, 2011). Strikingly, neither the precursor nor the cleaved Sre2N form was observed in *rbd2Δ* cells (Fig 2D, lane 3). Similar to Sre1, Sre2 activation was rescued in *rbd2Δ* cells by *rbd2*, but not by the *rbd2-S130A* or *rbd2-H182A* mutants (Fig 2D, lanes 4–6). Again, the Sre2 precursor accumulated in *dsc1Δ* cells, but was not detected in the absence of functional Rbd2 (Fig 2D).

In parallel to our genetic interaction studies, we identified point mutations of *rbd2* in a genetic selection to identify genes defective for Sre1 cleavage (Stewart *et al*, 2011). Briefly, each mutant that failed to activate an Sre1 reporter was tested for linkage to *rbd2Δ*, identifying three *rbd2*-linked mutants. DNA sequencing of *rbd2* in these mutants revealed mutation of *rbd2-A127D*, *rbd2-G128A*, and *rbd2-A186T*. These mutations reside in keystones III and IV, which are key architectural motifs conserved in all rhomboid proteases (Fig 1C, green residues) (Baker & Urban, 2012). Indeed, these *rbd2* mutants resembled *rbd2Δ* cells in Sre1 and Sre2 cleavage assays (Fig 2E and F), confirming that these residues are required for SREBP activation and independently demonstrating that SREBP activation requires Rbd2 rhomboid function.

Finally, Rbd2 was required for processing of a truncated Sre2 model substrate (Sre2MS, aa 423–793), previously shown to be processed like endogenous Sre2 (Cheung & Espenshade, 2013). Cleaved Sre2MS was not detected in *rbd2Δ* cells (Fig 2G, lanes 10–12). Unlike endogenous Sre2, a small amount of uncleaved Sre2MS was retained in the *rbd2Δ* cells, possibly due to differences in expression. Uncleaved Sre2MS accumulated in *dsc1Δ* cells as expected (Fig 2G, lanes 6–8) (Cheung & Espenshade, 2013). Taken together, these results show that Rbd2 activity is required for fission yeast SREBP activation and that precursor forms of SREBP decrease in the absence of functional Rbd2, unlike in *dsc1Δ* cells.

### SREBP precursor is degraded in a proteasome- and Dsc E3 ligase-dependent manner in the absence of *rbd2*

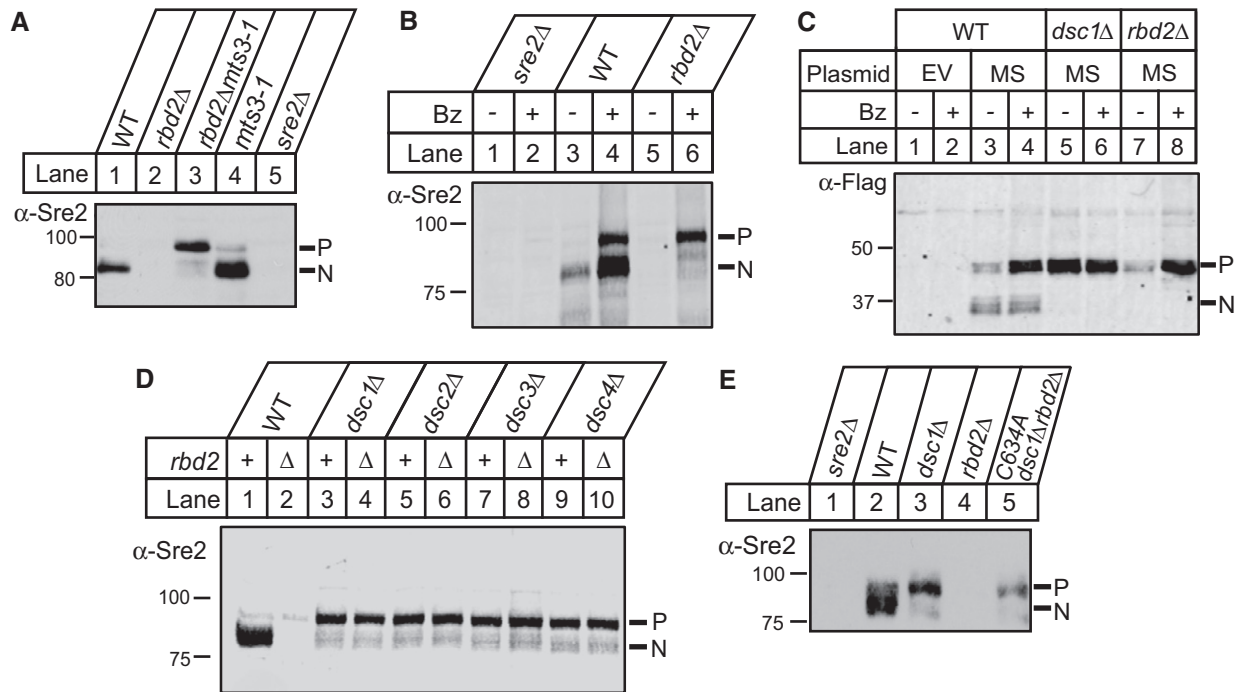
Uncleaved SREBP precursor decreased in *rbd2Δ* cells, but not in wild-type or *dsc1Δ* cells (Fig 2B–F). We next asked whether disappearance of the precursor was due to proteasomal degradation. To impair proteasome function, we used a temperature-sensitive allele of *mts3* that codes for the 19S proteasome subunit Rpn12 (Gordon *et al*, 1996). In wild-type cells, the majority of Sre2 existed as Sre2N, and we observed loss of Sre2 in *rbd2Δ* cells (Fig 3A, lanes 1–2). Inactivation of *mts3* in wild-type cells resulted in accumulation of

both Sre2P and Sre2N (Fig 3A, lane 4). Importantly, inactivation of *mts3* in *rbd2Δ mts3-1* double mutant cells showed recovery of the Sre2P but not Sre2N (Fig 3A, lane 3). Consistent with this result, treatment with the 20S proteasome inhibitor bortezomib (Bz) increased both Sre2P and Sre2N in wild-type cells, but only Sre2P accumulated in *rbd2Δ* cells (Fig 3B). Finally, the Sre2MS precursor accumulated in *rbd2Δ* mutant cells upon Bz treatment, but the cleaved form did not (Fig 3C, lanes 7–8). Bz treatment did not change the level of Sre2MS in *dsc1Δ* cells (Fig 3C, lanes 5–6). Deletion of the ERAD E3 ligases *hrd1* or *doa10* did not rescue Sre2 levels (Fig EV1A), indicating that these ER-localized E3 ligases do not target Sre2 for degradation. Taken together, these data indicate that Sre2 precursor is degraded by the proteasome in the absence of Rbd2. Furthermore, proteasome inhibition revealed that Rbd2 initiates SREBP proteolytic processing, because under these conditions, conversion of Sre2 precursor to Sre2N requires Rbd2 (Fig 3A, lanes 3–4 and Fig 3B, lanes 4 and 6).

Proteasome inhibition did not increase Sre2MS in *dsc1Δ* cells (Fig 3C, lanes 5–6), suggesting that the Golgi-localized Dsc E3 ligase may be required for SREBP degradation. Both the Dsc E3 ubiquitin ligase and Rbd2 are required for SREBP activation, and we performed epistasis tests to determine the order of action of these two pathway components. Sre2 decreased in *rbd2Δ* cells and the precursor accumulated in *dsc1Δ-dsc4Δ* cells (Fig 3D, lanes 1–3, 5, 7, 9). Deletion of *dsc1* in *rbd2Δ* cells restored Sre2 precursor levels to that of the single *dsc1Δ* mutant (Fig 3D, compare lanes 3 and 4). Identical results were seen for deletion of *dsc2-dsc4* (Fig 3D, lanes 5–10). Likewise, Sre1 precursor disappeared under low oxygen in *rbd2Δ* cells, but was recovered in *dsc1Δrbd2Δ* cells (Fig EV1B). Thus, the Dsc E3 ligase is required for proteasome-dependent degradation of SREBP in *rbd2Δ* cells, demonstrating that the Dsc E3 ligase acts prior to Rbd2 in the biochemical pathway to cleave SREBP. To test specifically whether Sre2 precursor degradation requires Dsc E3 ligase activity, we studied a previously characterized, catalytically dead mutant of Dsc1 with a mutation in the Dsc1 RING domain (*dsc1-C634A*) (Raychaudhuri & Espenshade, 2015). As observed for the *dsc1* deletion, mutation of the Dsc1 RING domain restored Sre2 precursor in *rbd2Δ* cells (Fig 3E), showing that the Sre2 precursor degradation requires Dsc RING E3 ligase activity. Collectively, these results suggest that uncleaved SREBP is ubiquitinated by the Dsc E3 ligase and degraded by the proteasome when Rbd2 function is impaired.

