Roles of Hsl1p and Hsl7p in Swe1p Degradation: beyond Septin Tethering

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The morphogenesis checkpoint in *Saccharomyces cerevisiae* couples bud formation to the cell division cycle by delaying nuclear division until cells have successfully constructed a bud. The cell cycle delay is due to the mitosis-inhibitory kinase Swe1p, which phosphorylates the cyclin-dependent kinase Cdc28p. In unperturbed cells, Swe1p is degraded via a mechanism thought to involve its tethering to a cortical scaffold of septin proteins at the mother-bud neck. In cells that experience stresses that delay bud formation, Swe1p is stabilized, accumulates, and promotes a G2 delay. The tethering of Swe1p to the neck requires two regulators, called Hsl1p and Hsl7p. Hsl1p interacts with septins, and Hsl7p interacts with Swe1p; tethering occurs when Hsl1p interacts with Hsl7p. Here we created a version of Swe1p that is artificially tethered to the neck by fusion to a septin so that Swe1p no longer requires Hsl1p or Hsl7p for its localization to the neck. We show that the interaction between Hsl1p and Hsl7p, required for normal Swe1p degradation, is no longer needed for septin-Swe1p degradation, supporting the idea that the Hsl1p-Hsl7p interaction serves mainly to tether Swe1p to the neck. However, both Hsl1p and Hsl7p are still required for Swe1p degradation, implying that these proteins play additional roles beyond localizing Swe1p to the neck.

Yeast cells in the environment are exposed to variations in temperature, osmolarity, and nutrient conditions that can delay bud formation. A common response to sudden (presumably stressful) environmental changes is to depolarize the actin cytoskeleton and transiently halt bud growth until the cells adapt to the stress (6, 11). During this interval, there is a concomitant delay of the cell cycle in G2 phase, enforced by the morphogenesis checkpoint, which maintains coordination between the budding and nuclear cycles and prevents the formation of binucleate cells (15). The morphogenesis checkpoint cell cycle delay requires the Wee1-family kinase Swe1p, which inhibits the mitotic Clb/cyclin-dependent kinase (CDK) complex by phosphorylating a conserved tyrosine residue on the CDK, Cdc28p (13, 16, 27). The delay is also thought to involve the inhibition of the counteracting Cdc25-family phosphatase Mih1p, which would otherwise dephosphorylate Cdc28p (12). The best-characterized aspect of the morphogenesis checkpoint concerns the degradation of Swe1p. Swe1p degradation is halted in response to stress (26), promoting a G2 delay, and this has served as a valuable assay to dissect how stresses impact the cell cycle. Here we have tested a prevailing hypothesis for the role of two proteins, Hsl1p and Hsl7p, that are thought to be primary transducers of stress signals to control Swe1p degradation (7, 19, 21, 28).

Swe1p accumulates periodically in the cell cycle, peaking in late S/G2 phase and quickly in G2/M phase. Swe1p abundance is controlled at the levels of transcription (*SWE1* mRNA accumulates in a sharp pulse in late G1 phase and disappears in S phase) and degradation (Swe1p is degraded slowly in G1/S phase and quickly in G2/M phase) (26, 27). The phosphorylation state of Swe1p also changes during the cell cycle, with multiple sites being phosphorylated in G1/M phase by Clb2p/CDK and the polo family kinase Cdc5p (1, 24). Phosphorylation by the CDK primes Swe1p for subsequent phosphorylation by Cdc5p, which is necessary for Swe1p degradation (1). Following its hyperphosphorylation, Swe1p is probably ubiquitylated by the redundant E3 ubiquitin ligases Dma1p and Dma2p, which target it for degradation by the proteasome (23).

