DOA1/UFD3 Plays a Role in Sorting Ubiquitinated Membrane Proteins into Multivesicular Bodies*

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Ubiquitin (Ub) is a sorting signal that targets integral membrane proteins to the interior of the vacuole/lysosome by directing them into lumenal vesicles of multivesicular bodies (MVBs). The Vps27-Hse1 complex, which is homologous to the Hrs-STAM complex in mammalian cells, serves as a Ub-sorting receptor at the surface of early endosomes. We have found that Hse1 interacts with Doa1/Ufd3. Doa1 is known to interact with Cdc48/p97 and Ub and is required for maintaining Ub levels. We find that the Hse1 Src homology 3 domain binds directly to the central PFU domain of Doa1. Mutations in Doa1 that block Hse1 binding but not Ub binding do not alter Ub levels but do result in the missorting of the MVB cargo GFP-Gps1. Loss of Doa1 also causes a synthetic growth defect when combined with loss of Vps27. Unlike the loss of Doa1 alone, the doa1Δ vps27Δ double mutant phenotype is not suppressed by Ub overexpression, demonstrating that the effect is not due to indirect consequence of lowered Ub levels. Loss of Doa1 results in a defect in the accumulation of GFP-Ub within yeast vacuoles, implying that there is a reduction in the flux of ubiquitinated membrane proteins through the MVB pathway. This defect was also reflected by an inability to properly sort Vph1-GFP-Ub, a modified subunit of the multiprotein vacuolar ATPase complex, which carries an in-frame fusion of Ub as an MVB sorting signal. These results reveal novel roles for Doa1 in helping to process ubiquitinated membrane proteins for sorting into MVBs.

One of the key sorting steps in sending integral membrane proteins to the lysosome for degradation is their incorporation into vesicles that form from the limiting membrane of early endosomes and bud into the lumen. This budding process results in the formation of a multivesicular body or multivesicular endosome that fuses to lysosomes to deliver its intralumenal vesicles and membrane cargo proteins to the hydrolytic lysosomal lumen (1). Integral membrane proteins gain access into the MVB sorting pathway by becoming attached to ubiquitin (Ub). Ub attachment serves as a sorting signal at the plasma membrane for internalization, at the endosome for incorporation into intralumenal vesicles that comprise MVBs, and at the trans-Golgi network, from where proteins are sorted into vesicles that are delivered to the forming MVB (1, 2). The processing and sorting of Ub cargo at the endosome is in part fulfilled by a set of proteins originally identified in yeast as class E Vps (vacuolar protein sorting) proteins. This set includes 18 proteins whose loss impairs lumenal vesicle formation and protein sorting and results in the accumulation of aberrantly large late endosomal (class E compartment) structures adjacent to the yeast vacuole (lysosome). The majority of the yeast class E proteins have clear functional homologs in mammalian cells and associate in distinct complexes, including ESCRT I, II, III, a Vps4 AAA ATPase complex, and the Vps27-Hse1 (has symptoms of class E) complex, whose mammalian ortholog is the Hrs-STAM complex (1, 2).

Several class E Vps protein components, including Vps27, Vps23 (ESCRT-I), and Vps36 (ESCRT-II), can noncovalently bind Ub via discrete functional domains and are thus believed to have a role in the recognition of Ub cargo. The Vps27-Hse1 complex largely localizes to endosomes via its association with phosphatidylinositol 3-phosphate, clathrin, and ESCRT-I (3–6). Current models suggest that after Ub cargo is captured by the Vps27-Hse1 complex, it passes through a series of steps that concentrate cargo within areas that will undergo vesicle formation (1, 2). At a late point in the sorting process, Ub is removed from cargo by the deubiquitinating enzyme Ubp4 (ubiquitin peptidase 4)/Doa4 (degradation of MATα2) before incorporation into intralumenal vesicles of the MVB (7). Loss of DOA4 depletes cellular Ub levels, which causes a number of cellular defects, including loss of efficient MVB sorting of a number of integral membrane proteins (8–11). However, despite the wealth of structural data that describe the architecture of these protein complexes and how they interact, little is known about the exact sorting steps they perform or how they are regulated.

The Vps27-Hse1/Hrs-STAM complex has emerged as at least one important focal point for regulation. In mammalian cells, the Hrs-STAM complex interacts with numerous effectors of protein trafficking and is the target of receptor-tyrosine

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3 The abbreviations used are: Ub, ubiquitin; MVB, multivesicular body; GFP, green fluorescent protein; DIC, differential interference contrast; SH3, Src homology 3; SD, synthetic dextrose; AAA, ATPases associated with various cellular activities; PIPES, 1,4-piperazinediethanesulfonic acid; GST, glutathione S-transferase.
kinases, which can modulate their own down-regulation by tyrosine-phosphorylating Hrs (12–18). This complex also associates with deubiquitinating enzymes and Ub-ligases, and recent studies in yeast suggest that this association serves to regulate the ubiquitination status of membrane protein cargo to either reverse or reinforce cargo trafficking along the MVB pathway (19, 20).

To understand more about how MVB sorting is controlled, we searched for protein machinery that interacts with the Vps27-Hse1 complex and found Ufd3 (ubiquitin fusion degradation 3)/Doa1. Doa1 was first found in genetic screens for mutants defective in degrading model proteasome substrates (21–23). Loss of Doa1 causes many phenotypes in yeast, including sensitivity to DNA-damaging agents, cycloheximide, caffeine, cadmium, canavanine, growth at high temperature, and volatile anesthetics (24–31). Other analysis has shown that loss of Doa1 dramatically decreases Ub levels (21). Many of the phenotypes of doa1Δ mutants can be suppressed by overexpressing Uf3, suggesting that many of the noted doa1Δ defects are solely an indirect consequence of lowered Ub levels (21, 26, 28, 31). It is not presently clear how Ub is depleted in the absence of Doa1; however, inhibiting delivery and degradation of ubiquitinated proteins to the proteasome suppresses doa1Δ growth defects, suggesting that more Ub may be degraded by the proteasome in the absence of Doa1 (24, 28, 31).

