

# A single ubiquitin is sufficient for cargo protein entry into MVBs in the absence of ESCRT ubiquitination

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**E**SCRTs (endosomal sorting complexes required for transport) bind and sequester ubiquitinated membrane proteins and usher them into multivesicular bodies (MVBs). As Ubiquitin (Ub)-binding proteins, ESCRTs themselves become ubiquitinated. However, it is unclear whether this regulates a critical aspect of their function or is a nonspecific consequence of their association with the Ub system. We investigated whether ubiquitination of the ESCRTs was required for their ability to sort cargo into the MVB lumen. Although we found that Rsp5 was the

main Ub ligase responsible for ubiquitination of ESCRT-0, elimination of Rsp5 or elimination of the ubiquitinatable lysines within ESCRT-0 did not affect MVB sorting. Moreover, by fusing the catalytic domain of deubiquitinating peptidases onto ESCRTs, we could block ESCRT ubiquitination and the sorting of proteins that undergo Rsp5-dependent ubiquitination. Yet, proteins fused to a single Ub moiety were efficiently delivered to the MVB lumen, which strongly indicates that a single Ub is sufficient in sorting MVBs in the absence of ESCRT ubiquitination.

## Introduction

Many ESCRT (endosomal sorting complex required for transport) components have Ubiquitin (Ub)-binding domains (UBDs) that contribute to their ability to sort ubiquitinated membrane proteins into multivesicular bodies (MVBs). Like other proteins that contain UBDs, ESCRTs can be ubiquitinated by a variety of mechanisms mediated by the UBD itself in a process termed “coupled ubiquitination” (Haglund and Stenmark, 2006). Moreover, ESCRT-0 and ESCRT-I associate with Ub-ligases and deubiquitinating enzymes that might influence ESCRT activity by modulating ESCRT ubiquitination (Kato et al., 2000; Amit et al., 2004; McCullough et al., 2004; Kim et al., 2007; Ren et al., 2007). Coupled ubiquitination has been proposed to drive critical aspects of ESCRT function by altering intermolecular interactions between ESCRT proteins that house UBDs, or triggering cargo release by causing intramolecular association between the UBD and attached Ub (Hicke et al., 2005; Komada and Kitamura, 2005; Piper and Katzmann, 2007; Williams and Urbé, 2007; Roxrud et al., 2010). However, definitive evidence that such UBD-dependent ubiquitination drives a physiologically relevant aspect of ESCRT function has been difficult to obtain. In addition, it remains unclear what type of Ub-based sorting motif actually serves as the signal for incorporation

into MVBs. Structural studies showing tandem arrays of UBDs within ESCRTs as well as functional studies in cells deficient in polyubiquitin assembly suggest that short K63-linked chains of Ub constitute the operative sorting signal recognized by ESCRT UBDs (Erpapazoglou et al., 2008; Kulathu et al., 2009; Lauwers et al., 2009; Paiva et al., 2009; Ren and Hurley, 2010). However, limitations from these correlative experiments complicate such conclusions about potential regulatory mechanisms that control how cargo is identified, recognized, and processed by the endocytic system. Using a variety of approaches, we found that ubiquitination of the ESCRT apparatus is not required for its ability to sort monoubiquitinated cargo into the MVB.

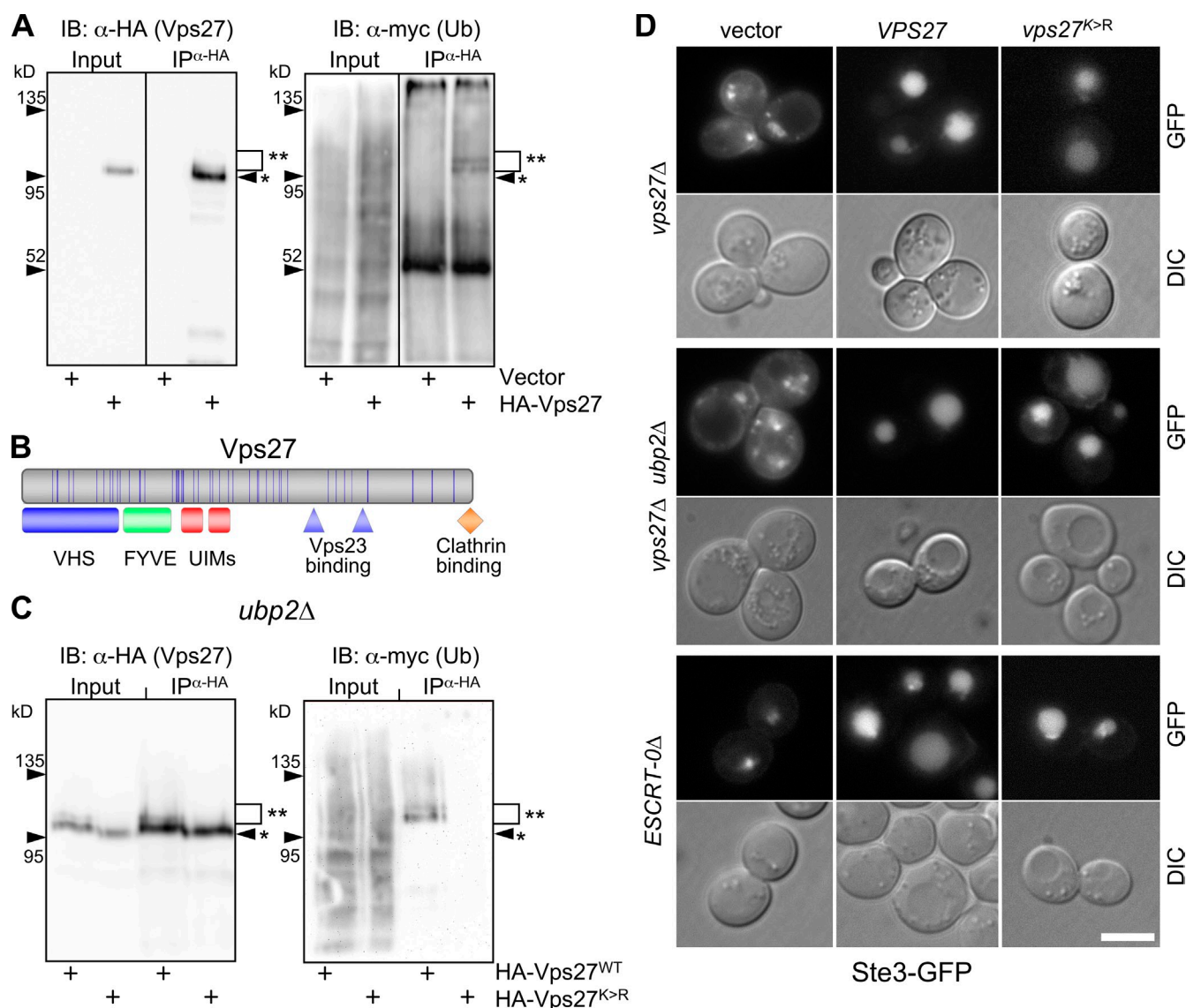
## Results and discussion

We found that the yeast Hrs homologue Vps27 is ubiquitinated by immunoprecipitating HA-Vps27 from denatured lysates and immunoblotting for Ub. The major proportion of HA-Vps27 migrated at ~96 kD, whereas a less prominent higher molecular weight band (~105 kD) corresponded to ubiquitinated HA-Vps27 (Fig. 1 A).

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Abbreviations used in this paper: DUB, deubiquitinating peptidase; ESCRT, endosomal sorting complex required for transport; MVB, multivesicular body; Ub, Ubiquitin; UBD, Ub-binding domain.

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**Figure 1. Vps27 lacking lysine residues is not ubiquitinated but still functions in MVB cargo sorting.** (A) Ubiquitination of Vps27 was assessed in cells expressing HA-Vps27 and myc-Ub. Vps27 immunoprecipitates (IP) from denatured cell lysates were immunoblotted (IB) for HA-Vps27 (anti-HA) or Ub (anti-myc). Input represents a 5% equivalent. Black lines indicate that dividing lanes have been spliced out. (B) Domain organization of yeast ESCRT-0 protein Vps27. Blue lines show positions of the 47 lysines. (C) Ubiquitination of wild-type and lysine-less Vps27 (HA-Vps27<sup>WT</sup> and HA-Vps27<sup>K>R</sup>) was assessed in *ubp2* $\Delta$  cells expressing myc-Ub. Anti-HA (Vps27) immunoprecipitates (IP) were immunoblotted (IB) for HA-Vps27 (anti-HA) or Ub (anti-myc). Input represents a 5% equivalent. \*, expected molecular weight; \*\*, ubiquitinated forms. (D) Sorting of Ste3-GFP in *vps27* $\Delta$ , *vps27* $\Delta$  *ubp2* $\Delta$ , or *vps27* $\Delta$  *hse1* $\Delta$  (*ESCRT-0* $\Delta$ ) cells transformed with plasmids expressing HA-Vps27<sup>WT</sup> or HA-Vps27<sup>K>R</sup>. Bar, 5  $\mu$ m.