Swe1p degradation is also correlated with the subcellular localization of Swe1p. In G1 phase, Swe1p is localized primarily in the nucleus and is stable (17, 26). As the cell builds a bud in S phase, some of the Swe1p accumulates at the bud neck. Neck localization relies on the interaction of Swe1p with Hsl7p (19, 21), which itself localizes to the neck through its interaction with the septin-binding kinase Hsl1p (17, 19, 25). The septins are conserved, filament-forming proteins that assemble a cortical scaffold at the cytoplasmic face of the plasma membrane at the mother-bud neck (29). The genetic perturbation of septin scaffold assembly results in a Swe1p-mediated delay of mitosis, suggesting that septin-mediated Swe1p neck localization is important for Swe1p degradation (3, 17).

Hsl1p and Hsl7p are both required for Swe1p degradation, and mutations that specifically disrupt the Hsl1p-Hsl7p or Hsl7p-Swe1p interactions result in Swe1p delocalization and stabilization (7, 19, 21). These findings suggest that Hsl1p and Hsl7p tether Swe1p to the bud neck, leading to its degradation. Because both Clb2p/CDK (2) and Cdc5p (1) also accumulate at the bud neck, it is attractive to speculate that tethering serves to co-concentrate Swe1p together with its upstream kinases, thereby enhancing the Swe1p phosphorylations that target it for degradation.

The tethering model for Swe1p degradation presents an elegant explanation for how yeast cells couple budding to nuclear division, as the neck exists only after bud emergence, so the degradation of the Swe1p mitotic inhibitor would depend on prior budding. After budding, some stresses can transiently displace Swe1p (and possibly also Hsl7p) from the neck (8, 17), potentially explaining why such stresses can stabilize Swe1p to delay mitosis even after bud emergence. However, actin depolymerization in budded cells does not immediately
displace Swe1p from the neck (17), and the basis for Swe1p stabilization in those cells remains unclear.

A prediction of the tethering model is that the need for Hsl1p and Hsl7p in Swe1p degradation could be bypassed if Swe1p were to be tethered directly to the septins at the bud neck. Here we tested this prediction by fusing Swe1p to the septin Cdc3p. The fusion protein was degraded normally in G2/M phase, but al-

### Materials and Methods

**Yeast strains and plasmids.**

Yeast strains are in the BF264-15DU background and are listed in Table 1. The creation of the following alleles was described previously:

- **swe1**
- **hsl1K110R**
- **hsl7**
- **CDC28E12K**

To create green fluorescent protein (GFP)-SWE1-myc and CDC3-GFP-SWE1-12×myc plasmids, we began with plasmid pDLB2340 (14), which contains GFP-coding sequences inserted at codon 13 of the Swel promoter, the Swe1p open reading frame (ORF) fused in frame to the septin-Cdc3p promoter (referred to here as “septin-Swe1p”) (Fig. 1A). We integrated this plasmid at an exogenous locus so that cells would arrest in the subsequent G1 phase. Samples for Western blotting were scored for budding and nuclear division on an Axioskop apparatus with a 100× objective equipped with epifluorescence and differential interference contrast (DIC) optics (Zeiss, Thornwood, NY). For DIC and GFP photography, cells were grown in complete synthetic medium containing dextrose and spotted onto a 2% agarose slab made with synthetic medium. Images were acquired with an Axioimager apparatus with a 100× objective equipped with epifluorescence and differential interference contrast optics (Zeiss) and photographed with an Orca ER monochrome cooled charge-coupled-device camera (Hamamatsu, Bridgewater, NJ). Images were captured using MetaMorph software (Millipore, Billerica, MA).