### Rbd2 is an active rhomboid intramembrane protease

We next used a heterologous assay with a panel of model rhomboid substrates to examine whether Rbd2 is able to catalyze intramembrane proteolysis directly. This assay relies on co-transfecting human HEK293T cells with a 3xHA-tagged rhomboid and a GFP-tagged substrate and using Western analysis to detect proteolysis. Although this approach has been used to study the protease activity of rhomboid proteases from bacteria, protozoa, plants, and animals, it has never been applied successfully to study yeast rhomboid activity. Transfecting cells with *3xHA-rbd2* resulted in low levels of protein expression (Fig 4A, lane 2). To circumvent this obstacle, we recoded the entire *rbd2* ORF to codon-optimize it for expression in human cells, and this resulted in > 35-fold higher expression of 3xHA-Rbd2 (Fig 4A, lanes 2–3). Despite achieving this robust level



**Figure 3. Uncleaved SREBP is degraded in a proteasome-dependent and Dsc E3 ligase-dependent manner in the absence of *rbd2*.**

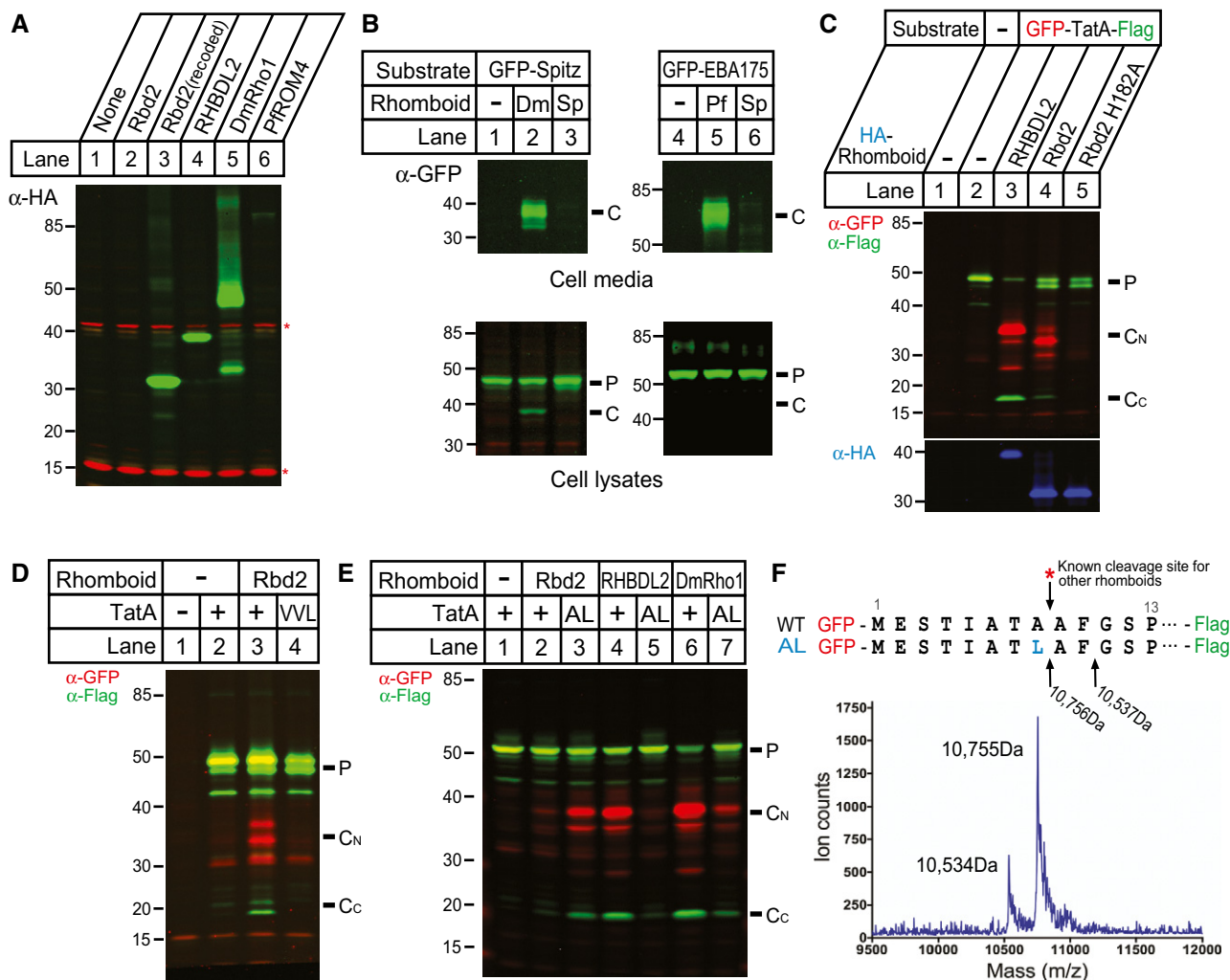
A Wild-type (WT) and the indicated mutant strains were grown for 3 h at the non-permissive temperature (36°C) to inactivate Mts3. Phosphatase-treated, whole-cell lysates were analyzed by Western blot with anti-Sre2 serum and imaged using chemiluminescence. P and N denote Sre2 precursor and cleaved nuclear forms, respectively.  
 B WT, *rbd2Δ*, and *sre2Δ* cells were treated with bortezomib (Bz) for 3 h, and phosphatase-treated, whole-cell lysates were analyzed by Western blot with anti-Sre2 serum.  
 C WT, *dsc1Δ*, and *rbd2Δ* cells carrying either empty vector (EV) or a plasmid expressing 3xFlag-Sre2MS (MS) were treated with bortezomib (Bz) for 3 h. Whole-cell lysates were analyzed by Western blot with anti-Flag antibody.  
 D Whole-cell lysates from the indicated strains were analyzed by Western blot with anti-Sre2 serum.  
 E Catalytically dead Dsc1 E3 ligase (*dsc1-C634A*) mutant and other indicated yeast strains were analyzed by Western blot probed with anti-Sre2 serum and imaged using chemiluminescence.

of expression (higher than other rhomboid proteases that we used as positive controls), Rbd2 was completely unable to cleave either the *Drosophila* signaling molecule Spitz, which is a canonical rhomboid substrate, or *Plasmodium* EBA-175, which is an “atypical” substrate that is cleaved by many rhomboid proteases that are unable to cleave Spitz (Fig 4B) (Urban & Freeman, 2003; Baker *et al*, 2006; Baxt *et al*, 2008).

We therefore next examined cleavage of a third substrate, TatA from the bacterium *Providencia stuartii*, which is perhaps the most efficient rhomboid substrate known (Moin & Urban, 2012). Co-transfecting *rbd2* resulted in consistent but weak TatA cleavage (Fig 4C, lane 4). To ensure that this was proteolysis, we tagged both termini of TatA and followed processing by Rbd2 or the human rhomboid protease RHBDL2 as a positive control. In each case, we detected both proteolytic fragments, and TatA proteolysis required the predicted catalytic histidine of Rbd2 (Fig 4C, lanes 4–5).

We next examined the substrate requirements for Rbd2 processing of TatA. Rhomboid proteases require helix-destabilizing residues in the substrate transmembrane segment, and mutating these residues of TatA (GSP, aa 11–13) to VVL abolished processing by Rbd2 (Fig 4D). All rhomboid proteases studied to date cleave after

small residues (glycine, alanine, serine) (Urban, 2010), and all rhomboid proteases tested cleave TatA after A8 (Strisovsky *et al*, 2009). We therefore tested the ability of Rbd2 to cleave TatA harboring the A8L mutant. To our surprise, we found that Rbd2 cleaved the A8L mutant substrate four-fold more efficiently than wild-type TatA (Fig 4E, lanes 2–3). However, this mutation nearly abolished processing by other rhomboid proteases (Fig 4E, lanes 4–7). Rhomboid proteases typically have only limited sequence dependence, and thus, mutations can increase proteolysis by shifting the site of cleavage to a new location. To examine whether the A8L mutant was truly being cleaved after a large hydrophobic residue, we immunopurified the C-terminal cleavage product and determined its size by mass spectrometry (Fig 4F). The predominant cleavage of the TatA A8L mutant was indeed after the leucine at position 8. Interestingly, we discovered a second, minor cleavage site (~20%) after F10, two residues deeper into the TatA transmembrane segment. Notably, this cleavage also occurred between a large and small residue, namely a naturally occurring phenylalanine and glycine. As such, Rbd2 is an active rhomboid intramembrane protease that can cleave after large hydrophobic residues. This unique property has not been observed for any rhomboid protease.



**Figure 4. Rbd2 is an active rhomboid protease.**

**A** Human HEK293T cells were transfected with plasmids expressing different 3xHA-tagged rhomboid proteases: *Schizosaccharomyces pombe* rbd2, *S. pombe* rbd2 recoded for human expression, human RHBDL2, *D. melanogaster* Rho1, and *P. falciparum* ROM4. Whole-cell lysates were analyzed by Western blot to detect the HA-tagged rhomboid. Asterisks indicate loading control bands detected with antibody ab179726 (Abcam).

**B** Human HEK293T cells were transfected with plasmids expressing the indicated rhomboid substrates and either recoded *S. pombe* rbd2, *D. melanogaster* Rho1, or *P. falciparum* ROM4. Conditioned media and cell lysates were analyzed by Western blot. P and C denote precursor and cleaved products, respectively.

**C** Human HEK293T cells were transfected with plasmids expressing the GFP-TatA-Flag substrate and the indicated rhomboid proteases. Whole-cell lysates were analyzed by Western blot. P, C<sub>N</sub>, C<sub>C</sub> denote precursor, N-terminal cleaved, and C-terminal cleaved products, respectively.

**D** Human HEK293T cells were transfected with plasmids expressing the indicated GFP-TatA-Flag substrates and in the absence or presence of recoded *S. pombe* rbd2. Whole-cell lysates were analyzed by Western blot.

**E** Human HEK293T cells were transfected with plasmids expressing the indicated GFP-TatA-Flag substrates and different rhomboid proteases. Whole-cell lysates were analyzed by Western blot.

**F** Human HEK293T cells were transfected with plasmids expressing the GFP-TatA (A8L)-Flag and recoded *S. pombe* rbd2. C-terminal cleavage products were purified using anti-Flag antibodies, and purified protein was analyzed by mass spectrometry.