To assess whether ESCRT-0 ubiquitination is required for MVB sorting, we generated a Vps27<sup>K>R</sup> mutant lacking all 47 lysine residues in the full-length protein (Fig. 1 B). All of the Lys residues were altered because coupled ubiquitination can promiscuously use a variety of lysines (Polo et al., 2002; Miller et al., 2004; Woelk et al., 2006; Uchiki et al., 2009). Ubiquitination of Vps27<sup>K>R</sup> was undetectable, even in cells lacking the deubiquitinating enzyme Ubp2, which antagonizes the HECT-type Ub ligase Rsp5 responsible for a wide variety of ubiquitination events within the endocytic pathway (Kee et al., 2005, 2006; Rotin and Kumar, 2009). In contrast, a sizable proportion of wild-type Vps27 was ubiquitinated in *ubp2* $\Delta$  cells because  $\sim$ 20% was found in the higher  $\sim$ 105-kD form (Fig. 1 C). We also found that both wild type and the Vps27<sup>K>R</sup> protein were equally stable (Fig. S1 A).

To determine if loss of Vps27 ubiquitination impairs MVB sorting, we examined the localization of Ste3-GFP, a GFP-tagged G protein-coupled receptor normally targeted to MVBs. Cells lacking Vps27 accumulated Ste3-GFP on the vacuole limiting membrane and in endosomes. However, cells with either wild-type Vps27 or Vps27<sup>K>R</sup> sorted Ste3-GFP into the vacuole interior (Fig. 1 D). Both Vps27 proteins functioned equally in cells also lacking Ubp2, which implies that although wild-type Vps27 is ubiquitinated to a greater extent in these cells, the MVB sorting process is normal. Under conditions where MVB sorting is less efficient, the Vps27<sup>K>R</sup> mutant performed like wild type, which indicates that loss of Vps27 ubiquitination does not increase sorting efficiency (Fig. S1 B).

Yeast ESCRT-0 contains both Vps27 and the STAM homologue Hse1 (Bilodeau et al., 2002). STAM also undergoes ubiquitination (Row et al., 2006), raising the possibility that ubiquitination of Hse1 may regulate ESCRT-0 activity. To test this, we used the SEY6210 parental yeast strain, which requires Vps27 but not Hse1 for MVB sorting (unpublished data and Emr, S.D., personal communication). We found that Ste3-GFP sorting was normal in *hse1Δ vps27<sup>K>R</sup>* cells where all the lysines of ESCRT-0 are eliminated (Fig. 1 D).

Hrs associates with HECT-type Ub ligases such as Nedd4 and AIP4, which promote ubiquitination of Hrs both in vivo and in vitro (Katz et al., 2002; Marchese et al., 2003; Woelk et al., 2006). The yeast homologue of Nedd4 is Rsp5, which interacts with ESCRT-0 (Ren et al., 2007), and is a prime candidate for the ligase that ubiquitinates Vps27. Vps27 ubiquitination was severely attenuated in *rsp5-1* temperature-sensitive cells at 37°C to ~10%, the level observed in wild-type cells (loss of Ub-Vps27 was 91% ± 2.7% SD, *n* = 3; Fig. 2 A). These data indicated that Rsp5 is the major Ub ligase responsible for ESCRT-0 ubiquitination, which suggests that Rsp5 inhibition might be a means of assessing the role of ESCRT ubiquitination.

We next examined MVB sorting of Mup1-GFP in cells with compromised Rsp5 activity. We first examined sorting in cells lacking many of the adaptors Rsp5 requires to modify cargo (Nikko and Pelham, 2009), and in which we found ubiquitination of Mup1-GFP to be severely compromised (Fig. 2 B and S1 C). We also assessed sorting in *rsp5-1* cells at 37°C (Fig. 2 C). Either method of inhibiting Rsp5 dramatically blocked sorting of Gap1-GFP and Mup1-GFP, which is consistent with previous studies demonstrating a requirement for Rsp5-mediated ubiquitination (Rotin and Kumar, 2009). However, simply adding Ub in frame restored sorting of Mup1-GFP, indicating that ESCRT-dependent MVB sorting operates normally as long as the requirement for cargo ubiquitination is satisfied.

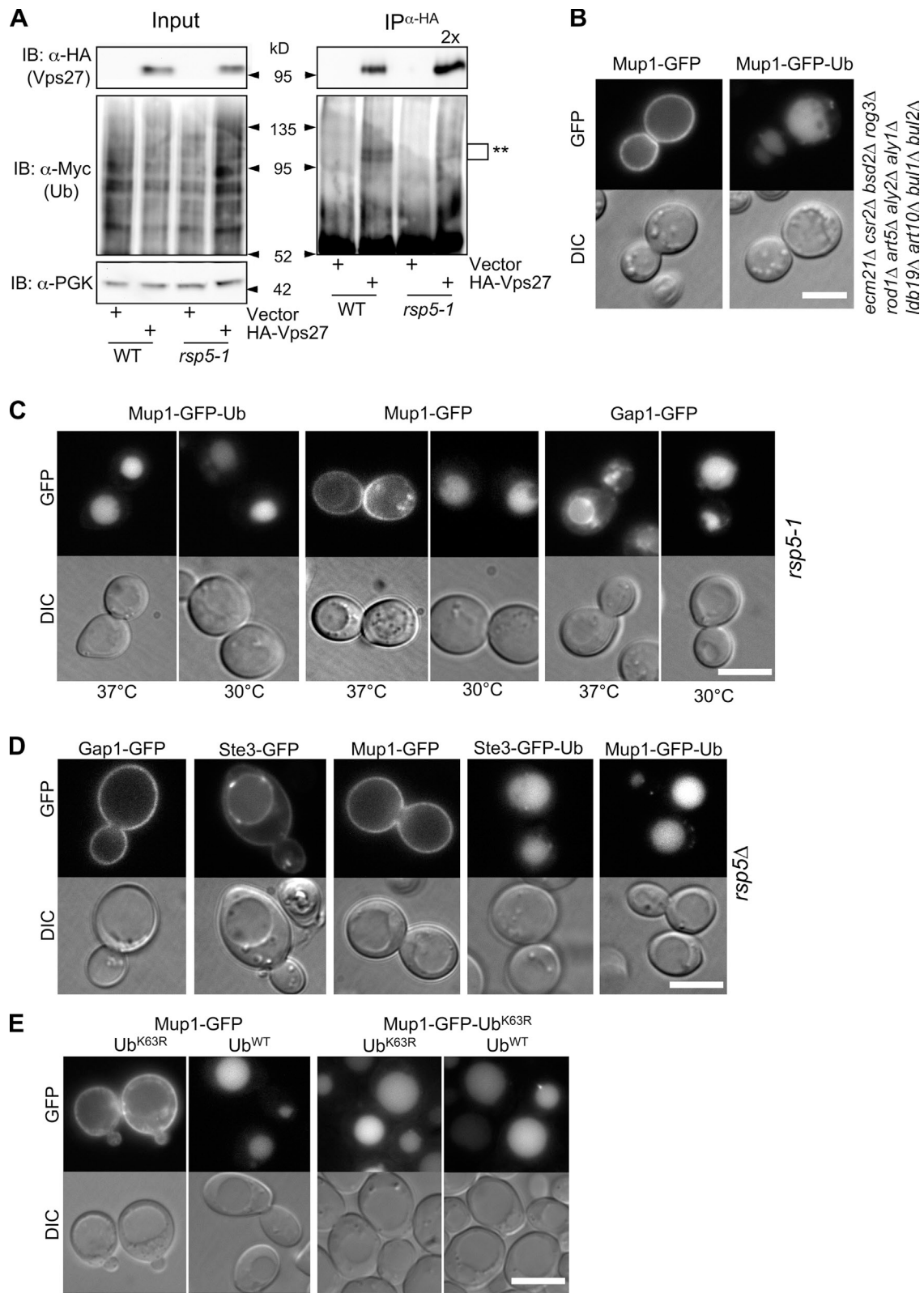
Previous studies indicate that *rsp5-1* cells might still harbor residual Rsp5 activity at 37°C (Kee et al., 2006), perhaps enough to mediate a low level of ESCRT ubiquitination. RSP5 is essential; however, its essential function can be bypassed by expressing a truncated version of the Mga2 transcription factor that undergoes Rsp5- and proteasome-dependent cleavage (Hoppe et al., 2000; Chellappa et al., 2001). We generated *rsp5Δ* cells, which grew slowly at 30°C and were inviable at 37°C (Fig. S1 D). Sorting of proteins such as Mup1-GFP, Ste3-GFP, and Gap1-GFP along the MVB pathway was completely blocked in *rsp5Δ* cells. In contrast, fusion of Ub onto the C terminus of either Ste3-GFP or Mup1-GFP restored sorting (Fig. 2 D), which indicates that ESCRT ubiquitination by Rsp5 is not required for MVB sorting.