**Western blotting.**

Cell lysis was performed according to the tricholoroacetic acid (TCA) method (14). Samples were subjected to SDS-PAGE on a 6% polyacrylamide gel. Proteins were transferred onto a nitrocellulose membrane (Pall, East Hills, NY) and blocked for 1 h with PBS containing 3% nonfat dry milk (Kroger, Cincinnati, OH). Antibodies were diluted in blocking buffer supplemented with 0.1% Tween 20 (Bio-Rad, Hercules, CA), and membranes were probed overnight with a 1:250 dilution of mouse anti-myc antibody 9E10 and a 1:50,000 dilution of rabbit anti-Cdc11p antibody (Santa Cruz Biotechnology, Santa Cruz, CA). Membranes were then washed three times with PBS–0.1% Tween 20 and probed with a 1:7,500 dilution of goat anti-mouse HRP (Invitrogen, Carlsbad, CA) or a 1:7,500 dilution of goat anti-rabbit Alexa Fluor 680 antibody (Invitrogen, Carlsbad, CA) for 1 h in blocking buffer containing 0.1% Tween 20 and 0.01% SDS. Blots were then washed as described above and visualized by using an Odyssey scanner (Li-Cor Biosciences, Lincoln, NE). Fluorescence was quantified by using Odyssey software (Li-Cor Biosciences), where the intensities of the myc bands were normalized to the Cdc11 loading controls.

**RESULTS**

If the role of Hsl1p and Hsl7p is simply to recruit Swe1p to the septin cortex at the neck, then the artificial linkage of Swe1p to a septin should bypass the need for Hsl1p and Hsl7p in Swe1p degradation. To test this hypothesis, we created a fusion protein that joins GFP-Cdc3p (a septin) with Swe1p-myc, under the control of the Swe1p promoter (referred to here as “septin-Swe1p”) (Fig. 1A). We integrated this plasmid at an exogenous locus so that septin-Swe1p did not replace endogenous Swe1p or Cdc3p. As anticipated, septin-Swe1p was localized to the bud neck independent of Hsl1p and Hsl7p (Fig. 1B).
Localization and degradation of a septin-Swe1p fusion. (A) Schematic of the Cdc3-GFP-Swe1p fusion protein. (B) Localization of Cdc3-GFP-Swe1p to the neck is independent of Hsl1p and Hsl7p. GFP-Swe1p is shown as a control. Cells were grown overnight at 30°C. Representative cells are shown, and over 200 cells of each strain were scored for bud neck signals. Strains containing GFP-Swe1p (with genotypes indicated in parentheses) were DLY12320 (wild type [WT]), DLY12369 ($hsl7^{+}$/H9004), DLY12237 ($hsl1^{+}$/H9004), and DLY12229 ($hsl1^{+}$/H9004 $hsl7^{+}$/H9004), and strains containing Cdc3-GFP-Swe1p were DLY12321 (wild type), DLY12238 ($hsl1^{+}$/H9004), DLY12370 ($hsl7^{+}$/H9004), and DLY12230 ($hsl1^{+}$/H9004 $hsl7^{+}$/H9004). (C) Swe1p and septin-Swe1p levels rise and fall in parallel as cells traverse the cell cycle. Cells containing both Swe1p-12×myc and septin-Swe1p-12×myc (DLY12321) were arrested in G1 phase with pheromone and released into fresh medium at 30°C to traverse the cell cycle. Pheromone was reintroduced 60 min after release to rearrest cells at the next G1 phase. Samples taken at 15-min intervals were analyzed by Western blotting with anti-myc antibody (Cdc11 as a loading control) and quantitated (graph). (D) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel C were scored for >200 cells per sample. (E) Swe1p degradation kinetics are reproducible between experiments. Five replicates of the experiment depicted in panel C were quantified, and an average degradation profile for wild-type Swe1p is plotted as a dotted black line. (F) Degradation of septin-Swe1p occurs independent of wild-type Swe1p. A single-cycle synchrony experiment, as described above for panel C, was performed with swe1Δ strain DLY15563. (G) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel F were scored for >200 cells per sample.
To directly compare Swe1p and septin-Swe1p levels, both proteins were myc tagged in the same strain. Cells were synchronized in G1 phase by using mating pheromone, released to traverse the cell cycle, and rearrested in the subsequent G1 phase by the read- dition of pheromone. The kinetics of Swe1p accumulation and degradation were highly reproducible by using this protocol (Fig. 1E). The septin-Swe1p fusion accumulated, became hyperphosphorylated, and then disappeared with a timing comparable to that of the wild-type Swe1p-myc control (Fig. 1C), suggesting that the normal Swe1p degradation machinery can target the full septin-Swe1p fusion for degradation. Septin-Swe1p degradation was similar whether or not wild-type Swe1p was present (compare Fig. 1F with Fig. 1C), indicating that its degradation was not due to dimerization with wild-type Swe1p.