### Rbd2 cleavage requires Lys743 in Sre2 luminal loop

Rhomboid proteases cleave type-I transmembrane substrates (Urban & Freeman, 2003; Ha, 2009) and consistent with this Rbd2 cleaved TatA within its transmembrane segment toward its N-terminus (Fig 4F). We next asked whether this unique substrate sequence preference was also observed for SREBP. Heterologous Rbd2 cleavage assays using multiple *S. pombe* SREBP substrate variants failed, possibly due to the lack of Dsc E3 ligase machinery in HEK293 cells.

To test directly the requirement for large hydrophobic residues, we assayed cleavage of alanine-substitution mutants in the N-terminus of the second predicted transmembrane segment of Sre2 (aa 748–754) (Fig 5A). Processing of each mutant (I748A/Y749A, L750A, F752A, L754A) was normal in wild-type cells and required both *rbd2* and *dsc1* (Fig 5B and Appendix Fig S1A), indicating that Sre2MS cleavage does not require these individual large hydrophobic residues. However, we cannot exclude the possibility that one or several of these mutations block cleavage at the wild-type site but

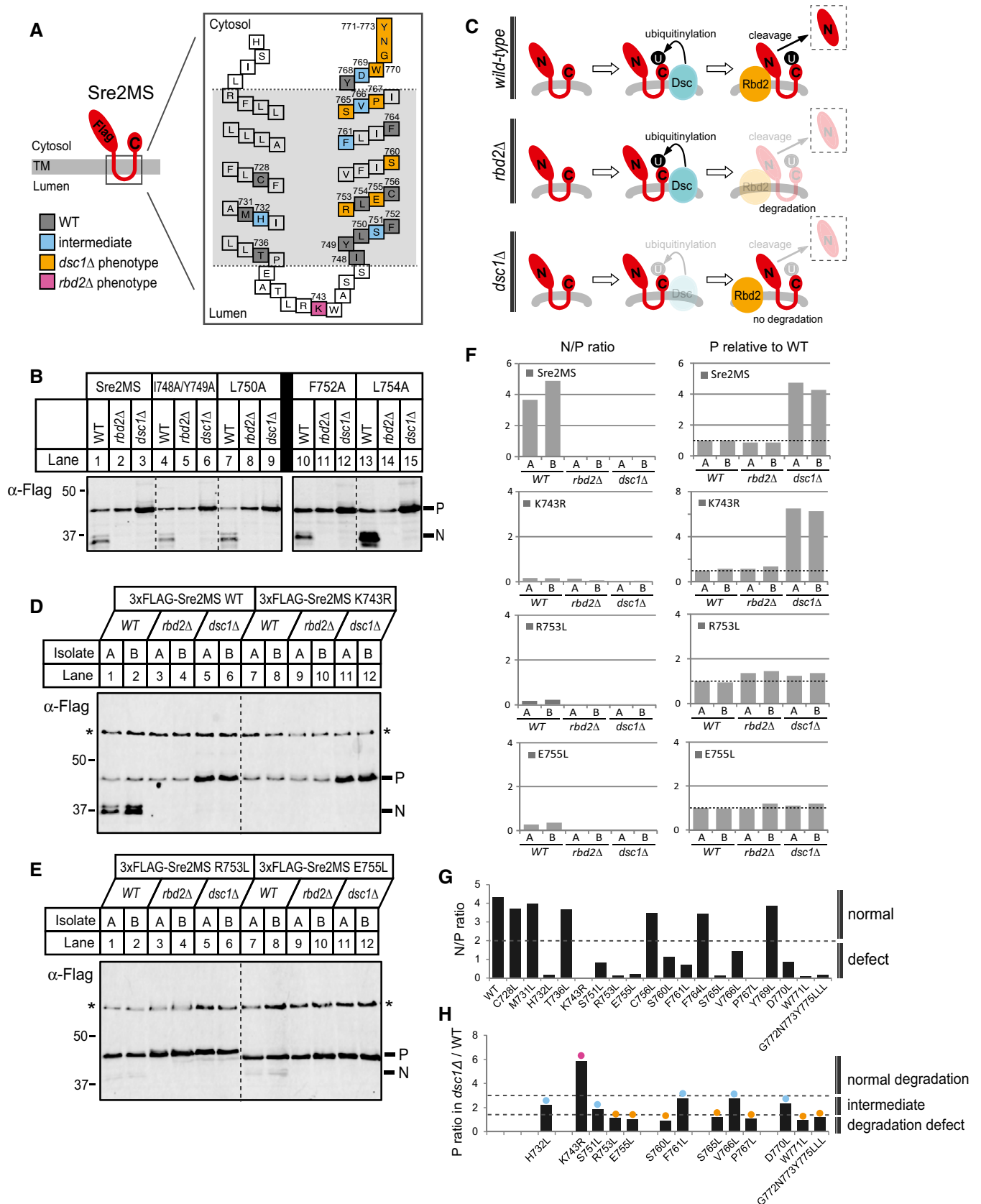


Figure 5.



**Figure 5. Rbd2 cleavage requires Lys 743.**

- A Diagram of Sre2MS substrate (aa 423–793). The enlarged model shows predicted positions of TM1 and TM2 denoting the 26 amino acids tested for mutagenesis by colored squares. Numbers indicate the number of amino acid residues. Mutants showing a block in cleavage due to defects in Dsc1 function are indicated in orange, Rbd2 function in pink, a combination of defects in Dsc1 and Rbd2 in blue (intermediate), and residues showing normal cleavage are indicated in gray.
- B Whole-cell lysates from WT, *rbd2Δ*, and *dsc1Δ* cells carrying a plasmid expressing 3xFlag-Sre2MS WT or indicated mutants were analyzed by Western blot with anti-Flag antibody. P and N denote Sre2MS precursor and cleaved forms, respectively.
- C Model outlining the predicted fate of Sre2MS in different yeast strains. U denotes either mono-ubiquitin or poly-ubiquitin.
- D, E Whole-cell lysates from WT, *rbd2Δ*, and *dsc1Δ* cells carrying a plasmid expressing 3xFlag-Sre2MS WT or indicated mutants were analyzed by Western blot with anti-Flag antibody. P and N denote Sre2MS precursor and cleaved forms, respectively. Asterisks indicate non-specific, loading control bands, which were used to normalize the intensities of P and N in (F). Two independent isolates, A and B, are shown for each strain.
- F Intensities of individual P and N bands from Western blots were quantified using LI-COR software. Relative intensity of P and N to corresponding non-specific band (asterisk) in each lane was normalized to that of P level in isolate A of WT strain for each mutagenesis. The ratio of N to P or relative P level was presented in bar graphs.
- G The average ratio of N to P level for different Sre2MS mutants in wild-type cells (Appendix Fig S2) was plotted. Cleavage was scored as normal if the ratio was > 2.
- H For Sre2MS mutants showing defects in cleavage, the average fold-increase in P level in *dsc1Δ* cells compared to WT cells from two isolates (Appendix Fig S2) was plotted. Mutants were grouped into three phenotypic classes (intermediate, *dsc1Δ*, and *rbd2Δ*) according to observed precursor accumulation. Precursor accumulation was defined as *dsc1Δ* if < 1.5 (orange), *rbd2Δ* if > 3 (pink), and intermediate if in between (blue).

induce a compensating shift in cleavage site, as has been observed in other rhomboid substrates (Strisovsky *et al*, 2009; Moin & Urban, 2012).

To identify residues important for proteolytic cleavage, we tested 19 additional Sre2MS mutants. We quantified cleavage of each mutant in three different backgrounds: wild type, *rbd2Δ*, and *dsc1Δ* (Fig 5C). Sre2MS precursor (P) was processed to a cleaved product (N) in wild-type cells (Fig 5D, lanes 1–2), and the average N/P ratio was 4–5 (Fig 5F). Unlike for endogenous Sre2 (Fig 2D), we consistently observed unprocessed Sre2MS precursor in wild-type cells, likely due to higher protein expression. In *rbd2Δ* cells, Sre2MS precursor was not cleaved to the N form (Fig 5D, lanes 3–4), and the precursor did not accumulate due to Dsc-dependent proteasomal degradation (Fig 5C). In contrast in *dsc1Δ* cells, Sre2MS precursor accumulated ~4.5-fold compared to wild-type cells (Fig 5D and F), because it is neither cleaved nor degraded due to lack of Dsc E3 ligase function (Fig 5C).