Rsp5 favors formation of K63-linked polyubiquitin chains (Kee et al., 2005, 2006), and several studies suggest that K63-linked Ub chains, rather than a single mono-Ub, serve as a necessary signal for entry into the MVB sorting pathway (Lauwers et al., 2010). Indeed, cargo fused to a single Ub, which induces MVB sorting, might still undergo additional K63 ubiquitination (Reggiori and Pelham, 2001; Urbanowski and Piper, 2001). However, our data demonstrate that fusion of a single Ub to a cargo protein is sufficient for MVB sorting in cells where

Rsp5-dependent K63 polyubiquitination is absent. We further tested the role of K63 Ub chains in cells that were entirely incapable of generating K63 polyubiquitin chains (Fig. 2 E). The sole source of Ub in these cells was Ub<sup>K63R</sup>, which was a condition previously shown to cause severe MVB sorting defects for membrane proteins such as Gap1, Jen1, and Sit1 (Erpapazoglou et al., 2008; Lauwers et al., 2009; Paiva et al., 2009). Similarly, we found that Mup1-GFP was localized exclusively to the cell surface and intracellular puncta in Ub<sup>K63R</sup> cells. However, Mup1-GFP-Ub<sup>K63R</sup>, containing a single in-frame Ub also lacking K63, sorted normally to the vacuole interior. Collectively, these results indicate that a single Ub is sufficient for cargo to sort into the MVB lumen. They also imply that the sorting defects resulting from depriving cells of K63-linked Ub chains may arise from global changes that disrupt trafficking pathways upstream of the MVB or perturb the dynamic cycle of cargo ubiquitination and deubiquitination, from which Mup1-GFP-Ub is immune. In addition, fusion of Ub may route cargo more efficiently to endosomes, circumventing many of the obstacles that other cargos might encounter in Ub-K63-deficient cells during their endocytic itinerary (see supplemental material). Our data clearly agree with previous studies establishing some type of role for K63-linked chains in regulating MVB sorting (Erpapazoglou et al., 2008; Lauwers et al., 2009; Paiva et al., 2009). However, our finding that fusing Ub<sup>K63R</sup> is sufficient for MVB sorting suggests that K63-linked chains need not be on cargo themselves. It still remains possible that the sorting of some cargo proteins benefits from the added avidity such chains have for the ESCRT Ub receptors. K63-linked chains may allow cargo to resist the effects of deubiquitinating enzymes as they travel through the endocytic pathway.

The preceding data indicate that neither removing ubiquitinatable lysine residues from ESCRT-0 nor eliminating the major Ub ligase responsible for the majority of ESCRT-0 ubiquitination critically interferes with ESCRT function. However, these approaches have caveats. First, although eliminating Rsp5 dramatically reduces ESCRT-0 ubiquitination, other Ub-ligases might still modify ESCRTs at undetectable levels. This is a distinct possibility because coupled ubiquitination in vitro can be mediated by many different ligases or by E2-conjugating enzymes alone (Hoeller et al., 2007; Uchiki et al., 2009). Second, although we eliminated all of the lysine residues within ESCRT-0, it is possible that another protein associated with the canonical ESCRT-0 may be a relevant Ub acceptor. Finally, Rsp5 is capable of ubiquitinating the N termini of proteins in vitro (Kim and Huibregtse, 2009), whereas other ligases can ubiquitinate cysteine, serine, and threonine residues (Cadwell and Coscoy, 2005; Wang et al., 2007). Thus, the prospect of removing all ubiquitinatable residues within ESCRT-0 and its associated proteins is daunting and leaves no way to control for structural perturbations caused by the mutations themselves. Therefore, we devised a different method to render proteins resistant to ubiquitination. Because in vivo removal of Ub is catalyzed by deubiquitinating peptidases (DUBs), we reasoned that it was possible to generate nonubiquitinatable proteins by fusing them to the catalytic domain of a DUB (Fig. 3 A). We chose three catalytic domains from cysteine-based DUBs that can be





**Figure 2. Cells lacking Rsp5 activity still sort ubiquitinated cargo proteins into vacuoles.** (A) Wild-type and temperature-sensitive *rsp5-1* cells expressing HA-Vps27 and Myc-Ub were shifted to 37°C for 1 h. Input lysates and Vps27 immunoprecipitates (IP) were immunoblotted (IB) for HA (Vps27) and Myc (Ub). Twice the amount of the anti-HA IP from *rsp5-1* cells was loaded. \*, expected molecular weight; \*\*, ubiquitinated forms. (B) Localization of Mup1-GFP and Mup1-GFP-Ub in mutant EN67 cells lacking multiple Rsp5 adaptors. Cells were grown in YPD and copper for 4 h to produce Mup1 fusion proteins from the *CUP1* promoter. (C) Sorting of Gap1-GFP, Mup1-GFP, and Mup1-GFP-Ub after Rsp5 inactivation. Cells (*rsp5-1*) were shifted to 37°C for 1 h and grown in YPD in the presence of 100  $\mu$ M CuCl<sub>2</sub> to induce production of GFP-tagged proteins under control of the *CUP1* promoter. (D) Localization of Gap1-GFP, Ste3-GFP, Mup1-GFP, Ste3-GFP-Ub, or Mup1-GFP-Ub in *rsp5Δ*-null cells grown in YPD + 25  $\mu$ M CuCl<sub>2</sub>. (E) Localization of Mup1-GFP or Mup1-GFP-Ub<sup>K63R</sup> in cells expressing wild-type (WT) Ub (*SUB492*) or K63R Ub (*SUB493*) as their sole source of Ub. Bars, 5  $\mu$ m.

inactivated by mutating their catalytic cysteine (Reyes-Turcu et al., 2009): the C-terminal catalytic domain from yeast Ubp7, the UL36 Ub-specific protease domain from type I Herpes virus (Gredmark et al., 2007; Kattenhorn et al., 2005), and the M48 Ub-specific protease domain of murine cytomegalovirus (Schlieker et al., 2007).

When these catalytic domains were fused to GFP-tagged membrane proteins that normally undergo MVB sorting, their localization to the vacuole interior was completely blocked, with localization primarily at the cell surface (Fig. 3 A). The effect on Fur4 and Gap1 was consistent with the strong plasma membrane localization of Fur4 and Gap1 lacking their ubiquitination sites, as described in previous studies (Blondel et al., 2004; Lauwers et al., 2009). Fusion of a DUB domain did not adversely affect the normal localization of Vph1 to the vacuole limiting membrane. In addition, coexpressing DUB-tagged membrane proteins did not affect the sorting of other proteins that normally follow a Ub-dependent pathway to the vacuole interior (Fig. 3 B). As expected, the stability of Ste3-GFP fused to an active DUB was substantially higher than when fused to a catalytically inactive DUB (Figs. 3 C and S1 E). Similarly, fusion of Ste3 to active UL36 catalytic domain abolished detectable ubiquitination (Fig. 3 D). Despite its steady-state localization to the cell surface, we found that Ste3-GFP-DUB was still internalized (Fig. 3 E) because it accumulated in endosomal compartments of *rcy1Δ* mutants where recycling is blocked (Wiederkehr et al., 2000; Galan et al., 2001). However, Ste3-GFP-DUB did not accumulate in the late endosome-like class E compartments that accumulate in *vps4Δ* cells (Babst et al., 1997). These data imply that Ste3 internalizes in a Ub-independent manner and that without Ub, it quickly recycles to the cell surface via early endosomal compartments without reaching late endosomal compartments.