The N-terminal region of Doa1 contains a seven-bladed WD40 repeat β-propeller, which in general is thought to mediate protein-protein interactions (32). Doa1 also has two other regions dubbed the PFU and PUL domains, based on their shared sequence homology with Doa1 orthologs throughout eukaryota (27, 33). The central PFU domain of Doa1 mediates interaction with Ub, whereas the C-terminal PUL domain mediates interaction with Cdc48 (cell division cycle 48), an AAA ATPase belonging to a family of ATPases associated with various cellular activities (27, 28). Cdc48 (and its mammalian ortholog p97) forms a hexameric ring that acts as a molecular chaperone for a variety of ubiquitinated proteins (34–36). Cdc48 is organized into an N-terminal portion, followed by two AAA ATPase domains. Recent biochemical studies confirm that Cdc48 probably works as a “segregate” that can dissociate aggregates of ubiquitinated proteins, which would help convey them to the proteasome for degradation (37). Cdc48/p97 associates with a variety of “adaptor” proteins that help program it for various functions. Adaptor proteins, such as the UBX family of proteins, bind to the N-terminal portion of Cdc48 (36), whereas Doa1, the Ub E3/E4 ligase Ufd2, and peptide:N-glycanase associate with the C terminus (28, 35, 38). Although the role for some of these Cdc48/adaptor complexes is known, little is known about the specific functions executed by a Doa1-Cdc48 complex.

Our analysis indicates that Doa1 plays a role in sorting Ub cargo into MVBs. Part of this activity is directed specifically by association with the Vps27-Hse1 complex, and this association is required for the efficient sorting of particular MVB substrates. However, Doa1 also appears to serve a more general role to help capture and process Ub cargo for concentration at endosomal subdomains and eventual incorporation into lumenal vesicles.

**EXPERIMENTAL PROCEDURES**

**Materials, Yeast Strains, and Plasmids—**Synthetic dextrose (SD) medium was made using yeast nitrogen base containing ammonia and 2% glucose. Yeast nitrogen base was purchased from RPI Research Products International Corp. Amino acid supplements were purchased from Bio 101, Inc. (La Jolla, CA). Glutathione-agarose beads were purchased from GE Healthcare. Zymolyase 100T was purchased from Seikagaku Corp. (East Falmouth, MA). Protease inhibitor mixture (Complete™) was purchased from Roche Applied Science. Endocytic tracer dye FM4-64 was purchased from Molecular Probes, Inc. (Eugene, OR). The pCR2.1, pYES2.1, and pET151 TOPO cloning kits were purchased from Invitrogen. Anti-Ub monoclonal antibody P4D1 was purchased from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). Monoclonal anti-CYP 10A5-B5 was a kind gift from Tom Stevens (University of Oregon). Anti-HA antibody was purchased from Covance Research Products, Inc. (Berkeley, CA); anti-V5 was from Amersham Biosciences.

Saccharomyces cerevisiae strains used in this study are listed in Table 1. The parental strain was BY4742 (MATa his3Δ1

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**TABLE 1** Yeast strains used in this study

| Strain     | Genotype            | Reference/Source |
|------------|---------------------|------------------|
| SF389-9D   | MATa leu2-3,112 ura3-52 his4-519 ade6 pep4-3 | Ref. 69          |
| BY4742     | MATa his3 leu2 lys2 | Ref. 70          |
| PLY2498    | MATa hse1Δ::Kan' his3, leu2, lys2, ura3 | Yeast Gene Deletion Project |
| PLY3705    | MATa otu1Δ::Kan' his3, leu2, lys2, ura3 | Yeast Gene Deletion Project |
| PLY3706    | MATa ufd2Δ::Kan' his3, leu2, lys2, ura3 | Yeast Gene Deletion Project |
| PLY3704    | MATa vps23Δ::Kan' his3, leu2, lys2, ura3 | Yeast Gene Deletion Project |
| PLY3709    | MATa doa1Δ::HIS3 his3, leu2, lys2, ura3 | This study       |
| PLY3462    | MATa doa1Δ::Kan' his3, leu2, lys2, ura3 | This study       |
| PLY3694    | MATa doa1Δ::Kan' vps27Δ::LEU2 leu2 lys2, ura3 | This study       |
| PLY3249    | MATa hse1Δ::URA3 his3, leu2, lys2, ura3 | Ref. 20          |
| PLY3175    | MATa vps27Δ::Kan' leu2 lys2, ura3 | Ref. 20          |
| PLY3556    | MATa doa1Δ::HIS3 vps27Δ::URA3 leu2 lys2, ura3 | This study       |
| PLY3700    | MATa vps27Δ::LEU2 ufd6Δ::Kan' his3, leu2, lys2, ura3 | This study       |
| PLY3699    | MATa vps27Δ::LEU2 otu1Δ::Kan' his3, leu2, lys2, ura3 | This study       |
| PLY3702    | MATa vps27Δ::LEU2 ufd2Δ::Kan' his3, leu2, lys2, ura3 | This study       |
| PLY3707    | MATa doa1Δ::HIS3 ufd2Δ::Kan' his3, leu2, lys2, ura3 | This study       |
| PLY3711    | MATa vps27Δ::LEU2 doa1Δ::HIS3 ufd2Δ::Kan' his3, leu2, lys2, ura3 | This study       |
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Doa1 Binds Hse1 and Is Required for Sorting MVB Substrates—Previous large scale two-hybrid screens identified Doa1 as potentially interacting with the SH3 domain of Hse1 (46). To confirm this interaction, we produced a C-terminally V5 epitope-tagged Doa1 in yeast and subjected the corresponding lysates to GST pull-down experiments using the SH3 domain of Hse1. Fig. 1A shows that the Doa1 specifically bound the Hse1 SH3 domain. However, Doa1 did not bind the SH3 domain of Pex13. Furthermore, binding of Doa1 was dramatically reduced when a critical tryptophan residue within the predicted hydrophobic binding surface of the Hse1 SH3 domain was altered to alanine.