Overall, we observed cleavage defects for 13 of the 19 mutants tested (Fig 5G and Appendix Figs S1 and S2). Since Sre2MS precursor has different fates in *rbd2Δ* and *dsc1Δ* cells (Fig 5C), quantification of mutant Sre2MS precursor levels allowed assignment of the step defective in processing to either Dsc-dependent ubiquitinylation or Rbd2 cleavage. If a Sre2MS mutant is not cleaved due to a failure in Dsc1 ubiquitinylation, we expect that the Sre2MS precursor will not accumulate further in *dsc1Δ* cells and that the *dsc1Δ*/wild-type ratio will equal 1, as seen for the R753L and E755L mutants (Fig 5E and F). In contrast, if mutant Sre2MS precursor is ubiquitinated by the Dsc E3 ligase but defective for Rbd2 cleavage, we expect the precursor to accumulate in *dsc1Δ* cells compared to wild-type cells, as seen for the K743R mutant (Fig 5D, lanes 7–12). To categorize the mutants, we plotted the ratio of Sre2MS precursor in *dsc1Δ*/wild-type cells (Fig 5F and H, and Appendix Fig S2). Many amino acid residues appeared to be required for Dsc E3 activity (Fig 5A and H, orange) or for both Dsc and Rbd2 activity (Fig 5A and H, blue). No transmembrane residues were specifically required for Rbd2 activity, but mutation of lysine 743 in the luminal loop to arginine prevented Rbd2-mediated cleavage (Fig 5A and H, pink). It is not clear whether the lysine is essential for substrate recognition or whether introducing arginine at position 743 interferes with cleavage. This extended mutagenesis study demonstrates that Sre2MS cleavage does not require large hydrophobic residues in the

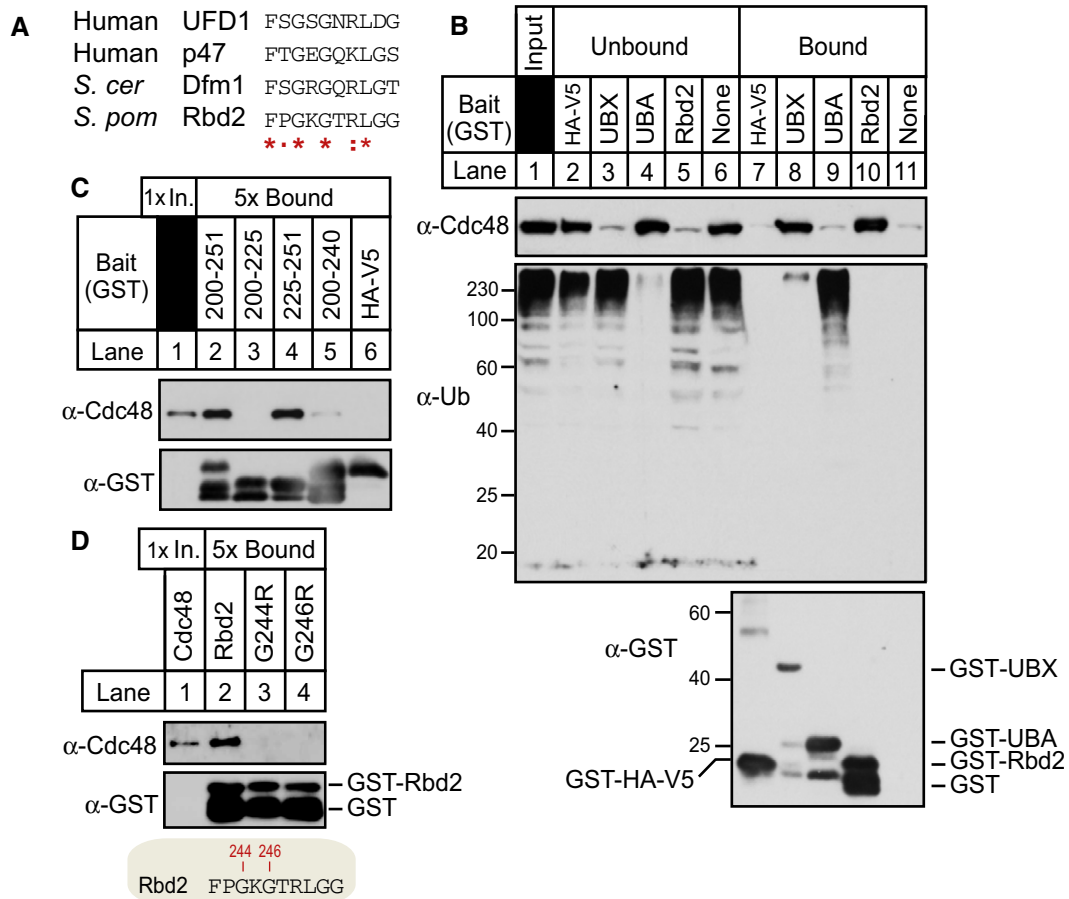
second transmembrane segment, but identifies a luminal lysine residue required for cleavage.

**Rbd2 binds Cdc48 through the C-terminal SHP box**

Bioinformatic analysis identified a putative Cdc48-interaction motif called a SHP box in the cytosolic C-terminus of Rbd2 (Fig 1D) (Sato & Hampton, 2006). SHP box motifs are present in Cdc48/p97 cofactors such as mammalian UFD1, p47, and yeast Dfm1 (Fig 6A). To test whether Rbd2 binds Cdc48, the Rbd2 C-terminus (aa 200–251) fused to GST was affinity purified from bacterial lysates and mixed with *S. pombe* cytosol. After washing, the bound fraction was analyzed for the presence of Cdc48 (Fig 6B). GST-Rbd2 (aa 200–251) bound Cdc48 to the same extent as the Dsc5 UBX domain, which is a known Cdc48-binding domain (Fig 6B, compare lanes 8 and 10) (Stewart *et al*, 2012). Dsc2 UBA domain served as a negative control that binds ubiquitin but not Cdc48 (Fig 6B, compare lanes 9–10) (Lloyd *et al*, 2013). Truncations of GST-Rbd2 that lacked the SHP box (Rbd2<sub>200–225</sub> or Rbd2<sub>200–240</sub>) failed to bind Cdc48, while a truncation containing the SHP box (Rbd2<sub>225–251</sub>) binds Cdc48 (Fig 6C). Mutational analysis of the Rbd2 SHP box demonstrated that at least two conserved residues, G244 and G246, are critical for Cdc48 binding (Fig 6D). These *in vitro* experiments established that the Rbd2 C-terminal SHP box binds Cdc48.

**APEX2 system detects Rbd2–Cdc48 interaction in cells**

To test whether Rbd2 and Cdc48 interact in cells, we performed an immunoprecipitation experiment. Flag-Rbd2 immunoprecipitated endogenous Cdc48 from yeast cells, while mutation of the SHP box (Rbd2-G246R) decreased Cdc48 binding (Fig 7A, lanes 4–6). As an independent confirmation of this interaction, we employed the APEX2 system (Rhee *et al*, 2013; Hung *et al*, 2016). Briefly, APEX2 is an engineered ascorbate peroxidase that generates biotin-phenoxy radicals upon addition of biotin-phenol and hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), resulting in covalent biotinylation of binding partners within a small radius (< 20 nm) (Fig 7B). After cell lysis, proteins labeled with biotin can be isolated with streptavidin beads for identification. We recently optimized the APEX2 system in yeast (Hwang & Espenshade, 2016). To label proximal and interacting proteins of Rbd2 using APEX2 technology, we fused Flag-APEX2 to



**Figure 6. Rbd2 binds Cdc48 through a C-terminal SHP box.**

**A** Alignment of SHP box sequences from human UFD1, human p47, *S. cerevisiae* Dfm1, and *Schizosaccharomyces pombe* Rbd2 C-terminal SHP box (aa 242–251). Asterisks denote residues identical in all sequences, colons mark conservative substitutions, and dots mark semi-conservative substitutions.

**B** Recombinant proteins GST-fused Rbd2 C-terminus (Rbd2<sub>200–251</sub>), Dsc5 UBX (Dsc5<sub>323–425</sub>), Dsc2 UBA (Dsc2<sub>298–372</sub>), and GST-HA-V5 control were bound to GST magnetic beads and incubated with *S. pombe* cytosol fraction from wild-type cells. Equivalent amounts of unbound and bound fractions were probed for anti-Cdc48, anti-ubiquitin, and anti-GST IgG.

**C** GST pull-down assay was performed as in (B) using truncated forms of Rbd2 C-terminus; Rbd2<sub>200–251</sub>, Rbd2<sub>200–225</sub>, Rbd2<sub>225–251</sub>, and Rbd2<sub>200–240</sub>. Unbound (1×) and bound (5×) fractions were analyzed by Western blotting

**D** Single residue mutants of conserved glycine residues in Rbd2<sub>200–251</sub> (G244R and G246R) were tested for Cdc48 binding as in (C).