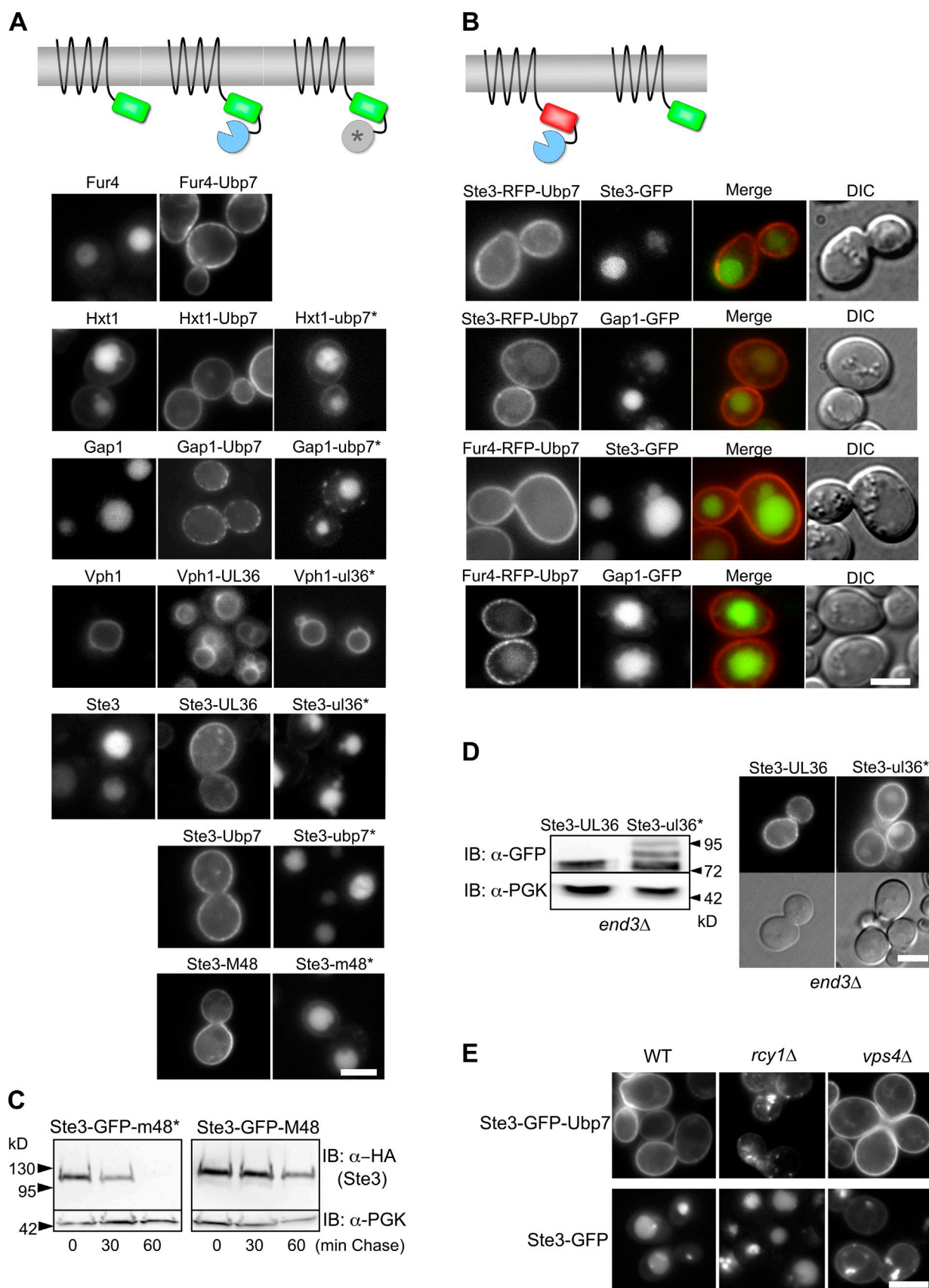
Equipped with a new method to immunize proteins against ubiquitination, we reexamined the role of ESCRT ubiquitination (Fig. 4). Fusing DUB catalytic domains to ESCRTs should not only prevent ubiquitination of the ESCRTs themselves, but if suitably robust, the DUB should also strip Ub from cargo proteins and block their entry into MVBs (Fig. 4 A). Moreover, if ESCRT ubiquitination is not required for the sorting process, cargo fused to Ub so that it cannot be removed by DUB activity should be able to sort normally. Fusion of UL36 catalytic domain to ESCRT-0 subunit Hse1 prevented accumulation of ubiquitinated Hse1 and Vps27 (Fig. 4 B). Expression of Hse1-DUB proteins in wild-type cells caused a dramatic block in the sorting of Ste3, Fur4, Gap1, and Mup1, all of which rely on Rsp5-dependent ubiquitination for sorting into the vacuole (Fig. 4 C). Importantly, both Ste3-GFP-Ub and Mup1-GFP-Ub, which contained a single “non-deubiquitinatable” Ub, were properly sorted to the vacuole interior. We also fused DUB catalytic domains to Vps23 and Mvb12, two ESCRT-I subunits thought to act as a downstream Ub cargo receptor and whose mammalian counterparts also undergo ubiquitination (Kim et al., 2007; Tsunematsu et al., 2010). The Vps23-DUB and Mvb12-DUB fusions also interrupted the Ub- and Rsp5-dependent sorting of membrane cargo to the MVB lumen yet did not block sorting of cargo with an in-frame fusion of Ub. All of the MVB sorting defects were abolished when the

catalytic activity of the DUB domains was inactivated (Fig. 4 C), and no nonspecific defects in MVB sorting were observed when we expressed catalytic domains fused to His3, a cytosolic protein within the histidine biosynthesis pathway (Struhl and Davis, 1977). All of the DUB fusion proteins were expressed to comparable levels (Fig. S1, F and G), and none caused a vacuolar protein secretion (VPS) phenotype (Fig. S2 A), which is consistent with ESCRT function being maintained. We also examined the effect of fusing a DUB catalytic domain to Gga2, a Ub-binding clathrin adaptor protein thought to sort ubiquitinated proteins from the TGN to endosomes (Scott et al., 2004). GGA proteins may also contribute to Ub-dependent transport of proteins from the cell surface to endosomes in yeast; however, evidence for a clear role in this pathway is obscured because ESCRT-0 also operates in this pathway in parallel (Erpapazoglou et al., 2008; Deng et al., 2009; Lauwers et al., 2009). The dominant-acting Gga2-DUBs dramatically blocked sorting of Gap1, Mup1, and Fur4 but did not affect that of Ste3 to the vacuole, which indicates that yeast Gga proteins may also mediate Ub-specific sorting of a subset of cargos as they travel between the cell surface (Puertollano and Bonifacio, 2004).

The ESCRT-DUB fusion proteins could also functionally substitute for their wild-type counterparts in the MVB sorting process (Fig. 5, A–C). Neither cells with Hse1-UL36 as their sole source of Hse1 nor cells with Vps23-UL36 as their sole source of Vps23 sorted Ste3-GFP or Gap1-GFP into the vacuole lumen. However, both cell types were able to sort Ste3-GFP-Ub and Fur4-GFP-Ub fusion proteins. Mvb12-Ubp7 and Mvb12-UL36 were also able to functionally substitute for wild-type Mvb12 to promote the sorting of Ste3-GFP-Ub, but they effectively blocked sorting of Fur4-GFP (Fig. 5 A). Because loss of Mvb12 results in only mild MVB sorting defects (Curtiss et al., 2007), we assessed Mvb12-DUB function in cells that also had mutations in the UBDs of Vps23 and Vps36, which yield a severe synthetic class E VPS phenotype (Shields et al., 2009). These data indicate that the ESCRT-fused DUB domains are robust enough to deubiquitinate cargo and prevent ubiquitination of ESCRTs, yet are still able to sort Ub-tagged cargo.

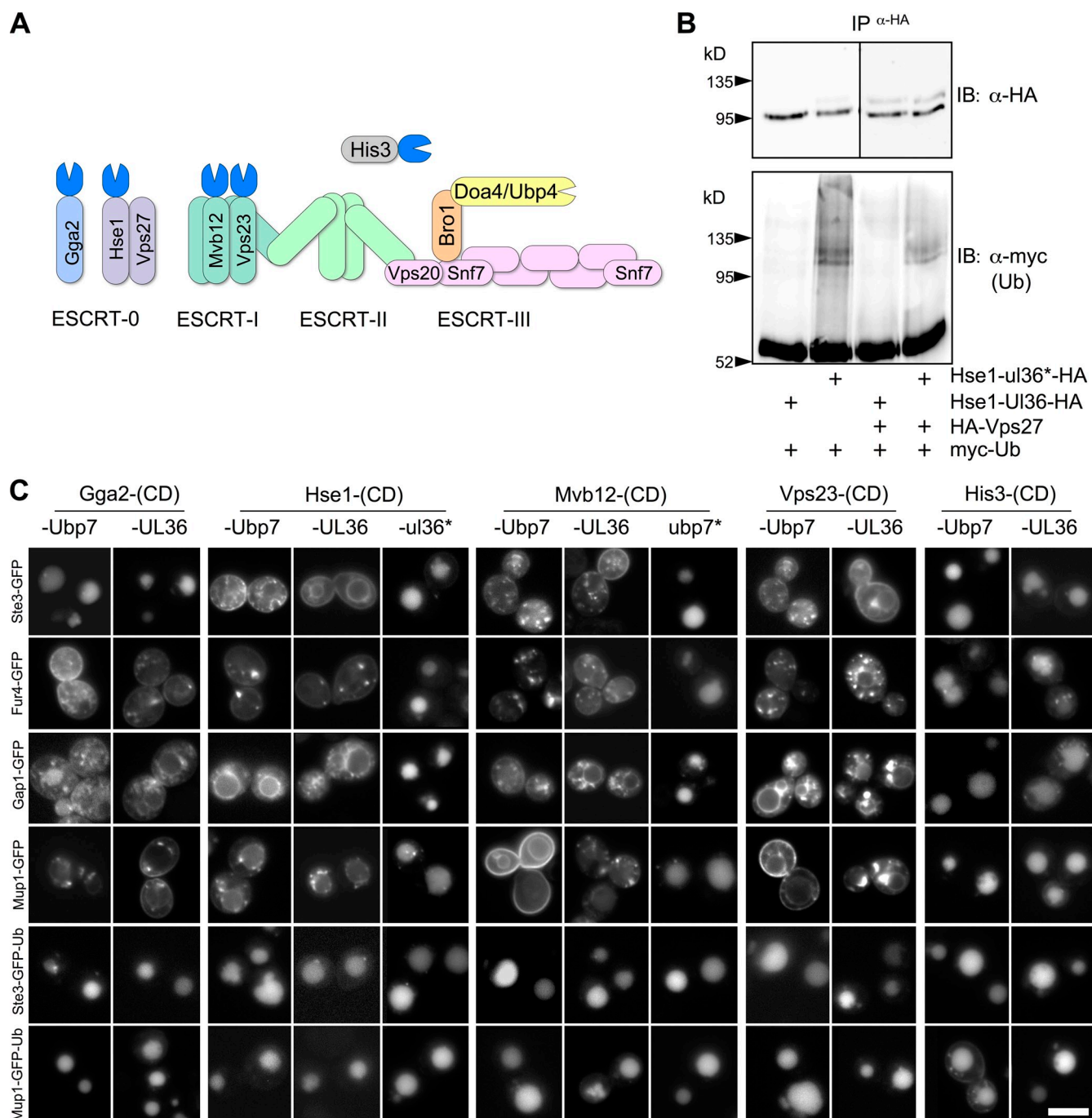
We then evaluated whether fusion of Rsp5 to a DUB catalytic domain would create a dominant “reversal” enzyme that could deubiquitinate Rsp5 substrates *in vivo*. Expressing Rsp5 fused to the catalytic domain of Ubp7 effectively blocked sorting of many Rsp5 substrates (Fig. 5 D). No effect was observed with Rsp5 fused to an inactive catalytic domain (Fig. 5 D) expressed at comparable levels (Fig. S2 B). Expressing Rsp5-DUB also inhibited growth, which is consistent with the essential role of Rsp5 for viability (Fig. S2 C). Although Rsp5-DUB removed Ub from endogenous Rsp5 substrates and blocked their sorting, cargo such as Ste3-GFP-Ub and Mup1-GFP-Ub sorted into the vacuole normally, which supports the idea that their single in-frame Ub is sufficient for their sorting. Furthermore, these data indicate that any deubiquitination of ESCRTs by Rsp5-DUB does not hamper ESCRT sorting activity.