We next asked whether Doa1 could localize to endosomes. To image Doa1, we fused GFP to its C terminus. The strategy was used previously in a genome-wide effort to C-terminally GFP-tag all yeast open reading frames systematically (47). We first verified that the DOA1-GFP-tagged strain from this collection, in which the only copy of Doa1 is fused to GFP, was able to grow like wild type cells at 37 °C, indicating that the Doa1-GFP was functional (data not shown). We then drove expression of Doa1-GFP from the moderately strong PRC1 promoter, which normally drives the expression of vacuolar carboxypeptidase Y. In wild type cells, Doa1-GFP was localized diffusely to the cytosol, was excluded from the vacuoles, and was slightly concentrated in the nucleus, consistent with previous observations (47). We then localized Doa1-GFP in mutants lacking Vps4, a class E Vps protein whose loss results in the accumulation of...

RESULTS

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A434AAKAAA to make doa1ΔHse1. PFU domain from pPL3371 (deletion of residues 433–445) tagged C-terminally with V5 and His6 tag in 2Hse1. Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. This study

pPL3601 GALL driving DOA1 tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1 Contains mutations P99PLPK → A99 ALKA. This study

pPL3637 GALL driving DOA1 tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1 Contains mutations F434ILKNTN → A434 AAKAAA to make doa1ΔHse1. Fragment derived from pPL367 (F434ILKNTN → A434 AAKAAA) residues 301–466 with N-terminal V5 epitope tag in pET151. This study

pPL3638 Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. This study

pPL3639 Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. This study

pPL3498 DOA1-V5 expressed from DOA1 promoter in pRS316. This study

pPL3499 Doa1ΔHse1 (F434ILKNTN → A434 AAKAAA) expressed from DOA1 promoter in pRS316. This study

pPL3501 Doa1ΔHse1 (deletion of residues 433–445) expressed from DOA1 promoter in pRS316. This study

pUG27 Plasmid containing gene disruption cassettes with his5 (S. pombe) as selection marker. Ref. 40

pPL2161 HA-tagged HSE1 in pRS315 Ref. 3

pPL1124 VPS9 dominant negative in pRS426 Ref. 20

gEX6-1 GST expression vector

pET151 T7 promoter-based bacterial expression vector.

TABLE 2
Plasmids used in this study

| Plasmid | Description | Reference/Source |
|---------|-------------|------------------|
| pGO45   | pRS426 carrying GFP-CPS1 | Ref. 71 |
| pPL1867 | LEU2 conversion of pGO45 | This study |
| pPL3267 | HA-Ub-GFP-CPS1 in pRS316 | Ref. 20 |
| pPL3453 | HA-Ub-GFP-CPS1 in pRS315 | This study |
| pPL1124 | pRS316 containing dominant negative VPS4 | GenBank™ accession number U78872 |
| pGEX6-1 | GST expression vector | This study |
| pPL2710 | GST-Hse1-SH3 | Ref. 20 |
| pPL2831 | GST-Pex13-SH3 | Ref. 20 |
| pPL3164 | GST-Hse1-SH3. GST-Hse1-SH3 plasmid containing in which the codons encoding W254A and W255A mutations. | Ref. 20 |
| pPL1978 | GST-Vps27 C terminus | Ref. 3 |
| pPL833 | JLU34 | This study |
| pGFP-Ub | GFP-UB expressed from the PRCl promoter in pRS426. | D. Katzmann Mayo Clinic (Rochester, MN) |
| pPL3601 | LEU2 marker convert of pGFP-Ub. | This study |
| pPL3155 | YEp351 carrying URA4 | This study |
| pPL3327 | DOA1-GFP downstream of PRC1 promoter and upstream of the PHO8 3′-UT in pRS316. | This study |
| pPL2794 | GALL driving full-length DOA1 tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3270 | GALL driving truncated DOA1I (residues 1–300) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3273 | GALL driving truncated DOA1I (residues 1–470) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3359 | GALL driving truncated DOA1I (residues 1–414) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3363 | GALL driving truncated DOA1I (residues 1–433) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3341 | GALL driving truncated DOA1I (residues 1–445) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3323 | GALL driving a fragment of DOA1I (residues 301–470) tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. | This study |
| pPL3300 | GALL driving DOA1 tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. Contains mutations P99PLPK → A99 ALKA. | This study |
| pPL3367 | GALL driving DOA1I tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. Contains mutations F434ILKNTN → A434 AAKAAA to make doa1ΔHse1. | This study |
| pPL3371 | GALL driving DOA1I tagged C-terminally with V5 and His, tag in 2µ plasmid pYES2.1. Contains deletions of residues 433–445 to make doa1ΔHse1. | This study |
| pPL2967 | Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. | This study |
| pPL3604 | Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. | This study |
| pPL3603 | Expression plasmid for PFU domain from wild type DOA1. Residues 301–466 with N-terminal V5 epitope tag in pET151. | This study |
| pPL3498 | DOA1-V5 expressed from DOA1 promoter in pRS316. | This study |
| pPL3499 | Doa1ΔHse1 (F434ILKNTN → A434 AAKAAA) expressed from DOA1 promoter in pRS316. | This study |
| pPL3501 | Doa1ΔHse1 (deletion of residues 433–445) expressed from DOA1 promoter in pRS316. | This study |
| pUG27 | Plasmid containing gene disruption cassettes with his5 (S. pombe) as selection marker. | Ref. 40 |
| pPL2161 | HA-tagged HSE1 in pRS315 | Ref. 3 |
| pPL1124 | VPS9 dominant negative in pRS426 | Ref. 20 |
| gEX6-1 | GST expression vector | GenBank™ accession number U78872 |
| pPL1556 | VPH1-GFP-UB in pRS316 | Ref. 41 |
| pYES2.1 | High copy (2µ) URA3-containing GALI expression plasmid. | Invitrogen |
| pET151 | T7 promoter-based bacterial expression vector. | Invitrogen |

aberrantly large endosomes (class E compartments), which can trap a variety of proteins that transiently associate with endosomes (48, 49). Here we saw colocalization of Doa1-GFP within membrane patches adjacent to the vacuole that also accumulated the endocytic tracer dye FM4-64 (Fig. 1B). These data indicated that Doa1-GFP was capable of localizing to endosomes.