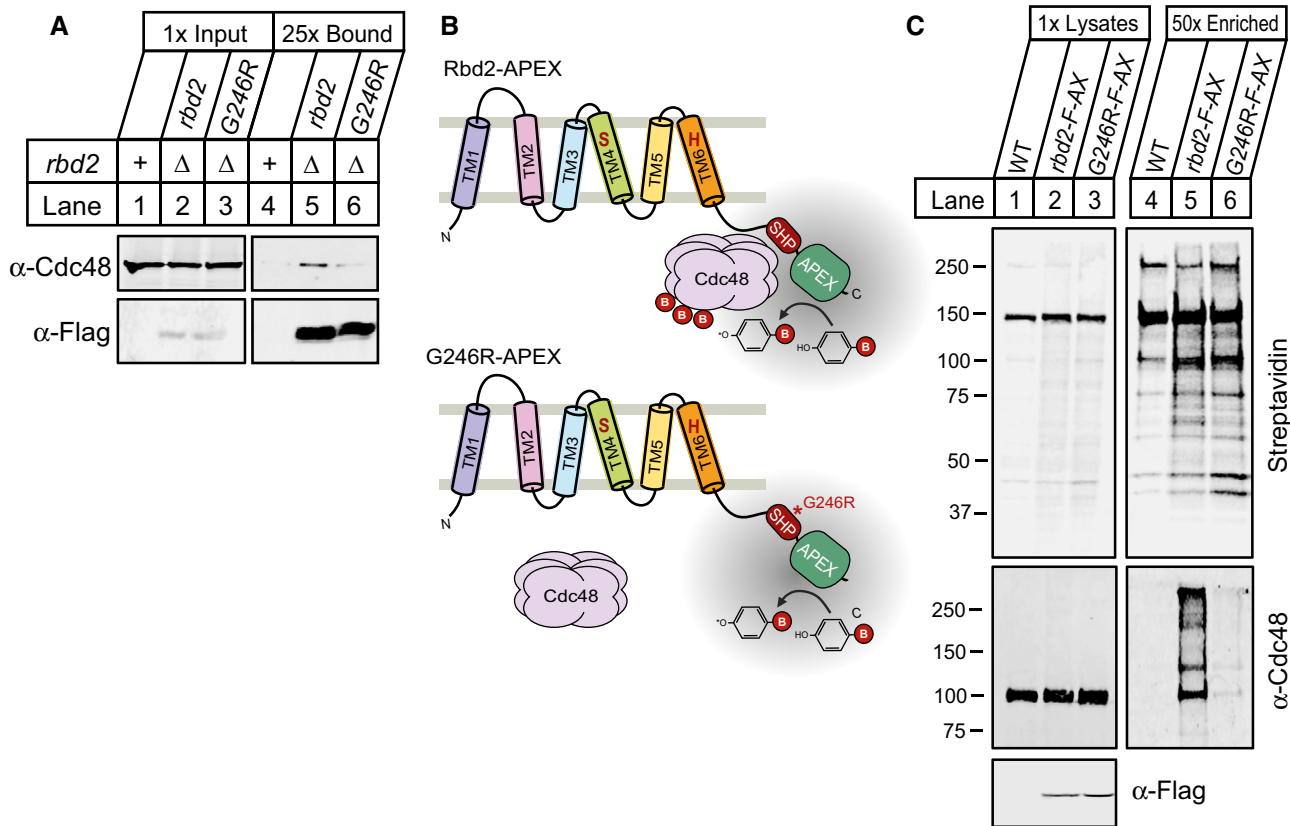
Data information: Western blots (B–D) were images using chemiluminescence.

the C-terminus of Rbd2 and overexpressed the fusion protein from a plasmid in *rbd2Δ* cells. Rbd2-Flag-APEX2 supported Sre1 cleavage demonstrating that the fusion protein was functional (Fig EV2A). Wild-type cells or *rbd2Δ* cells expressing Rbd2-Flag-APEX2 or Rbd2-G246R-Flag-APEX2 were subjected to biotin-phenol labeling for 1 min, biotinylated proteins were recovered using streptavidin beads, and bound proteins were visualized by Western blotting. Compared to wild-type cells which contain multiple naturally biotinylated proteins, cells expressing Rbd2-Flag-APEX2 and G246R-Flag-APEX2 had additional labeled proteins (Fig 7C, lanes 2–3 and 5–6) (Koffas *et al*, 1998). Western blotting of these samples revealed that Cdc48 was specifically labeled in Rbd2-Flag-APEX2 cells (Fig 7C, bottom panels). Mutation of the SHP box prevented labeling of Cdc48, confirming that the SHP box mediates Rbd2 binding to Cdc48. In contrast, the catalytically inactive Rbd2-S130A mutant still binds Cdc48 (Fig EV2B). Collectively, these *in vitro* and *in vivo*

experiments demonstrate that the SHP box mediates Cdc48 binding to Rbd2.

**SREBP activation requires Rbd2 binding to Cdc48**

To test whether SREBP activation requires Cdc48 binding to Rbd2, we used four different assays with both Rbd2 SHP box mutants (G244R and G246R): (i) growth on medium containing the hypoxia mimetic CoCl<sub>2</sub> (Fig 8A), (ii) Sre1 cleavage assay in the absence of oxygen (Fig 8B), (iii) Sre2 cleavage assay (Fig 8C), and (iv) Sre2MS cleavage assay (Fig 8D). *rbd2-G244R* cells exhibited slow growth on CoCl<sub>2</sub>, reduced Sre1 cleavage, and decreased Sre2N, consistent with partial loss of Rbd2 function. *rbd2-G246R* cells showed more severe defects and phenocopied the catalytic mutant Rbd2-S130A, demonstrating that SREBP activation requires a functional SHP box.



**Figure 7. APEX2 system detects Rbd2–Cdc48 binding in cells.**

- A Detergent-solubilized yeast extracts from cells containing chromosomal Flag-tagged *rbd2* or *rbd2*-*G246R* in *rbd2*Δ background were prepared, and proteins associated with Rbd2-Flag were immunopurified using anti-Flag magnetic beads. Equal amount of total proteins (lanes 1–3) along with 25× bound fractions (lanes 4–6) were analyzed by Western blot with anti-Cdc48 serum and anti-Flag IgG.
- B Illustration outlining APEX2-catalyzed biotin-phenol labeling. An Rbd2-Flag-APEX fusion protein oxidizes biotin-phenol to biotin-phenoxyl radical that covalently labels binding partners.
- C Biotin-phenol labeling was performed for 1 min with H<sub>2</sub>O<sub>2</sub> treatment as described in Materials and Methods. WT cells or *rbd2*Δ cells carrying *rbd2*-Flag-APEX2 (*rbd2*-F-AX) or *rbd2*-*G246R*-Flag-APEX2 (*G246R*-F-AX) plasmid were lysed after biotin-labeling reaction, and proteins were denatured by heating the cells in lysis buffer containing 1% SDS. Biotinylated proteins were then enriched using streptavidin magnetic beads. Lysates and 50×-enriched eluates were analyzed by Western blot with IRDye 800CW Streptavidin, anti-Cdc48 serum, or anti-Flag IgG.

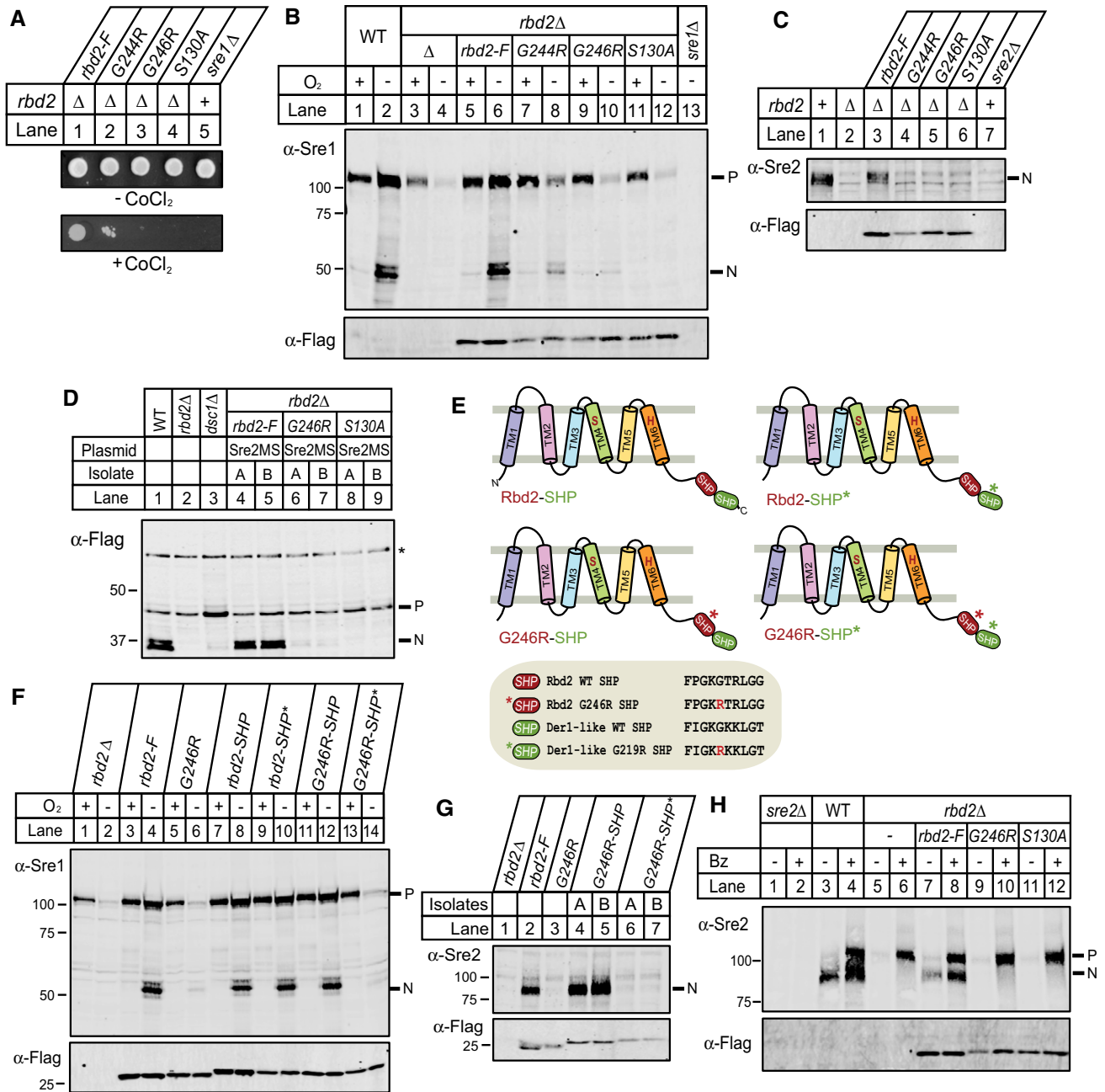
We reasoned that if the Rbd2 SHP box functions to recruit Cdc48, we might restore SREBP cleavage by appending a heterologous SHP box to the Rbd2-*G246R* mutant. Fusion of a SHP box from the fission yeast Der1-like protein (SPBC365.08c) to the C-terminus of Rbd2-*G246R* restored its ability to produce Sre1N to wild-type levels (Fig 8E, *G246R*-SHP; Fig 8F, compare lanes 5–6 to 11–12). In contrast, fusion of a mutant Der1-like SHP box at the equivalent position (*G246R*-SHP\*) failed to rescue Sre1 cleavage (Fig 8F, lanes 13–14). Parallel results were obtained in Sre2 cleavage assays (Fig 8G). Furthermore, fusion of the non-homologous Dsc5 UBX domain that recruits Cdc48 to the Dsc E3 ligase partially rescued Sre1 cleavage (Fig EV3) (Stewart *et al.*, 2012). Thus, SREBP cleavage requires Cdc48 recruitment to Rbd2.