Potentially, the organization of the ESCRT apparatus could be regulated by ubiquitination by altering inter- and intramolecular interactions among its UBD-containing components. Cumulative loss of multiple UBDs within the ESCRTs



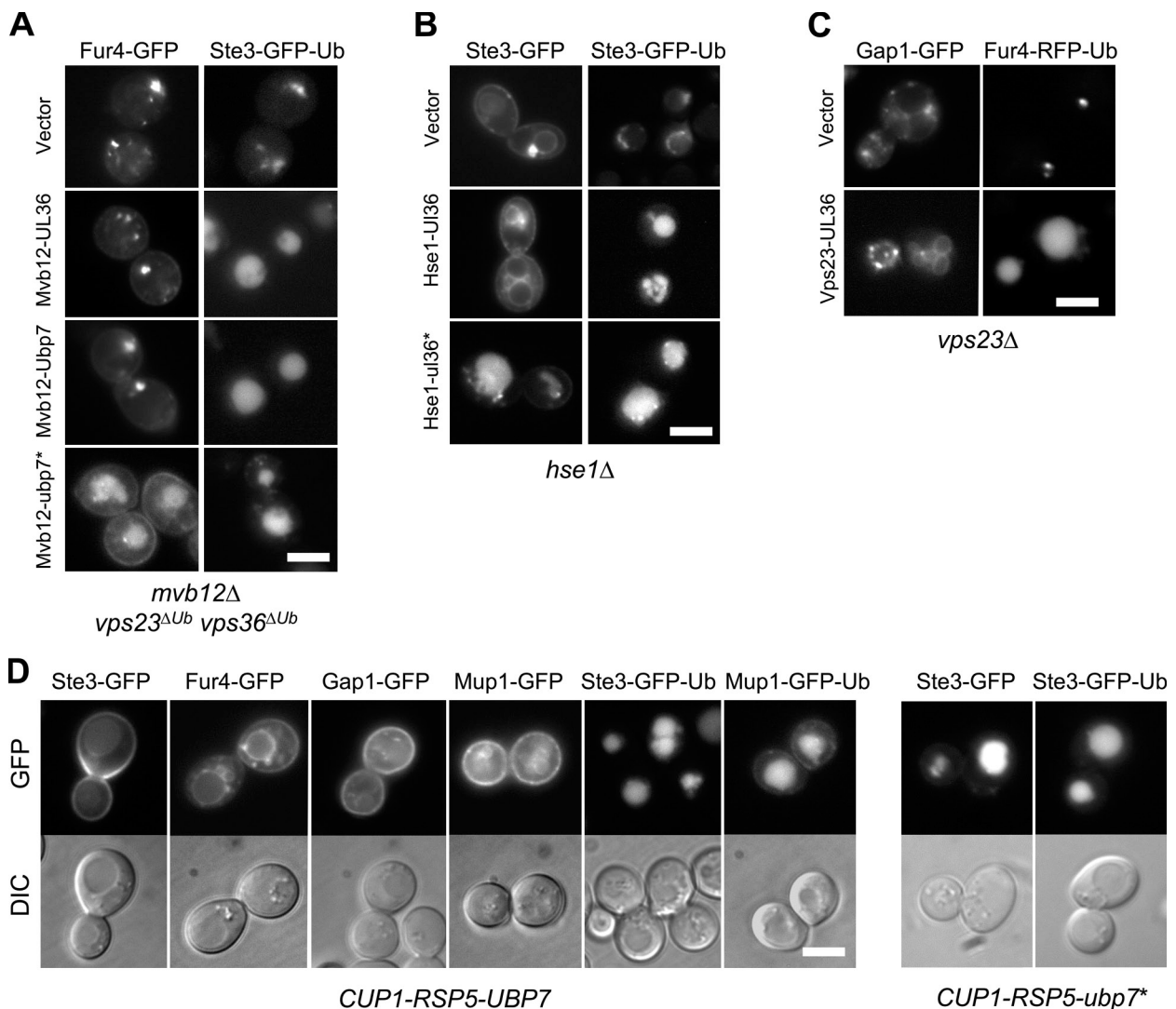
**Figure 3. Fusing cargo to the catalytic domain from deubiquitinating enzymes blocks ubiquitination and trafficking to the vacuole.** (A) Schematic of DUB catalytic domain fusion proteins (top). Localization of the indicated GFP-tagged proteins fused to either enzymatically active or inactive (\*) catalytic domains of Ubp7, HSV-1 UL36, and mCMV M48 (bottom). (B) The effect of DUB fusion proteins (Ste3-RFP-Ubp7 or Fur4-RFP-Ubp7) on coexpressed Gap1-GFP or





**Figure 4. Fusion of DUBs to ESCRT proteins inhibits their ubiquitination but allows vacuolar sorting of Ub fusion cargo proteins.** (A) Schematic of the ESCRT sorting apparatus showing subunits fused to different DUB catalytic domains. (B) Cells expressing myc-Ub and either Hse1-UL36-HA or Hse1-ul36\*-HA (inactive) were transformed with vector alone or with plasmids expressing HA-Vps27. Lysates were immunoprecipitated (IP) with anti-HA antibodies. Immunoprecipitates were immunoblotted (IB) for HA (top) to confirm isolation of HA-tagged proteins; immunoprecipitates were immunoblotted with anti-myc to assess ubiquitination. Fusion of active UL36 abolished ubiquitination of either HA-tagged Hse1 or Vps27 proteins. Black lines indicate that dividing lanes have been spliced out. (C) ESCRT subunits, GGA, or His3 fused to the indicated active (Ubp7 or UL36) or inactive (ubp7\* or ul36\*) catalytic domains (CD) were expressed in wild-type cells. Cells coexpressed the indicated GFP-tagged MVB cargo. Bar, 6  $\mu$ m.

Ste3-GFP. (C) Cycloheximide chase with cells expressing HA-tagged Ste3-GFP-m48\* and Ste3-GFP-M48, which carried active and inactive DUB domains, respectively. Aliquots were removed at the indicated times after cycloheximide addition (100  $\mu$ g/ml) and immunoblotted for HA and PGK. (D) Ste3-GFP-UL36 or Ste3-GFP-ul36\* (with an inactive catalytic domain) were expressed in *end3Δ* cells and immunoblotted with anti-GFP antibodies (left) or localized (right). (E) Localization of Ste3-GFP or Ste3-GFP-Ubp7 in wild type, *rcy1Δ* cells, and *vps4Δ* cells. Bars, 5  $\mu$ m.



**Figure 5. The effect of localizing DUB domains on ESCRTs and the Ub ligase Rsp5.** (A) Complementation of *mvb12Δ* by Mvb12-UL36. The localization of Fur4-GFP and Ste3-GFP-Ub in *mvb12Δ vps23Δ<sup>Ub</sup> vps36Δ<sup>Ub</sup>* cells carrying vector plasmid alone or plasmids expressing Mvb12 fused to active (UL36, Ubp7) or inactive (ubp7\*) DUB domains is shown. (B) Complementation of *hse1Δ* by Hse1-UL36 and Hse1-ul36\* (inactive). The localization of Ste3-GFP and Ste3-GFP-Ub in *hse1Δ* cells carrying the indicated plasmids is shown. (C) Complementation of *vps23Δ* by Vps23-UL36. Localization of Gap1-GFP and Fur4-RFP-Ub in *vps23Δ* cells carrying indicated plasmids is shown. (D) Wild-type cells expressing Rsp5 fused to active (Ubp7) or inactive (ubp7\*) catalytic DUB domain and the indicated GFP-tagged MVB cargo proteins. Expression of Rsp5 fusions were from the *CUP1* inducible promoter, induced 5 h before microscopy (for Rsp5-Ubp7) or overnight (for Rsp5-ubp7\*). Bars, 5  $\mu$ m.

results in a general loss of function that is not confined to cargo recognition alone (Shields et al., 2009). These data imply that UBDs may contribute to the productive organization of the ESCRT apparatus in a manner similar to what has been proposed for the internalization apparatus (Dores et al., 2010). Alternatively, ESCRT ubiquitination might mediate an intramolecular interaction that prevents interaction with Ub cargo, providing a mechanism for cargo release. However, ESCRTs as well as many other Ub-binding proteins undergo ubiquitination by virtue of simply having UBDs, and this may not reflect a genuine regulatory mechanism. We found that preventing ubiquitination of ESCRTs did not diminish or enhance their ability to sort Ub cargo or affect their ability to release cargo because cargo did not accumulate on endosomes, nor did nonubiquitinatable ESCRT-0

accumulate in the vacuole lumen (Fig. S2 D). It remains possible that future experiments will demonstrate that other ESCRT components also become ubiquitinated. However, the best candidate to mediate such ubiquitination is Rsp5, and we have taken several approaches to demonstrate that Rsp5-dependent ubiquitination is not required for the MVB sorting process once a single Ub modifies cargo. Determining whether such coupled ubiquitination is correlative or indeed causes a specific and physiologically relevant regulatory mechanism in vivo requires that the process of coupled ubiquitination be uncoupled. The multiple approaches used here, including the approach of tethering DUB activity to particular protein complexes, provides a comprehensive complementary strategy for assessing a physiological role for such ubiquitination events.