We also tested whether loss of DOA1 caused defects in MVB sorting. For this, we used two MVB marker proteins. The first was GFP-tagged Cps1, a vacuolar protease that gains access to the vacuolar lumen via the MVB pathway (50). GFP-Cps1 is a Type I membrane protein containing GFP fused to its cytosolic N terminus. Cps1 is ubiquitinated by Rsp5 and Tull and is then sorted by the class E Vps machinery into the vacuolar lumen (51, 52). The second cargo was Ub-GFP-Cps1, in which GFP-Cps1 is translationally fused to Ub (20). This cargo behaves similarly to GFP-Cps1 except that it does not require Rsp5-dependent ubiquitination to undergo MVB sorting, since it contains its Ub sorting motif constitutively (51). Fig. 1C shows that although wild type cells efficiently sorted GFP-Cps1 into the vacuolar lumen, doa1Δ cells did not and accumulated GFP-Cps1 on the limiting membrane of the vacuole. In contrast, Ub-GFP-Cps1, which does not require ubiquitination to undergo Ub-dependent MVB sorting, was sorted correctly to the vacuolar lumen in doa1Δ cells.

These results indicated that Doa1 might indeed provide a function with regard to MVB sorting, presumably by promoting the processing and ubiquitination of cargo proteins like GFP-Cps1. The major caveat to this interpretation is that doa1Δ mutants have lowered levels of Ub, which occurs through an unknown mechanism (21). Lower Ub levels could indirectly affect a number of processes. Therefore, we went on to further characterize the specificity of these effects and examine the functional relevance of the Hse1-Doa1 interaction.

Doa1 Contains a Novel Binding Motif for the Hse1 SH3 Domain—Previous studies indicated that the SH3 domain of Hse1 was the relevant portion of Hse1 that could interact with Doa1 (46). Our previous studies showed that this domain is critical for Hse1 functions but also mediates interactions with other proteins, including UbP7 and the Rsp5-Ubp2-Rup1-
Hua1 complex (20). Thus, phenotypic analysis of Hse1 lacking its SH3 domain would not specifically ablate the Hse1-Doa1 interaction. Therefore, we undertook a series of mapping experiments to find the motif within Doa1 that mediates its interaction with the Hse1 SH3 domain. Fig. 2A shows a set of Doa1 deletion and substitution mutants subjected to GST pull-down experiments using GST alone or GST fused to the SH3 domain of Hse1. As a further control, we also used GST fused to the SH3 domain of Hse1 in which the predicted SH3 ligand was mutated (20, 53). We found that full-length Doa1 and Doa1 lacking its C-terminal 270 residues (Doa1 1–445) also specifically bound to the Hse1 SH3 domain. As a further control, we also used GST fused to the SH3 domain from Pex13 (Pex13 SH3). Also shown are the GST fusion proteins used in this analysis. B, Doa1-GFP (expressed from pPL3327) was localized in wild type cells (WT), vps4Δ cells, and vps27Δ cells. Cells were counterlabeled with the endocytic tracer dye FM4-64. C, GFP-Cps1 was correctly localized to the vacuole lumen in wild type cells but not doa1Δ mutant cells, the latter of which showed accumulation of GFP-Cps1 at the limiting membrane of the vacuole. The lower panel shows sorting of Ub-GFP-Cps1 to the vacuole lumen in doa1Δ cells. Also shown are corresponding DIC images.

To confirm the interaction of Doa1 with Hse1, we exploited our previous observation that previously showed that Hse1 associates with the C-terminal residues (positions 353–622) of Doa1-Ufd3 Involved in MVB Sorting domain itself was sufficient for Ub binding. The PFU domain is not homologous to any other described domain, and the structural basis for how the PFU contributes to Ub binding is unknown. Importantly, Mullally et al. (27) found that when two conserved phenylalanine residues in the PFU domain were altered to alanine, both Ub binding and known Doa1-dependent functions were abolished. These residues were Phe426 and Phe434, which conform to a canonical polyproline SH3 ligand. Since our mapping analysis was performed by expressing Doa1 mutants in yeast, we then performed experiments to show that the interaction with the Hse1 SH3 domain was direct and independent of other yeast proteins. We were unable to produce full-length Doa1 in bacteria; however, we were able to produce Doa1 truncation mutants encompassing residues 1–433 and 1–445. Consistent with the above mapping experiments, we found that bacterially produced recombinant Doa1 1–445 was able to interact with the Hse1 SH3 domain, whereas the Doa1 1–433 (lacking the F434I/KNTXG motif) was unable to interact.

The novel SH3 ligand that we found lies within central PFU domain of Doa1. Previous experiments have shown that this domain is required for binding Ub (27, 28). These conclusions were based on deletion analysis showing that both N-terminal and C-terminal fragments of Doa1 could bind to Ub, provided they contain the central PFU domain. However, these studies did not show that the PFU domain was direct and independent of other yeast proteins. We were unable to produce full-length Doa1 in bacteria; however, we were able to produce Doa1 truncation mutants encompassing residues 1–433 and 1–445. Consistent with the above mapping experiments, we found that bacterially produced recombinant Doa1 1–445 was able to interact with the Hse1 SH3 domain, whereas the Doa1 1–433 (lacking the F434I/KNTXG motif) was unable to interact.

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The novel SH3 ligand that we found lies within central PFU domain of Doa1. Previous experiments have shown that this domain is required for binding Ub (27, 28). These conclusions were based on deletion analysis showing that both N-terminal and C-terminal fragments of Doa1 could bind to Ub, provided they contain the central PFU domain. However, these studies did not show that the PFU domain was direct and independent of other yeast proteins. We were unable to produce full-length Doa1 in bacteria; however, we were able to produce Doa1 truncation mutants encompassing residues 1–433 and 1–445. Consistent with the above mapping experiments, we found that bacterially produced recombinant Doa1 1–445 was able to interact with the Hse1 SH3 domain, whereas the Doa1 1–433 (lacking the F434I/KNTXG motif) was unable to interact.