*rbd2*-*G246R* cells failed to generate a detectable SREBP cleavage product despite the presence of an intact catalytic dyad (Fig 8B and D). The inability to detect a cleavage product could be due to a failure to cleave SREBP, or alternatively, a cleaved product may be degraded when Rbd2 fails to recruit Cdc48. To discriminate between these two possibilities, we treated cells with a proteasome inhibitor

and examined Sre2 cleavage. As expected, proteasome inhibition resulted in the accumulation of uncleaved Sre2 precursor in both *rbd2*Δ and *rbd2*-*S130A* cells (Fig 8H, lanes 5–6 and 11–12). *rbd2*-*G246R* cells also accumulated Sre2 precursor, but not a cleaved product, indicating that Cdc48 binding to Rbd2 is required for SREBP cleavage (Fig 8H, lanes 9–10). Thus, the function of Cdc48 is distinct from that shown for p97 with RHBDL4 where p97 acts after cleavage (Fleig *et al.*, 2012). Taken together, these results demonstrate a new function for Cdc48 in mediating Rbd2-dependent cleavage of yeast SREBPs.

#### Cdc48 acts as a substrate adaptor for Rbd2

Rhomboid proteases exhibit weak binding affinity for substrates (Dickey *et al.*, 2013). Given that Cdc48 cofactors bind ubiquitin (Stolz *et al.*, 2011), we hypothesized that Cdc48 plays a role in SREBP recognition by recruiting SREBP substrate to Rbd2 through cofactor binding. If this was true, we reasoned that increasing the concentration of Cdc48-binding defective Rbd2 enzyme would



**Figure 8. SREBP activation requires Cdc48 binding to Rbd2.**

A Yeast strains (200 cells) containing chromosomal Flag-tagged *rbd2*, *rbd2-G244R*, *rbd2-G246R* or *rbd2-S130A* in *rbd2Δ* background, and *sre1Δ* yeast were grown on rich medium in the absence or presence of cobalt chloride (CoCl<sub>2</sub>).

B Western blot of phosphatase-treated, whole-cell lysates from strains in (A) grown for 3 h in the presence or absence of oxygen was probed with anti-Sre1 or anti-Flag IgG. P and N denote Sre1 precursor and cleaved nuclear forms, respectively.

C Sre2 cleavage was assayed by Western blot using anti-Sre2 serum or anti-Flag IgG of phosphatase-treated, whole-cell lysates from the indicated strains. N denotes Sre2 cleaved nuclear form.

D Whole-cell lysates from indicated yeast strains carrying a plasmid expressing 3xFlag-Sre2MS were analyzed by Western blot with anti-Flag antibody. Two independent isolates, A and B, are shown for each strain. Asterisk denotes non-specific band.

E Diagram of four Rbd2 chimeras fused to the SHP box from *Schizosaccharomyces pombe* Der1-like protein (SPBC365.08c). The SHP boxes of Rbd2 and Der1-like protein are shaded red and green, respectively. Asterisks denote point mutation in the SHP box.

F Yeast strains containing Flag-tagged *rbd2-SHP*, *rbd2-SHP\**, *rbd2-G246R-SHP*, and *rbd2-G246R-SHP\** were generated in *rbd2Δ* background by chromosomal integration of the *rbd2* fusion proteins depicted in (E). Indicated yeast strains were then assayed for Sre1 cleavage as in (B).

G Indicated strains were assayed for Sre2 cleavage as in (C).

H Wild type, *sre2Δ*, or strains containing chromosomal Flag-tagged *rbd2*, *rbd2-G246R*, or *rbd2-S130A* in *rbd2Δ* background (lane 7–12) were treated with bortezomib (Bz) for 2 h, and phosphatase-treated, whole-cell lysates were analyzed by Western blot with anti-Sre2 serum or anti-Flag IgG.

bypass the requirement for Cdc48. To test our hypothesis, we performed Sre1 cleavage analysis in yeast cells expressing *rbd2-Flag-APEX2*, *rbd2-G246R-Flag-APEX2*, or *rbd2-S130A-Flag-APEX2* under control of the constitutive *adh1<sup>+</sup>* promoter in *rbd2Δ* cells. Indeed, overexpression of *rbd2-G246R* fully restored Sre1 cleavage, while overexpression of catalytically dead *rbd2-S130A* did not (Fig 9A). In addition, Sre1 precursor levels increased with time in *rbd2-G246R* overexpressing cells, indicating that cleaved Sre1N was transcriptionally active and promoted *sre1* transcription through a well-characterized positive feedback loop (Todd *et al*, 2006). Consistent with the Western blot results, *rbd2-G246R* overexpression rescued growth of *rbd2Δ* cells on CoCl<sub>2</sub> while *rbd2-S130A* did not (Fig 9B). Compared to *rbd2-Flag* expressed from its endogenous promoter, we observed ~14-fold higher expression of *rbd2-Flag-APEX2*, *rbd2-G246R-Flag-APEX2*, and *rbd2-S130A-Flag-APEX2* using the *adh1<sup>+</sup>* promoter (Fig 9C).

Lastly using the proximity biotinylation technique, we tested whether Rbd2 and SREBP interact using catalytically dead Rbd2 to capture substrate-enzyme binding. Proximity biotinylation experiments using endogenous expression of Rbd2-APEX2 failed to detect an interaction with the known binding partner Cdc48, so we overexpressed Rbd2 fusion proteins using the *adh1<sup>+</sup>* promoter. Rbd2-S130A-APEX2 labeled both Cdc48 and GFP-Sre2MS (Fig 9D, lane 5). Sre2 labeling by Rbd2-S130A-APEX2 was specific inasmuch as we observed 10-fold lower labeling of the Golgi mannosyltransferase Anp1 (Fig 9D, lane 4). Consistent with the ability of overexpressed Rbd2-G246R to support SREBP cleavage, we detected equal labeling of GFP-Sre2MS in cells overexpressing Rbd2-S130A/G246R-APEX2 (Fig 9D, lanes 5–6). As expected, Rbd2-S130A/G246R-APEX2 showed reduced Cdc48 labeling (Fig 9D, lane 6). Collectively, these results demonstrate that increasing the concentration of Rbd2 enzyme bypasses the requirement of Cdc48 binding for Sre1 cleavage, defining a role for Cdc48 in substrate recruitment.

## Discussion

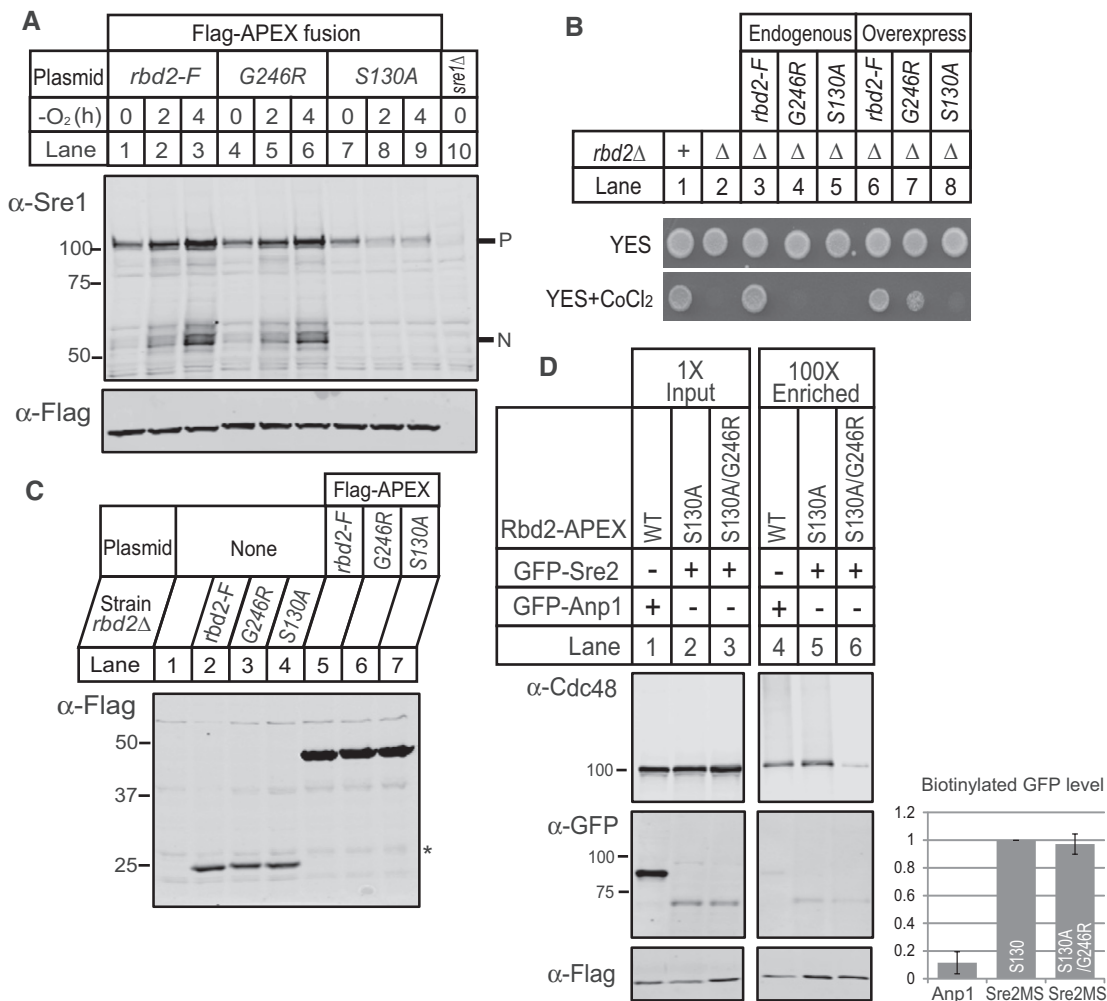
Rbd2 is the first SREBP protease identified in fission yeast (Kim *et al*, 2015). Using genetic interaction mapping, we independently identified Rbd2 as a Golgi-localized rhomboid protease required for proteolytic activation of yeast SREBPs. Deleting *rbd2* or mutating its catalytic residues prevented cleavage of both Sre1 and Sre2 (Figs 2 and 3). Our mammalian cell-based, rhomboid cleavage assays using heterologous substrates showed that Rbd2 is a *bona fide* rhomboid protease with a conserved, Ser-His catalytic dyad (Fig 4). Notably, Rbd2 showed novel substrate specificity: Rbd2 cleaved neither the canonical *D. melanogaster* substrate Spitz nor an alternative *P. falciparum* substrate EBA175, but cleaved *P. stuartii* TatA, the most efficient rhomboid substrate (Strisovsky *et al*, 2009; Moin & Urban, 2012). To date, characterized rhomboid proteases cleave substrates in the membrane after residues with small side chains (Strisovsky *et al*, 2009; Urban, 2010). Although Rbd2 cleavage of TatA required substrate helix-destabilizing residues, Rbd2 preferentially cleaved TatA substrates after large hydrophobic side chains (Leu and Phe) (Fig 4F), revealing a unique property not observed for any other rhomboid protease.