## Materials and methods

### Materials, yeast strains, and plasmids

Chemicals, growth methods, and other general procedures were used as described previously (Shields et al., 2009). Growth in the presence and absence of copper was done with plates containing 100  $\mu$ M CuCl<sub>2</sub> or bathocuproine disulfonate (BCS), respectively. Anti-CPY, GFP, HA, and Pho8 were used as described previously (Ren et al., 2007). Purchased antibodies were: anti-myc monoclonal (A00704; GenScript), anti-myc polyclonal (18826-01; QED Biosciences), anti-Ub (P4D1, sc-8017; Santa Cruz Biotechnology, Inc.), and anti-HA monoclonal (MMS-101R; Covance). Yeast strains are described in Table I and plasmids are described in Table II. The *vps27Δ ubp2Δ* strain was created by inserting the *HIS5* gene (Guldener et al., 2002) into *VPS27* within the *ubp2Δ* mutant from the BY4742 MAT $\alpha$  yeast deletion collection (Giaever et al., 2002). The *hse1Δ vps27Δ* strain was made by inserting the *TRP1* gene into the *HSE1* locus (Ren et al., 2007). The *rsp5Δ*-null strains were made from a parental W303 strain, which contained a disruption of the genomic *RSP5* gene and contained a *URA3*-based low copy plasmid housing the wild-type *RSP5* gene (pRS416-*RSP5*). This strain was transformed with a low copy plasmid expressing a truncated Mga2 protein (Chellappa et al., 2001), and an *ADE2* selectable plasmid expressing a GFP-tagged reporter protein. Strains that lost the plasmid-borne *RSP5* gene were selected on plates containing 5-fluoroorotic acid and oleic acid that were incubated at 25°C (Wang et al., 1999), and loss of *RSP5* was confirmed by lack of growth at 37°C (Hoppe et al., 2000; Chellappa et al., 2001). DUB catalytic domains were residues 561–1,071 of yeast Ubp7, the N-terminal residues 15–260 (UL36) of the type I Herpes virus VP1/2 tegument protein (Kattenhorn et al., 2005; Gredmark et al., 2007), and residues 4–234 of M48 Ub-specific protease (Schlieker et al., 2007). M48 is active toward both K48- and K63-linked Ub chains, whereas UL36 was originally reported to show specificity to K48-linked Ub chains (Kattenhorn et al., 2005). However, recent studies demonstrate that UL36 cleaves both K48- and K63-linked chains (Kim et al., 2009), which is consistent with our own observations (Fig. S2 E).

For experiments requiring detection of intact Ubp7 fusions, we used an altered form of the Ubp7 catalytic domain. The catalytic domain of Ubp7 undergoes autocleavage, resulting in a clipped form that apparently remains associated and active. The catalytically dead Ubp7 C<sub>618</sub>S mutant does not undergo this cleavage (Fig. S2 B). This characteristic has been observed for other catalytic domains such as Usp1, but is not required for its enzymatic activity (Huang et al., 2006). Ubp7 has a GG motif that is localized in a predicted loop that would lie between  $\beta$ 11 and  $\beta$ 12 of the USP fold “fingers” subdomain of the USP catalytic domain. Mutation of G to S (corresponding to G<sub>932</sub> in full-length Ubp7) abolished autocleavage while preserving enzyme activity (Fig. S2 B).

### Fluorescence microscopy

Cells were grown to mid-log phase, briefly centrifuged, and resuspended in 0.2% Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, 0.2% NaF, and 100 mM Tris, pH 8.0. GFP and mCherry in cells at 22°C were imaged with an epifluorescence microscope (BX60; Olympus) with a 100 $\times$  objective lens, NA 1.4. Images were captured with a cooled charge-coupled device camera (Orca R2; Hamamatsu Photonics) using iVision-Mac software (Biovision Technology) and scaled to the brightest and darkest pixel intensity with Photoshop software (Adobe). For cells containing copper-inducible expression plasmids (containing the *CUP1* promoter), cells were grown overnight, diluted in fresh media, and then induced with 100  $\mu$ M CuCl<sub>2</sub>. Induction of *CUP1*-dependent expression in *rsp5-1* cells was similar except cells were grown at permissive (30°C) and nonpermissive temperatures (37°C) during copper-triggered induction lasting 2 h. Cells were grown in media replete with methionine (yeast peptone dextrose [YPD] or synthetic defined [SD] media with 20  $\mu$ g/ml methionine [Met]) for all Mup1 localization experiments except where noted. Like other transporters such as Gap1, Mup1 can follow a Ub-dependent route from the cell surface into the MVB lumen that is triggered by the addition of its substrate methionine. Alternatively, Mup1 can also bypass the plasma membrane on its route from the Golgi to MVB/endosomes via a GGA-dependent sorting step if synthesized in the presence of methionine (Fig. S2 F). Unlike Gap1, however, fusion of Ub to the C terminus of Mup1-GFP still allowed for proper folding and exit out of the ER, thus providing a tool to bypass the need for ligase-mediated ubiquitination.

### Immunoblotting and immunoprecipitation

Whole cell denatured lysates were generated by resuspending pelleted cells in 0.2 M NaOH. After 2 min, cells were pelleted and resuspended in

8 M urea, 5% SDS, 10% glycerol, and 50 mM Tris, pH 6.8, and heated at 70°C for 5 min as described previously (Ren et al., 2008). Samples for immunoblotting Mup1-GFP proteins were prepared similarly, although they were not heated. For immunoprecipitations, whole cell denatured lysates were diluted with 10 volumes of 100 mM Tris, pH 7.5, and 0.4% Triton X-100. Samples were incubated with polyclonal anti-HA for 90 min at 25°C. Immune complexes were captured on fixed *Staphylococcus aureus* cells (IgG-sorb; Enzyme Center). Beads were washed thrice in 0.1 M Tris, pH 7.5, 0.4% Triton X-100, and 0.1% SDS, and analyzed by SDS-PAGE and immunoblotting. Cells carrying the *CUP1*-myc-Ub plasmid were grown overnight in 50  $\mu$ M CuCl<sub>2</sub> before lysis. Immunoprecipitation from *rsp5-1* cells was performed as in Fig. 1 except that cell cultures were divided and grown at 37°C for 1 h before lysis. To assess ubiquitination of Ste3-GFP fused to active and inactive UL36, experiments were performed in *end3Δ* cells defective for internalization, which facilitated our comparison because they accumulate proteins in a ubiquitinated state at the cell surface (Kölling and Hollenberg, 1994).

### Protease treatment

Recombinant UL36 catalytic domain was generated by expressing UL36 from pET151 in BL21 (DE3) star cells. Enzyme was purified from bacterial lysates (French press; Slm-Aminco) using Cobalt Talon resin (Takara Bio, Inc.). In vitro deubiquitinating assays were performed at 37°C for 30 min in PBS containing 0.2 mM DTT. Ub chains (1  $\mu$ g of K48 or K63; Boston Biochem) were mixed with recombinant UL36<sup>CD</sup> in the presence and absence of 1  $\mu$ M Ub-Aldehyde (Boston Biochem). Reactions were terminated by adjusting samples to 2% SDS and heating at 100°C for 10 min.

For protease protection assays, cells (~15 OD) were grown to mid-log phase. Cells were spheroplasted in 1.4 M sorbitol, 50 mM Tris, pH 8, 10 mM Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, and 1 mM DTT containing Zymolyase (AMS Biotechnology) for 1 h. Cells were purified over a cushion of 1.4 M sucrose by centrifugation and lysed at 4°C with 12 passes through a 26-g needle in sorbitol and 50 mM Tris, pH 8.0. Debris was removed from lysates by centrifugation at 1,000 g for 5 min. The resulting supernatant was then incubated in the presence and absence of trypsin (Sigma-Aldrich). Samples were adjusted to 2.5% SDS and 4 M urea, and heated to 100°C for 5 min to halt protease activity.