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Vps27 (3). Therefore, we used GST alone and GST fused to the Vps27 C terminus to see if Doa1 expressed in yeast lysates could indeed associate with a Vps27-Hse1 complex. Fig. 2E shows that although Doa1 could specifically associate with GST-Vps27, the mutant Doa1ΔHse1 protein could not.

MVB Sorting Defects Caused by Loss of Doa1-Hse1 Association—We next functionally characterized the Doa1ΔHse1 and Doa1Δ2Hse1 mutants. Wild type V5 epitope-tagged DOA1 as well as the doa1ΔHse1 and doa1Δ2Hse1 alleles were cloned into low copy centromere-based plasmids under the control of the DOA1 promoter and transformed into doa1Δ mutants. Fig. 3A shows that the expression level of the three proteins was comparable, indicating that the overall integrity of the proteins was intact. We next examined Ub levels within these transformants by immunoblotting cell extracts with anti-Ub antibodies and examining the level of Ub conjugates. Consistent with previous studies, loss of Doa1 significantly reduced the level of Ub (21, 27, 31). Using our particular lysis protocol, we found very little unconjugated Ub overall, making the comparison of Ub conjugates more informative for assessing the overall level of Ub. Ub levels were restored with the V5-tagged DOA1 as well as the V5 epitope-tagged doa1ΔHse1 and doa1Δ2Hse1 alleles. These data showed that association of Doa1 with the Hse1 SH3 domain was not important for maintaining Ub levels. Furthermore, they suggested that loss of Ub binding by the PFU domain of Doa1 also did not significantly alter cellular Ub levels.

We then examined the effect the doa1ΔHse1 alleles had on sorting GFP-Cps1. Fig. 3B shows that although cells expressing the wild type V5 epitope-tagged Doa1 sorted GFP-Cps1 into the vacuole normally, cells expressing either of the doa1ΔHse1 alleles were defective and accumulated GFP-Cps1 on the limiting membrane of the vacuole. Thus, the persistence of MVB sorting defects in the doa1ΔHse1 cells despite the normal complement of cellular Ub indicated that the Hse1-Doa1 interaction fulfilled a specific function.

To further confirm that the doa1ΔHse1 alleles could fulfill other Doa1 functions, we tested them for their ability to complement the temperature-sensitive growth defect of doa1Δ mutants. Fig. 4A shows that doa1Δ cells grow poorly at elevated temperature (37 °C), consistent with previous studies. Normal growth was restored by overexpressing Ub using a 2 µ copy plasmid carrying the UBI4 gene, which encodes a tandem fusion of 5 Ubs. Furthermore, normal growth was also restored by the V5 epitope-tagged wild type DOA1 as well as both doa1ΔHse1 alleles. Thus, these data show that we were able to genetically uncouple the role of Doa1 on Ub levels and generalized stress intolerance from its effect on MVB sorting, which supports the idea that Doa1 may be directly involved in some aspect of MVB sorting.

DOA1 Interacts Genetically with VPS27—Although the above experiments indicated that Doa1 performed a specific role by its association with Hse1, we also wondered whether Doa1 might fulfill a more general role in the processing and/or sorting of Ub cargo at the MVB. Indeed, although we observed
Doa1-GFP localization to class E compartment endosomes in *vps27Δ* mutants, we also observed some localization to endosomes in *vps27Δ* mutants (Fig. 1B). Our previous studies showed that Hse1 associates with class E compartment endosomes by virtue of its association with Vps27 (54). Thus, the fact that Doa1-GFP could still associate with endosomes in the absence of Vps27-Hse1 on endosomes implied that Doa1 can use other means to associate with endosomes and thus perform functions independent of its association with Hse1.

Further genetic analysis revealed that loss of Vps27 in *doa1Δ* cells caused a severe growth phenotype (Fig. 4A). This was consistent with the idea that Doa1 function helped protect cells from the potentially toxic effects that cells might otherwise endure when MVB formation and degradation of integral membrane proteins in the vacuole is compromised. Importantly, this synthetic effect was not suppressed by overexpressing Ub, suggesting that the synthetic effect reflected a direct contribution of Doa1 function and not an indirect effect of depleted Ub levels compromising the health of *vps27Δ* mutants. As expected, transforming the *doa1Δ vps27Δ* double mutant strain with either wild type V5 epitope-tagged *DOA1* or the two *doa1Δ Hse1* alleles restored growth at high temperatures, indicating that even under these more stringent conditions, these mutant alleles largely complemented the functions of *DOA1* related to stress tolerance. Furthermore, they demonstrated that the synthetic effect we observe between *doa1Δ* and *vps27Δ* reflects a function outside of the interaction between *DOA1* and *HSE1*. We also found that the loss of Vps23, another class E Vps protein that is part of the ESCRT-I complex, as well as expression of dominant negative *VPS4* caused a similar synthetic growth defect in combination with loss of Doa1 (Fig. 4B).

**Loss of Doa1 Alters the Flux of Cellular Ubiquitin**—The specific synthetic interaction between *doa1Δ* and class E *vps* mutants that fail to sort proteins into the MVB prompted us to examine other aspects for how Doa1 could contribute to the process of lysosomal degradation. Our previous experiments focused on a subset of MVB marker proteins, such as GFP-Cps1, which represent an idealized model MVB substrate, which accesses in the MVB pathway as part of its normal mechanism for delivering itself to the vacuole lumen, where it functions (50). Likewise, other MVB marker proteins that are widely used, such as the transporters Gap1, Fur4, and Ste6, as well as the G-protein-coupled receptors Ste3 and Ste2 become rapidly ubiquitinated as part of their natural strategy for down-regulation (55). Moreover, none of these proteins is thought to access the MVB degradative pathway, because they are misfolded or damaged, as would be the case for a proportion of a wide variety of cell surface proteins that might undergo a small but steady rate of vacuolar degradation. Thus, to get a better indication on the flux of Ub cargo in general, we examined the localization of GFP-tagged Ub, which has been used in other systems to monitor the use and distribution of cellular Ub (56, 57). The GFP-Ub construct we used contained GFP fused to the N terminus of wild type Ub replete with its normal C-terminal diglycine tail required for covalent modification of proteins. Production of GFP-Ub was via the moderately strong *PRC1* promoter, which was housed on a low copy centromere-based plasmid. Fig. 5B shows that expression of GFP-Ub completely suppressed the temperature-sensitive growth defect of *doa1Δ* cells, demonstrating that GFP-Ub can functionally substitute for Ub to at least some degree.