Mammalian SREBPs are activated by regulated intramembrane proteolysis (Brown *et al*, 2000). The serine protease S1P cleaves

SREBP in the luminal loop to produce a type-II (N<sub>in</sub>-C<sub>out</sub>), single transmembrane intermediate that is subsequently cleaved in the membrane by the zinc metalloprotease S2P. Rhomboid proteases preferentially cleave transmembrane substrates with a type-I (N<sub>out</sub>-C<sub>in</sub>) orientation (Urban & Freeman, 2003; Ha, 2009). Despite the sequence requirements observed for Rbd2 in TatA cleavage, mutation of any single large hydrophobic residue in the N-terminal end of TM2 did not block Sre2MS cleavage (Fig 5B). However, the inability to map the Sre2MS cleavage site(s) prevented us from determining whether the mutants support cleavage by the shifting cleavage site to other large, hydrophobic residues. Such cleavage site shifts in substrate mutants are common (Strisovsky *et al*, 2009; Moin & Urban, 2012). Our previous mutagenesis studies showed that Sre2MS cleavage required multiple residues in its type-I oriented TM2, while we found no essential residues in the type-II oriented TM1 (Cheung & Espenshade, 2013). Here, we assigned functions for these residues to either Dsc E3 ligase activity or Rbd2 cleavage. Lysine 743 in the luminal loop was the only residue required for Rbd2-mediated Sre2MS cleavage, while others were involved in Dsc E3 ligase function (Fig 5 and Appendix Figs S1 and S2). Precisely how this luminal lysine residue in Sre2 promotes cleavage remains to be determined.

Should Rbd2 cleave SREBP between the transmembrane segments, it would functionally replace S1P by initiating SREBP activation and yield a type-II intermediate requiring additional processing. Importantly, the active Sre1N and Sre2N terminate at a cytosolic position, ~10 amino acids prior to TM1 (Stewart *et al*, 2011; Cheung & Espenshade, 2013). These observations suggest that SREBP activation requires a second protease. We previously proposed that SREBP activation required the proteasome (Stewart *et al*, 2011), but results presented in this study suggest otherwise. In the earlier study, we used a temperature-sensitive mutation in *mts3* to impair 19S proteasome function, and we observed decreased levels of Sre1N under low oxygen (Stewart *et al*, 2011). However, in that experiment, *mts3-1* cells required a 3-h preincubation at non-permissive temperature, and Sre1 precursor levels also decreased during the course of this experiment, suggesting that proteasome inactivation had indirect effects on SREBP cleavage. In the current study, we observed that cleavage of Sre2 was not inhibited in *mts3-1* cells grown for three hours at the non-permissive temperature (Fig 3A). Furthermore, acute proteasome inhibition with bortezomib did not block Sre2 processing (Fig 3B). Collectively, these data suggest that an undiscovered protease, rather than the proteasome, functions in SREBP proteolytic processing downstream of Rbd2, and that fission yeast may employ a new variation of RIP to activate SREBPs.

Parallel studies in *A. fumigatus* recently identified an Rbd2 homolog, RbdA/B, as required for SREBP activation, demonstrating that this function is conserved in ascomycetous fungi (Dhingra *et al*, 2016; Vaknin *et al*, 2016). In addition, studies in *Aspergillus nidulans* found that SREBP activation required both the Dsc E3 ligase and the signal peptide peptidase SppA, an aspartyl protease (Bat-Ochir *et al*, 2016). Taken together, these recent *Aspergillus* studies support a model for SREBP cleavage in which Rbd2 initiates activation followed by SppA cleavage. However, unlike *A. nidulans* SREBP cleavage, *S. pombe* Sre1 cleavage did not require Ypf1, the *S. pombe* signal peptide peptidase homolog, or two other aspartyl proteases Yps1 or Sxa1 (Fig EV4), leaving unknown the identity of the second SREBP protease in fission yeast.



Prior to these studies, it was known that SREBP cleavage required *cdc48*, but Cdc48's role was unclear because its binding to the Dsc E3 ligase was not required for SREBP cleavage (Stewart et al, 2012). Here, we demonstrate that SREBP cleavage requires Cdc48 binding to the C-terminal SHP box in Rbd2 (Fig 8). Cells expressing an Rbd2 mutant defective for Cdc48 binding accumulated Sre2 precursor with proteasome inhibition rather than cleaved

products (Fig 8H). Thus, Cdc48 functions in Rbd2-dependent cleavage. The human rhomboid protease RHBDL4 binds ubiquitinated substrates through a ubiquitin-interacting motif and cleaves them in the ER to promote ERAD (Fleig et al, 2012). Interestingly, RHBDL4 also binds to p97/Cdc48 but through a different motif, called VBM. p97/Cdc48 binding to RHBDL4 facilitates extraction of cleaved products for proteasomal degradation. Cdc48 and its cofactors bind to

ubiquitin (Stolz et al, 2011). In a function similar to that for the RHBDL4 ubiquitin-interacting motif, we propose that Cdc48 with its cofactors is a receptor for ubiquitin, thereby recruiting ubiquitylated SREBPs to Rbd2. While p97 functions after RHBDL4 cleavage, here Cdc48 functions in recognizing substrate prior to the cleavage event. Supporting this model, overexpression of Rbd2 that cannot bind Cdc48 rescued Sre1 cleavage, bypassing the requirement for Cdc48 (Fig 9).

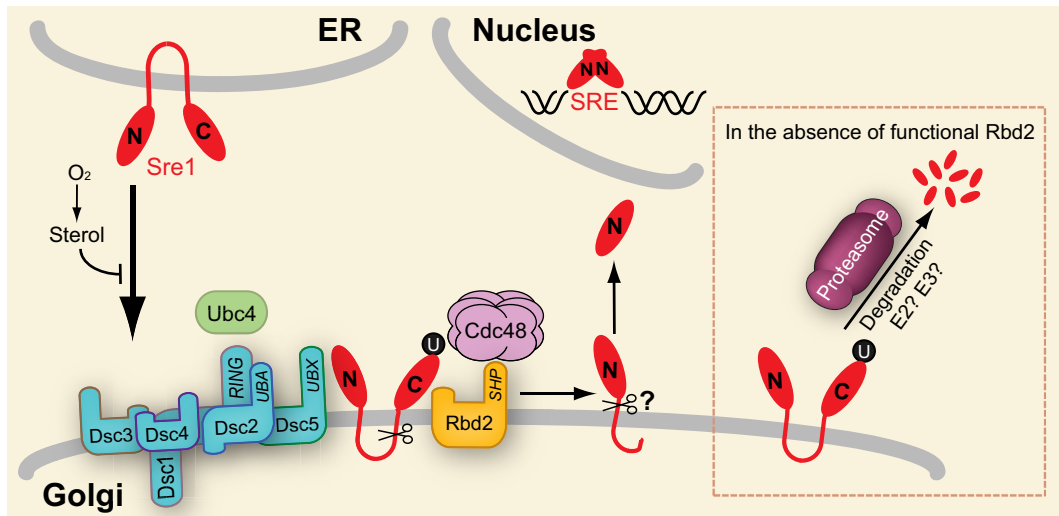
As noted, activation of *A. fumigatus* SREBP requires a rhomboid protease (Dhingra et al, 2016; Vaknin et al, 2016). Our bioinformatic analysis identified a conserved SHP box in the cytosolic C-terminus of *A. fumigatus* RbdB (Fig EV5A) (Sato & Hampton, 2006). GST-RbdB (aa 207–272) bound to Cdc48 *in vitro*, but the binding was ~7-fold lower than GST-Rbd2 (Fig EV5B). Reduced binding to RbdB may be due to an asparagine substitution at the G246 position in the Rbd2 SHP box (Fig EV5A). *S. cerevisiae* Rbd2 which does not contain a conserved SHP box was used as a negative control. Testing whether Cdc48 binding to RbdB is also required for *A. fumigatus* SREBP, cleavage activation is an important next step to further understand how Cdc48 functions with rhomboid proteases.