### Online supplemental material

Fig. S1 describes the additional effects of Vps27<sup>K>R</sup>, the effect of Rsp5 adaptors on ubiquitination of Mup1, and the expression levels of the ESCRT-DUB fusion proteins. Fig. S2 shows the effect of ESCRT-Dubs on CPY sorting, the autocleavage of the Ubp7 catalytic domain, the activity of UL36 toward both K48- and K63-linked Ub chains, and the intracellular itinerary of key cargo proteins used in this study. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.201008121/DC1>.

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Table I. Plasmids used

Plasmids	Description	Use	Source
pRS315	Centromere containing low copy yeast shuttle plasmids ( <i>LEU2</i> and <i>URA3</i> )	Figs. 1 (A, C, and D), 2 A, 5 (A–C), and S2 A	Sikorski and Hieter, 1989
pAC376	Yep351 (2 $\mu$ <i>LEU2</i> ) expressing myc-tagged Ub from <i>CUP1</i> promoter	Figs. 1 (A and C), 2 A, and 4 B	A. Cooper (Garvan Institute, Sydney Australia)
pRSmga2N $\Delta$ tm	<i>HIS3</i> low copy plasmid containing a truncated MGA2 gene lacking C-terminal 329 codons that include the membrane-spanning domain.	Figs. 2 D and S1 D	Chellappa et al., 2001
pYM-N36	nourseothricin-resistance marker natMX		Janke et al., 2004
Genomic DNA	HSV-1 (strain F)		Ejercito et al., 1968
pPL4149	pRS316 expressing HA-Vps27 from <i>MET25</i> promoter	Figs. 1 (A, C, and D), 2 A, 4 B, and S1 (A and B)	This study
pPL4150	pRS316 expressing HA-Vps27 <sup>K-R</sup> (where all K residues are altered to R) from <i>MET25</i> promoter.	Figs. 1 (C and D) and S1 (A and B)	This study
pPL967	pRS315 expressing Ste3-GFP from <i>STE3</i> promoter	Figs. 1 D, 2 D, 3 (A and E), 4 C, 5 (B and D), and S2 F	Urbanowski and Piper, 2001
pPL991	pRS313 expressing Ste3-GFP from <i>STE3</i> promoter	Fig. 1 D	Urbanowski and Piper, 2001
pPL2334	pRS315 expressing Gap1-GFP from <i>CUP1</i> promoter	Figs. 2 (C and D), 3 (A and B), 4 C, and 5 (C and D)	This study
pPL3484	pRS315 expressing Ste3-GFP-Ub from <i>STE3</i> promoter	Figs. 4 C and 5, A, B, and D	This study
pPL3797	pRS315 expressing Fur4-GFP from <i>CUP1</i> promoter	Figs. 3 A, 4 C, and 5, A and D	This study
pPL3931	ADE2 conversion of pPL3484 (ADE2 replacement of <i>LEU2</i> ) expressing Ste3-GFP-Ub from <i>STE3</i> promoter	Fig. 2 D	This study
pPL3962	pRS315 expressing Hxt1-GFP from <i>HXT1</i> promoter	Fig. 3 A	This study
pPL4144	pRS315 expressing Fur4-mCherry-Ub from <i>CUP1</i> promoter	Fig. 5 C	This study
pPL4146	pRS315 expressing Mup1-GFP from <i>CUP1</i> promoter	Figs. 2 (B, C, and E), 4 C, and 5D	This study
pPL4147	pRS315 expressing Mup1-GFP-Ub from <i>CUP1</i> promoter	Figs. 2 (B–D), 4 C, 5 D, and S2 F	This study
pPL4214	ADE2 conversion of pPL967 (ADE2 replacement of <i>LEU2</i> ) expressing Ste3-GFP from <i>STE3</i> promoter	Fig. 2 D	This study
pPL4215	ADE2 conversion of pPL2334 (ADE2 replacement of <i>LEU2</i> ) expressing Gap1-GFP from <i>CUP1</i> promoter	Fig. 2 D	This study
pPL4216	ADE2 conversion of pPL4146 (ADE2 replacement of <i>LEU2</i> ) expressing Mup1-GFP from <i>CUP1</i> promoter	Fig. 2 D	This study
pPL4217	ADE2 conversion of pPL4147 (ADE2 replacement of <i>LEU2</i> ) expressing Mup1-GFP-Ub from <i>CUP1</i> promoter	Fig. 2 D	This study
pPL4252	natMX conversion of pPL4146 (natMX replacement of <i>LEU2</i> ) expressing Mup1-GFP from <i>CUP1</i> promoter	Figs. 2 E and S2 F	This study
pPL4320	pRS315 expressing Mup1-GFP-Ub <sup>K63R</sup> from <i>CUP1</i> promoter		This study
pPL4342	natMX conversion of pPL4320 (natMX replacement of <i>LEU2</i> ) expressing Mup1-GFP-Ub <sup>K63R</sup> from <i>CUP1</i> promoter	Fig. 2 E	This study
pPL3584	pRS316 expressing Mvb12-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter; contains residues 561–1071 of Ubp7	Figs. 4 C, 5 A, and S1 F	This study
pPL3609	pRS316 expressing Ste3-mCherry-Ubp7 <sup>CD</sup> from <i>STE3</i> promoter	Fig. 3 B	This study
pPL3669	pRS316 expressing Fur4-GFP-Ubp7 <sup>CD</sup> from <i>CUP1</i> promoter	Fig. 3 B	This study
pPL3810	pRS316 expressing Fur4-mCherry-Ubp7 <sup>CD</sup> from <i>CUP1</i> promoter	Fig. 3 B	This study
pPL3876	pRS316 expressing Ste3-GFP-Ubp7 <sup>CD</sup> from <i>STE3</i> promoter	Figs. 3 A and S1 E	This study
pPL3900	pRS316 expressing Gap1-GFP-Ubp7 <sup>CD</sup> from <i>CUP1</i> promoter	Fig. 3 A	This study
pPL3964	pRS316 expressing Hxt1-GFP-Ubp7 <sup>CD</sup> from <i>HXT1</i> promoter	Fig. 3 A	This study
pPL3986	pRS316 expressing Ste3-GFP-Ubp7 <sup>CD(G&gt;S)</sup> -3xHA from <i>STE3</i> promoter; contains residues 561–1,071 of Ubp7 with mutations G932S	Figs. S1 E and S2B	This study
pPL3890	pRS316 expressing Ste3-GFP-3xHA from <i>STE3</i> promoter	Fig. S1 E	This study
pPL3722	pRS316 expressing His3-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter. Contains residues 561–1071 of Ubp7 followed by 3xHA epitope	Figs. 4C, S1 F, and S2, A and C	This study
pPL3723	pRS316 expressing Hse1-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, S1 F, and S2 A	This study
pPL3724	pRS316 expressing Vps23-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, S1 F, and S2 A	This study
pPL3742	pRS316 expressing Rsp5-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 5 D, and S2 B	This study
pPL3746	pRS316 expressing Gga2-Ubp7 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, S1 F, and S2 A	This study

Table 1. **Plasmids used** (Continued)