We then examined the distribution of GFP-Ub in wild type and mutant cells. Fig. 5A shows that in wild type cells, GFP-Ub was localized diffusely to the cytosol but could also be found accumulated within the vacuole. These data are consistent with previous studies that showed that although the bulk of Ub is thought to be removed from cargo before its irreversible sorting into the MVB interior, some of it persists and is ultimately delivered to the vacuole interior (7, 8, 54, 58, 59). The significant level of GFP-Ub in the vacuoles of wild type cells implies that the MVB pathway may play an important role in regulating the
overall Ub levels within cells. Previous studies on the deubiquitinating enzyme Doa4 have supported a role for this enzyme in the latter stages of sorting Ub cargo into MVB vesicles by removing Ub from MVB cargo (8, 58, 59). Using GFP-Ub, we found direct data supporting this model, since doa4/H9004 mutants showed a massive accumulation of GFP-Ub within the yeast vacuole with concomitant depletion of Ub in the cytosol. We next examined the distribution of GFP-Ub in vps27/H9004 cells, which accumulate large endosomes as well as cargo that would otherwise be delivered to the vacuolar lumen. In these mutants we found that the large “class E compartment” endosomes, marked by the endocytic tracer dye FM4-64, also accumulated high levels of GFP-Ub. These data indicated that in vps27Δ mutants, Ub cargo becomes highly concentrated within endosomes and remains ubiquitinated. These results are consistent with similar analysis in mammalian cells, where depletion or disruption of Class E Vps protein function blocks lysosomal degradation of ubiquitinated membrane proteins and leads to the accumulation of ubiquitinated proteins on endosomes (60). Together, these data demonstrated that the GFP-Ub fusion protein reflected the expected distribution of Ub well and supported its use to monitor Ub distribution. A major difference was found when we examined the distribution of GFP-Ub in doa1Δ cells. Here we found that GFP-Ub was excluded from the vacuole, indicating that the bulk of Ub cargo that would otherwise be delivered to the vacuole was blocked by the absence of Doa1. Importantly, these experiments were done where GFP-Ub was exogenously expressed under conditions that suppress the temperature-sensitive growth defect of doa1Δ cells and thus probably do not reflect the nonspecific effect of Ub depletion. The effect of doa1Δ on Ub was not, however, observed in cells expressing the doa1ΔHse1 protein, which is unable to associate with Hse1. This is consistent with the idea that the global effects of Doa1 with regard to the overall distribution and flux of GFP-Ub into the vacuole reflect a function independent of its association with Hse1.

Effects of Doa1 Are Independent of Ufd2 and Otu1—Although Doa1 has been implicated in Ub-dependent processes, specific functions for Doa1 have been difficult to determine. Recent studies have implicated various functions for Doa1 genetically with Cdc48 and its associated co-factors Otu1 and Ufd2 (28). For instance, overexpression of Otu1 suppresses the growth defects of doa1Δ mutants, as does deletion of Ufd2. Otu1 is a Ub-specific cysteine protease, which is a member of the ovarian tumor family of deubiquitinating enzymes (61). Although the precise mechanism for how its overexpression suppresses doa1Δ growth defects is unknown, the most plausible explanation is that it can complex with Cdc48 to deubiquitinate substrates targeted to the proteasome, and increased levels of Otu1 would program more Cdc48 complexes for this purpose and decrease the flux of ubiquitinated substrates to the proteasome. Ufd2 is a “E4” Ub ligase that preferentially extends polyubiquitin chains (62). Overexpression of Ufd2 enhances proteasomal degradation, supporting the idea that...
loss of Doa1 leads to more Cdc48 complexes programmed with Ufd2, which would increase flux through the proteasome, whereas loss of Ufd2 would decrease flux through the proteasome (28). Thus, the suppression observed by Otu1 overexpression or loss of Ufd2 can be explained by a general inhibition of proteasomal degradation. Under some circumstances, particular in ubp6 mutants, proteasomes can rapidly degrade much of the cellular Ub, and inhibiting flux to the proteasome in doa1Δ mutants would probably restore Ub levels and viability at high temperature (63). Consistent with this idea are the observations that mutations which compromise proteasome catalytic activity or Cdc48 activity also suppress the growth defect of doa1Δ mutants (24, 27).

To determine whether these alternate interactions of Doa1 affected the phenotypes we observed, we analyzed the effect of deleting UFD2 and OTU1 (ovarian tumor family 1). Fig. 6A confirmed that deletion of UFD2 suppresses the temperature-sensitive phenotype of doa1Δ mutants. However, ufd2Δ mutants were unable to suppress the growth defect of doa1Δ vps27Δ double mutants. These data are consistent with the idea that the synthetic effect of doa1Δ and vps27Δ goes beyond simply the effect of lower Ub levels, since altering Ub levels directly (by overexpressing UBI4) or indirectly (by inhibiting degrada-

**FIGURE 5. Visualization of Ub distribution.** A. Wild type cells (WT) along with the indicated mutant strains were transformed with low copy plasmid expressing GFP-Ub (pGFP-Ub) and imaged. Cells were co-labeled with the endocytic tracer dye FM4–64. Shown are the green channel (GFP-Ub), red channel (FM4–64), DIC image, and a merge of the fluorescence signals. Also shown are doa1Δ mutant yeast transformed with the doa1ΔHse1 allele. B, wild type and doa1Δ mutant cells were transformed with vector only or the GFP-Ub expression plasmid. Serial dilutions of cells were plated and grown at 37 °C for 2 days.