The Dsc1 subunit of the Dsc E3 ligase contains a RING domain and functions as an ubiquitin ligase with the E2 ubiquitin-conjugating enzyme Ubc4 (Raychaudhuri & Espenshade, 2015). Mutation of the Dsc1 RING domain, deletion of *dsc1-dsc5*, or mutation of *ubc4* blocks SREBP cleavage (Stewart et al, 2011, 2012). In addition, Sre2 binds to the Dsc E3 ligase (Stewart et al, 2011; Lloyd et al, 2013). Together, these data suggest that SREBPs are substrates for the Dsc E3 ubiquitin ligase. However, recently we found that inhibition of Dsc E3 ligase activity prevents ER-to-Golgi transport of the Dsc complex, confounding the interpretation of these *dsc* mutant phenotypes (Raychaudhuri & Espenshade, 2015). Here, we observed that in *rbd2Δ* cells, SREBPs

are degraded by the proteasome in a Dsc-dependent manner (Figs 3 and EV1). Although SREBP ubiquitylation remains to be shown, these data provide additional support for SREBPs as Dsc E3 ligase substrates. Interestingly, preliminary studies indicate that cytosolic lysines in Sre2MS are not required for cleavage, suggesting that SREBP ubiquitylation may involve non-classical sites. Ubc4 conjugates mono-ubiquitin to proteins, while proteasomal degradation typically requires poly-ubiquitin chains (Seino et al, 2003; Rodrigo-Brenni & Morgan, 2007). Whether other ubiquitylation enzymes are required for SREBP degradation in *rbd2Δ* cells and whether these enzymes are also required for SREBP cleavage remains to be tested. However, our studies using *hrd1Δ* and *doa10Δ* cells indicate that these two ERAD E3 ligases are not required for cleavage or degradation (Fig EV1A).

Combined with our past work, these new findings lead us to speculate on a new model for the activation of Sre1 (Fig 10). Following oxygen-regulated transport by the SCAP homolog Scp1 to the Golgi, Sre1 is ubiquitylated by the Dsc E3 ligase complex. Rbd2 cleaves ubiquitylated Sre1 in the second transmembrane segment or luminal juxtamembrane region in a reaction that requires substrate recruitment by Cdc48. This Sre1 intermediate is then cleaved by a second, unidentified protease to produce the soluble, active transcription factor. Sre2 activation follows a similar pathway, but ER-to-Golgi transport is constitutive. In the absence of Rbd2 cleavage (Fig 10 dashed box), ubiquitylated SREBPs are degraded by the proteasome. Thus, changes in Rbd2 activity will control the fate of SREBP substrates between protein processing and degradation. This mechanism provides a new control point for the regulation of SREBP activity in fungi. In mammals, a related regulatory mechanism exists: when S1P is inhibited, SCAP is targeted to the lysosome for degradation, leading to reduced pathway activity (Shao & Espenshade, 2014).

In summary, convergent evidence from studies in *S. pombe* and *A. fumigatus* demonstrates that SREBP activation requires a



**Figure 10. Model for yeast SREBP cleavage.**

Under low sterol and oxygen conditions, Sre1 exits the ER and travels to the Golgi. Golgi-localized Sre1 is ubiquitylated by E2 Ubc4 and Dsc E3 ligase. Ubiquitylated Sre1 is recruited by Cdc48 and cleaved by Rbd2. The intermediate cleaved product is further processed by a second, unknown protease to generate Sre1N that traffics to the nucleus to activate transcription. When Rbd2 is inactive (dashed box), ubiquitylated Sre1 is subjected to proteasomal degradation. SRE, sterol regulatory element; U, either mono-ubiquitin or poly-ubiquitin.

rhomboid protease, Rbd2 and RbdA/B, respectively (Bat-Ochir *et al*, 2016; Dhingra *et al*, 2016; Vaknin *et al*, 2016). Fungal SREBPs are required for growth under hypoxic conditions and for virulence of human opportunistic pathogens (Bien & Espenshade, 2010). In addition, SREBP is required for virulence of *Penicillium digitatum*, a major citrus fruit fungal pathogen (Liu *et al*, 2015) and *P. digitatum* contains an *rbd2* homolog (ORF name, PDIP\_45580). Thus, identification of a fungal SREBP protease opens the door to development of rhomboid inhibitors that could act as antifungals to improve both clinical and agricultural outcomes.

## Materials and Methods

Materials and standard methods (cell culture, plasmid construction, linkage analysis, Western blotting, immunoprecipitation, *in vitro* binding assay, microscopy, and mass spectrometry) are described in the Appendix Supplementary Methods. Yeast strains and oligonucleotide primers used in this study may be found in Appendix Tables S1 and S2, respectively.

### Yeast strain construction

The *rbd2*-integrating plasmids contained *rbd2*<sup>+</sup>, *rbd2-S130A-3xFlag*, *rbd2-H182A-3xFlag*, *rbd2-G244R-3xFlag*, *rbd2-G246R-3xFlag*, *rbd2-SHP-3xFlag*, or *rbd2-SHP\*-3xFlag* coding sequence flanked by 600 bp of upstream genomic sequence and 600 bp of downstream genomic sequence with a *his3*<sup>+</sup> marker for chromosomal reconstitution of histidine auxotrophy. Plasmids containing *rbd2* wild type and mutants were linearized with *AscI* and transformed into *rbd2Δ* strain (PEY1681) and selected for minimal medium lacking histidine to generate strains carrying integrated wild-type and mutant alleles of *rbd2* at the *his3*<sup>+</sup> locus. Isolates expressing Rbd2-3xFlag were confirmed by Western blotting. The *rbd2-Flag-APEX2* plasmid and its derivatives were transformed into *rbd2Δ* strain and selected for minimal medium lacking leucine to generate strains carrying *rbd2-Flag-APEX2*, *rbd2-S130A-Flag-APEX2*, *rbd2-G246R-Flag-APEX2*, or *rbd2-G246R/S130A-Flag-APEX2*.

### Genetic interaction study

The genetic interaction study was performed as described (Frost *et al*, 2012). Included in the query set for the study were the *dsc* genes and multiple alleles of *cdc48*. Hierarchical clustering profiles were analyzed for genetic interaction profiles similar to the *dsc* genes.

### Biotin-phenol labeling

The original biotin-phenol labeling protocol for mammalian cells was modified for yeast (Hung *et al*, 2016; Hwang & Espenshade, 2016). Exponentially growing cells ( $5 \times 10^8$ ) were harvested and washed with water. Cell pellets were resuspended in 1 ml of 1.2 M sorbitol dissolved in H<sub>2</sub>O. Biotin-phenol (2.5 mM) was added, and the cells were incubated for another 1 h at room temperature. H<sub>2</sub>O<sub>2</sub> (1 mM) was added for 1 min to initiate biotin-phenol labeling, and cells were quickly spun down. To stop the reaction, the solution was aspirated off and cell pellets were then washed four times with a

quenching solution consisting of 5 mM Trolox, 10 mM sodium azide, and 10 mM sodium ascorbate in 1.2 M sorbitol dissolved in H<sub>2</sub>O. Cell pellets were washed with 1.2 M sorbitol once more and lysed by addition of 27 mM NaOH, 1% (v/v) 2-mercaptoethanol for 10 min on ice. Total protein was precipitated with trichloroacetic acid (20% w/v) followed by acetone wash. Proteins were solubilized in 200 μl lysis buffer (25 mM Tris pH 7.4, 150 mM NaCl, 1% SDS) containing Complete Protease Inhibitor EDTA free by sonication. After denaturation by heating at 75°C for 15 min, protein samples were either directly analyzed by Western blotting or subjected to enrichment of biotinylated proteins. For enrichment, proteins (1–2 mg) were first dialyzed in dialysis buffer (25 mM Tris pH 7.4, 150 mM NaCl, 0.2% SDS) using 3.5K MWCO dialysis tubing (Thermoscientific) for 2–3 h at room temperature, and then incubated with 50–100 μl streptavidin-coated magnetic beads for 2–3 h at room temperature. Streptavidin beads were then washed with 0.3% SDS-containing wash buffer (25 mM Tris pH 7.4, 150 mM NaCl, 0.3% SDS) six times followed by 2 × wash with 2 M urea/50 mM Tris pH 7.4, 1 × wash with 1 M KCl, and 1 × with wash buffer. Biotinylated proteins were eluted by incubating the beads with 30 μl 2 × SDS sample loading buffer and heating to 75°C for 15 min.

**Expanded View** for this article is available online.

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### Author contributions

JH, DR, and PJE designed and performed experiments except as noted. SR and HG contributed to Figs 3 and 6, respectively. SU and LC performed experiments in Fig 4. AF performed genetic interaction study. JH, DR, SU, and PJE wrote the manuscript with contributions from the other authors.

### Conflict of interest

The authors declare that they have no conflict of interest.

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