Plasmids	Description	Use	Source
pPL3643	pRS316 expressing Ste3-GFP-ubp7 <sup>CD</sup> * from <i>STE3</i> promoter; contains residues 561–1071 of Ubp7 with C618S mutation	Fig. 3 A	This study
pPL3776	pRS316 expressing Mvb12-ubp7 <sup>CD</sup> *-3xHA from <i>CUP1</i> promoter	Figs. 4 C, 5 A, S1 F, and S2 A	This study
pPL3901	pRS316 expressing Gap1-GFP-ubp7 <sup>CD</sup> * from <i>CUP1</i> promoter	Fig. 3 A	This study
pPL4371	pRS316 expressing Hxt1-GFP-ubp7 <sup>CD</sup> * from <i>HXT1</i> promoter	Fig. 3 A	This study
pPL3994	pRS316 expressing Ste3-GFP-UL36 <sup>CD</sup> from <i>STE3</i> promoter; contains 561–591 of Ubp7 (Ubp7 linker)-and residues 15–260 of HSV1 UL36	Fig. 3, A and D	This study
pPL4012	pRS316 expressing Hse1-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 (B and C), 5 B, S1 G, and S2 A	This study
pPL4013	pRS316 expressing Mvb12-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, 5 A, S1 G, and S2 A	This study
pPL4014	pRS316 expressing His3-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, S1 G, and S2 A	This study
pPL4015	pRS316 expressing Vps23-GFP-UL36 <sup>CD</sup> from <i>VPS23</i> promoter	Fig. 5 C	This study
pPL4017	pRS316 expressing Gga2-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 C, S1 G, and S2 A	This study
pPL4140	<i>HIS3</i> conversion of pPL4012 expressing Hse1-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Fig. 4 B	This study
pPL4247	pRS316 expressing Vps23-UL36 <sup>CD</sup> -3xHA from <i>CUP1</i> promoter	Figs. 4 and S2, B and C	This study
pPL4058	pRS316 expressing Ste3-GFP-ubp7 <sup>CD</sup> * from <i>STE3</i> promoter; contains 561–591 of Ubp7 (Ubp7 linker) and residues 15–260 of HSV1 UL36 with mutation of C40S in UL36	Fig. 3, B and D	This study
pPL4131	pRS316 expressing Hse1-ubp7 <sup>CD</sup> *-3xHA from <i>CUP1</i> promoter	Figs. 4 C, 5 B, and S2, B and C	This study
pPL4141	<i>HIS3</i> conversion of pPL4131 expressing Hse1-ubp7 <sup>CD</sup> *-3xHA from <i>CUP1</i> promoter	Fig. 4 C	This study
pJLU40	pRS316 expressing Vph1-GFP from <i>VPH1</i> promoter	Fig. 3 A	This study
pPL4056	pRS316 expressing Vph1-GFP-UL36 <sup>CD</sup> from <i>VPH1</i> promoter	Fig. 3 A	This study
pPL4202	pRS316 expressing Vph1-GFP-ubp7 <sup>CD</sup> * from <i>VPH1</i> promoter	Fig. 3 A	This study
pPL4218	pRS316 expressing Ste3-GFP-M48 <sup>CD</sup> from <i>STE3</i> promoter; contains 561–591 of Ubp7 (Ubp7 linker) AGDRDIV-mCMV M48 (residues 4–234)-VAITHPLR.	Fig. 3 A	This study
pPL4219	pRS316 expressing Ste3-GFP-m48 <sup>CD</sup> * from <i>STE3</i> promoter; contains linker residues 561–591 of Ubp7, residues 15–21 of UL36, residues 4–234 of mCMV M48, and the mutations C23A, L110A, Y113A, and Y192A followed by residues (253–260 of UL36)	Fig. 3 A	This study
pPL3974	pET151D expressing UL36 <sup>CD</sup> (residues 15–260 of HSV1 UL36)	Fig. S2 E	This study
pPL2554	<i>HIS3</i> conversion of pPL2356 ( <i>HIS3</i> replacement of <i>URA3</i> ) expressing GFP-Cps1 from <i>Cps1</i> promoter	Fig. S1 B	This study
pPL4500	pRS316 expressing Ste3-GFP-M48-3xHA from <i>STE3</i> promoter	Fig. 3 C	This study
pPL4502	pRS316 expressing Rsp5-ubp7*-3xHA from <i>CUP1</i> promoter	Figs. 5 D and S2 B	This study
pPL4513	pRS316 expressing Ste3-GFP-m48*-3xHA from <i>STE3</i> promoter	Fig. 3 C	This study
pPL4515	pRS16 expressing Ste3-GFP-ubp7*-3xHA from <i>STE3</i> promoter	Fig. S1 E	This study
pPL4253	<i>natMX</i> conversion of pPL4147 ( <i>natMX</i> replacement of <i>LEU2</i> ) expressing Mup1-GFP-Ub from <i>CUP1</i> promoter	Fig. S2 F	This study
pPL4248	<i>natMX</i> conversion of pPL967 ( <i>natMX</i> replacement of <i>LEU2</i> ) expressing Ste3-GFP from <i>STE3</i> promoter	Fig. S2 F	This study
pPL4249	<i>natMX</i> conversion of pPL3484 ( <i>natMX</i> replacement of <i>LEU2</i> ) expressing Ste3-GFP-Ub from <i>STE3</i> promoter	Fig. S2 F	This study



Table II. Yeast strains used

Yeast strain	Description	Use	Source
BY4742	MAT $\alpha$ <i>his3<math>\Delta</math>0 leu2<math>\Delta</math>0 lys2<math>\Delta</math>0 ura3<math>\Delta</math>0</i>	Throughout	Brachmann et al., 1998
SF838-9D	MAT $\alpha$ <i>leu2-3,112 ura3-52 his4-519 ade6 pep4-3</i>	Fig. S2 D	Raymond et al., 1992
SEY6210	MAT $\alpha$ <i>leu2-3,112 ura3-52 his3-<math>\Delta</math>200 trp1-<math>\Delta</math>901 lys2-801 suc2-<math>\Delta</math>9 mel</i>	Throughout	Banta et al., 1988
PLY3909	MAT $\alpha$ <i>hse1<math>\Delta</math>::TRP1 vps27<math>\Delta</math>::LEU2 ura3-52 his3-<math>\Delta</math>200 lys2-801 suc2-<math>\Delta</math>9 pep4</i>	Fig. 1 D	This study
PLY1776	MAT $\alpha$ <i>vps27<math>\Delta</math>::LEU2 leu2-3,112 ura3-52 his3-<math>\Delta</math>200 trp1-<math>\Delta</math>901 lys2-801 suc2-<math>\Delta</math>9 mel</i>	Fig. S1 B	This study
PLY2858	MAT $\alpha$ <i>vps27<math>\Delta</math>::LEU2 ura3-52 his3-<math>\Delta</math>200 lys2-801 suc2-<math>\Delta</math>9 pep4</i>	Fig. S1 B	This study
GW004	MAT $\alpha$ <i>rsp5<math>\Delta</math>::LEU2, pRS416-RSP5 leu2-3112,ade2-,his3-11, trp1-1,can1-100; derived from W303</i>	Fig. 2 D	Wang et al., 1999
PLY441 vpl23-5 (vps27 $\Delta$ )	MAT $\alpha$ <i>leu2-3,112, ura3-52, his4, ade6 vpl23-5 (vps27<math>\Delta</math>) pep4-3</i>	Fig. S2 D	Raymond et al., 1992
BY4742 YOR124C	MAT $\alpha$ <i>ubp2<math>\Delta</math>::Kan<sup>r</sup> his3<math>\Delta</math>0, leu2<math>\Delta</math>0, lys2<math>\Delta</math>0, ura<math>\Delta</math>0</i>	Fig. 1 C	Winzeler et al., 1999
PLY3827	MAT $\alpha$ <i>ubp2<math>\Delta</math>::Kan<sup>r</sup> vps27<math>\Delta</math>::HIS3 leu2<math>\Delta</math>0, lys2<math>\Delta</math>0, ura<math>\Delta</math>0</i>	Fig. 1 D	This study
RH1800	MAT $\alpha$ <i>leu2 his4 ura3 bar1</i>	Fig. 2 A	Munn et al., 1999
LHY23	MAT $\alpha$ <i>rsp5-1 leu2 ura3 his3 trp1 lys2 bar1 GAL</i>	Fig. 2, A and C	Dunn and Hicke, 2001
KLY201	MAT $\alpha$ <i>end3<math>\Delta</math>::Kan<sup>r</sup> his3<math>\Delta</math>0 leu2<math>\Delta</math>0 lys2<math>\Delta</math>0 ura3<math>\Delta</math>0</i>	Figs. 3 D and S2 F	Liu et al., 2007
BY4742 YJL204C	MAT $\alpha$ <i>rcy1<math>\Delta</math>::Kan<sup>r</sup> his3<math>\Delta</math>0 leu2<math>\Delta</math>0 lys2<math>\Delta</math>0 ura3<math>\Delta</math>0</i>	Fig. 3 E	Winzeler et al., 1999
vpl4-21	MAT $\alpha$ <i>vps4<math>\Delta</math> leu2-3,112, ura3-52, his4, ade6 (SF838-9D)</i>	Fig. 3 E	Raymond et al., 1992
PLY3623	MAT $\alpha$ <i>mbv12<math>\Delta</math> vps36-<math>\Delta</math>NZF::TRP1::vps36<math>\Delta</math>::HIS vps23-<math>\Delta</math>Ub (F52Q117W125) leu2-3,112 ura3-52 his3-<math>\Delta</math>200 trp1-<math>\Delta</math>901 lys2-801 suc2-<math>\Delta</math>9 mel</i>	Fig. 5 A	Shields et al., 2009
BY4742 YCL008C	MAT $\alpha$ <i>vps23<math>\Delta</math>::KanR his3<math>\Delta</math>0 leu2<math>\Delta</math>0 lys2<math>\Delta</math>0 ura3<math>\Delta</math>0</i>	Fig. 5 C	Winzeler et al., 1999
BY4742 YHL002W	MAT $\alpha$ <i>hse1<math>\Delta</math>::KanR his3<math>\Delta</math>0 leu2<math>\Delta</math>0 lys2<math>\Delta</math>0 ura3<math>\Delta</math>0</i>	Fig. 5 B	Winzeler et al., 1999
EN67	MAT $\alpha$ <i>ecm21::G418 csr2::G418 bsd2 rog3::natMX rod1 ygr068c aly2 aly1 ldb19 ylr392c bul1 bul2::HIS his3 ura3 leu2</i>	Figs. 2 B and S1C	Nikko and Pelham, 2009
SUB492	MAT $\alpha$ <i>lys2-801 leu2-3, 112 ura3-52 his3-200 trp1-1[am] ubi1-1::TRP1 ubi2-2::ura3 ubi3-<math>\Delta</math> ubi4-<math>\Delta</math>::LEU2 [pUB39][pUB100]; pUB39 is 2<math>\mu</math> plasmid with CUP1-Ub (wild type); pUB100 the is plasmid that expresses RPL40A tail</i>	Fig. 2 E	Spence et al., 1995
SUB493	Isogenic to SUB492 except pUB197 replaces pUB39; pUB197 is 2 $\mu$ plasmid with CUP1-Ub (K63R)	Fig. 2 E	Spence et al., 1995
PLY3378	MAT $\alpha$ <i>end3::KANR gga1<math>\Delta</math>::TRP1 gga2<math>\Delta</math>::HIS3</i>	Fig. S2 F	Bilodeau et al., 2004

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