loss of Ufd2-dependent substrates) does not suppress the growth defect. Similarly, deletion of OTU1 did not have a synthetic effect on vps27Δ mutants. We also examined deletion of UBP6, which encodes a proteasome-associated deubiquitinase whose absence accelerates the degradation of Ub by the proteasome (63). Despite the adverse effects on Ub pools that deletion of UBP6 causes, we found no synthetic effect when ubp6Δ was combined with a vps27Δ mutation. Fig. 6B shows that the distribution of GFP-Ub remained relatively unperturbed in ubp6Δ, otu1Δ, and ufd2Δ mutants. All of these cells showed both cytosolic and intravacuolar GFP-Ub, in contrast to doa1Δ cells, which excluded GFP-Ub from the vacuole. The exclusion of GFP-Ub from the vacuole of doa1Δ was also seen in doa1Δ ufd2Δ cells, showing that UFD2 loss did not suppress this defect. Interestingly, ufd2Δ mutants occasionally showed clumps of ubiquitinated proteins, which may represent aggregates of ubiquitinated cytosolic and/or membrane proteins. However, we did not pursue this observation further.

**Doa1 Helps Ubiquitinated Cargo Proteins Concentrate on Endosomes**—We next investigated how loss of Doa1 was toxic to vps27Δ mutants. Loss of Vps27, like the loss of some other class E Vps proteins, blocks delivery of ubiquitinated membrane proteins to the vacuole interior and blocks the formation of the endosomal intraluminal vesicles themselves. Despite this profound block in degrading membrane proteins, vps27Δ cells are remarkably healthy, and their viability does not significantly suffer from the accumulation of damaged proteins that would otherwise be degraded in the vacuole. Fig. 5A showed that vps27Δ mutants accumulate ubiquitinated proteins in large endosomal class E compartments, which previous studies have found also concentrate proteins destined for vacuolar degradation. Fig. 7 confirmed this by showing that the same compartments that accumulate GFP-Ub in vps27Δ mutants also accumulate Ste3 tagged with the Cherry red fluorescent protein. In stark contrast, we found that loss of DOA1 dramatically blocked the ability of ubiquitinated proteins to be concentrated in an enlarged endosomal compartment. Instead, only small puncta accumulated GFP-Ub, which showed some but not extensive co-distribution with endocytic markers. Likewise, Ste3-Cherry did not accumulate within large perivacuolar structures and was instead found at the cell surface as well as numerous small puncta, which probably corresponded to endosomes. Collectively, these data indicate that in the absence of Vps27 function, where ubiquitinated proteins are blocked from delivery to internal lumenal vesicles, they are still collected and consolidated on endosomal structures. Furthermore, they indicate a specific role for Doa1 in fostering this accumulation and imply that loss of such consolidation may be toxic.

**Certain Ubiquitinated Membrane Proteins Are Defective for MVB Sorting in Doa1 Mutants**—If Doa1 were somehow required for processing ubiquitinated membrane proteins so that they were capable of moving into intraluminal vesicles, one might expect that specific types of proteins might be affected. These proteins might include damaged or denatured proteins prone to aggregation or bulky multisubunit complexes that need disassembly before sorting. We also searched for particular proteins that could also show such a dependence on Doa1. For this, we examined a GFP-tagged form of Vph1, which
is a 100-kDa integral membrane protein of the larger vacuolar ATPase H^+ pump complex. Vph1 associates with other membrane proteins that comprise a V₀ sector of the pump, which in turn associates with a large soluble multisubunit V₁ sector (64). Although GFP-tagged Vph1 localizes exclusively to the limiting membrane of the yeast vacuole, our previous studies have shown that translational fusion of Ub to the C terminus converts this protein into an MVB cargo, which localizes to the vacuolar lumen in wild type cells (41). Thus, this protein gains access to the MVB pathway by virtue of its in-frame C-terminal Ub and is not dependent on cellular ubiquitination for its sorting. Fig. 7C shows that unlike wild type cells, doa1Δ cells were highly defective in properly sorting Vph1-GFP-Ub to the vacuole interior and instead accumulated on the limiting membrane in doa1Δ cells. Upon examining 50 cells, we found that only 4% of wild type cells showed detectable rim fluorescence on the surface of the vacuole. In contrast, 94% of the doa1Δ cells showed rim fluorescence. These data indicate that although Vph1-GFP-Ub carries its own Ub sorting signal, it is inefficiently recognized or processed in doa1Δ mutants so that it fails to be delivered to the vacuole lumen.

DISCUSSION

Doa1 is a Ub-binding Cdc48 adaptor that has been implicated in a number of biological responses to a variety of cellular stresses. Many phenotypes have been found for doa1Δ mutants, but most of these phenotypes probably stem from an indirect effect of lowered Ub levels in the cell (21). The defects we observe with regard to proper MVB sorting of ubiquitinated cargo are independent of lower Ub levels and indicate a more direct role for Doa1 in controlling some aspect(s) of the MVB sorting process. As such, we envisage two roles for Doa1: one that is served by physical connection with the Hse1-Vps27 Ub-sorting receptor and another more gen-

FIGURE 6. Specificity of doa1Δ defects. A, cells with the indicated mutations were serially diluted and plated onto minimal media and grown at the indicated temperature. B, the indicated mutants were transformed with the low copy GFP-Ub-expressing plasmid. Transformants were co-labeled with the endocytic tracer dye FM4–64 and imaged. WT, wild type.
Doa1/Ufd3 Involved in MVB Sorting

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A.

GFP-Ub  FM4-64  DIC  Merge

vps27Δ

doa1Δ vps27Δ

B.

GFP-Ub  Ste3-Ch  DIC  Merge

vps27Δ

doa1Δ vps27Δ

C.

Vph1-GFP-Ub  DIC

WT

doa1Δ

General role independent of this association that helps to process Ub cargo so that it can be concentrated on endosomes and efficiently delivered to the ESCRT sorting machinery for incorporation into lumenal vesicles of the forming MVB.

An Hse1-dependent role for Doa1 is mediated by its direct association with the SH3 domain of Hse1. The SH3 ligand we discovered in Doa1 did not conform to a canonical proline-rich motif (65) but instead a novel motif that contained bulky hydrophobic residues that probably mediate interaction with the SH3 domain surface. This region is conserved in not only other yeast orthologs of Doa1 but also the mammalian Doa1 ortholog, PLAP. Previously, we have shown that the Hse1 SH3 domain also interacts with Ubp7, a deubiquitination enzyme, and Hua1, a component of complex containing the Ub ligase Rsp5 (20). These studies indicated that both deubiquitination and Ub ligation activities associate with Hse1 help determine the sorting efficiency of cargo proteins, such as GFP-Cps1, which requires Ubiquitination to access the MVB sorting pathway. Furthermore, when association of Hse1 to Rsp5 is severed, MVB sorting of GFP-Cps1 is defective, whereas sorting is restored when Ub is fused in frame to the N terminus of GFP-Cps1. Likewise, we find that mutants of Doa1 lacking their ability to bind Hse1 have a similar defect in sorting GFP-Cps1 but no defect in sorting the Ub-GFP-Cps1 reporter protein. Thus, the function of the Doa1-Hse1 interaction is consistent with a role for Doa1 in helping potentiate the ubiquitination status of GFP-Cps1 rather than affecting the general ability of the Ub-sorting ESCRT apparatus to deliver cargo to the endosomal intraluminal vesicles. These data further show that since binding of Doa1, Hua1, and Ubp7 to the Hse1 SH3 domain is sensitive to the same mutations in the SH3 ligand binding site, all of these activities may compete at some level, implying that there may be a mechanism that regulates access to the Hse1-Vps27 complex by these factors.

Previous studies indicated that the central PFU domain of Doa1 is required for efficient binding to Ub. Rumpf and Jentsch (28) showed that strong Ub binding was contained within residues 1–494 of Doa1, whereas Mullally et al. (27) showed Ub binding with the N-terminal 450 residues as well as a C-terminal fragment encompassing residues 354–715. Combining these overlapping fragments suggests that the central PFU domain defined by residues 354–450 is required for Ub binding. In particular, Mullally et al. (27) altered Phe426 and Phe434 and showed loss of Ub binding by Doa1. We show here that a bacterially expressed PFU domain is indeed sufficient for Ub binding. Since Ub binding is a common feature among many Cdc48-associated proteins, it would be predicted to be a key functional feature of Doa1. However, our data are unclear as to what function Ub binding by the PFU domain itself confers. The SH3 ligand we identified also lies within the PFU domain and contains one of the residues, Phe434, that Mullally et al. (27) identified as important for Ub binding. We find that although the isolated PFU domain of the Doa1<sup>Hse1</sup> mutant retained its ability to bind Ub, the PFU domain from the Doa1<sup>Hse1</sup> mutant did not. However, both Doa1 mutants were able to complement the growth defect of doa1Δ mutants, in contrast to the F426A/F434A mutant previously analyzed. We do not, however, conclude that Ub binding by Doa1 is dispensable for its function, since we have found other regions within Doa1 that are also sufficient for Ub binding. Thus, as Mullally et al. (27) cautioned in their study, the F426A/F434A mutation used may have abrogated other features of the protein.

A broader function of Doa1 at the MVB, outside of its association with Hse1, was first indicated by our observation that doa1Δ cells show a severe synthetic growth defect when combined with loss of VPS27. We also found that the general flux of

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Ub into the vacuole was defective in doa1Δ cells as was the MVB sorting Vph1-GFP-Ub, a component of the multisubunit V-ATPase complex, which we modified by translationally fusing Ub onto the C terminus. Importantly, all of these defects were largely independent of the nonspecific effect of lowered Ub levels that result from Doa1 loss. For instance, the synthetic growth defect of doa1Δ vps27Δ mutants was not suppressed by Ub overexpression or loss of Ufd2, which competes with Doa1 for Cdc48 binding and may accelerate proteasomal degradation and Ub loss (28). Likewise, the block in GFP-Ub accumulation in the vacuole was performed under conditions where the GFP-Ub suppresses the Ub-dependent temperature-sensitive toxicity perhaps by allowing their aberrant functions to compromise a host of other cellular functions.

One model we favor is that Doa1 is somehow required to help process a range of Ub cargo so that it can be efficiently recognized by the downstream ESCRT machinery and packaged into intraluminal vesicles. This processing would involve cooperation with the Cdc48 ATPase, which could help tease apart ubiquitinated membrane proteins as they are delivered to the endosome. Such activity would not be required for some proteins, such as Ub-GFP-Cps1 or Ste3-GFP, which are reporters based on proteins that access the MVB pathway as part of their natural biological program for down-regulation (55). Instead, we speculate that Doa1-Cdc48 would be required by larger protein complexes as well as damaged or aggregated proteins that are rendered nonfunctional or potentially toxic. Such processing would make such cargo more accessible to the sorting machinery and help cargo concentrate on endosomal subdomains that mediate formation of lumenal vesicles. A further observation consistent with this model is that Doa1 was required for Ub cargo to accumulate in large endosomal structures in vps27Δ mutants. The doa1Δ vps27Δ mutants showed a dramatic dispersal of Ub cargo throughout the cell. Indeed, the inability to consolidate Ub cargo on endosomal subdomains may explain the severe growth defect we observe in doa1Δ vps27Δ mutants. This effect would be analogous to the toxic effects of aggresomes and other protein aggregates observed in mammalian cells, which accumulate if proteasomal degradation is compromised. Normally, these proteins are localized to the microtubule organizing center by the action of Cdc48 and HDAC6 (66, 67). HDAC6 also binds ubiquitinated proteins and mediates their transport via microtubule motors to the cell center, where they are concentrated (68). Blocking the ability of HDAC6 to consolidate these damaged proteins increases their toxicity perhaps by allowing their aberrant functions to compromise a host of other cellular functions.

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