If Hsl1p and Hsl7p simply tether Swe1p to the neck for degradation, then septin-Swe1p should be degraded in strains lacking those proteins even though wild-type Swe1p is stable. However, a complication arises in strains containing both wild-type Swe1p and septin-Swe1p, because Swe1p degradation also requires active CDK (26). If wild-type Swe1p is stable, it would inhibit the CDK, and that inhibition could indirectly stabilize septin-Swe1p. To avoid this complication, we introduced an E12K mutation into the CDK Cdc28p. Cdc28pE12K is highly resistant to Swe1p-dependent inhibition, unlike the nonphosphorylatable Cdc28p Y19F, which is still susceptible to inhibition by binding to Swe1p (20). We used this CDC28E12K background for all Swe1p stability experiments, and as shown for the controls below, cell cycle progression in this background was unaffected by any of the manipulations of Swe1p, Hsl1p, or Hsl7p.

In hsl1Δ or hsl7Δ deletion strains, we found that both Swe1p and septin-Swe1p were stabilized in G2/M phase (Fig. 2). These findings do not support the idea that Hsl1p and Hsl7p act simply by tethering Swe1p to the neck: at minimum, they must play additional roles in promoting Swe1p degradation.

Hsl1p has an N-terminal kinase domain and a long C-terminal tail containing several regions with sequence motifs that are conserved among the Saccharomyces. In particular, region 8 mediates Hsl1p-Hsl7p interactions (Fig. 3A) (10). In a strain where HSL1 is replaced with hsl1/H9004Δ, neither Hsl7p nor Swe1p is recruited to the bud neck, and Swe1p is stable. However, we found that septin-Swe1p was degraded normally in this strain (Fig. 3B), showing that septin tethering bypasses the need for the Hsl1p-Hsl7p interaction in Swe1p degradation.

Like region 8, the kinase activity of Hsl1p is important for recruiting Hsl7p to the bud neck (28). In a strain where hsl1Δ is replaced with the kinase-dead hsl1K110R mutant, Swe1p is not detectably enriched at the bud neck, and it is stable. Although septin-Swe1p was effectively localized to the bud neck in an hsl1K110R strain, it was not degraded in G2/M phase (Fig. 4). Thus, the kinase activity of Hsl1p is needed not only for the effective recruitment of Swe1p to the bud neck but also for some subsequent step in Swe1p degradation.

A previous screen for SWE1 mutants that caused a lethal G2 arrest in the absence of MIH1 identified many point mutants in a

FIG 2 Hsl1p and Hsl7p are required for degradation of septin-tethered Swe1p in G2/M phase. (A) Swe1p and septin-Swe1p are both stabilized upon the deletion of HSL1. A single-cycle synchrony experiment, as described in the legend of Fig. 1C, was performed with hsl1Δ strain DLY12238. (B) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel A were scored for >200 cells per sample. (C) Swe1p and septin-Swe1p are both stabilized upon the deletion of HSL7. A single-cycle synchrony experiment, as described in the legend of Fig. 1C, was performed with hsl7Δ strain DLY12370. (D) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel C were scored for >200 cells per sample.
A small conserved region in the Swe1p N-terminal regulatory domain (21). The deletion of that region (in a protein designated Swe1p/H9004) did not impair Swe1p function but greatly reduced its interaction with Hsl7p and eliminated detectable neck localization. The fusion of Swe1p/H9004 to the septin Cdc3p restored neck localization, but septin-Swe1p was not degraded in G2/M phase (Fig. 5). The simplest interpretation of this result is that the Swe1p-Hsl7p interaction is required for Swe1p degradation, even when Swe1p is tethered to the neck.

**DISCUSSION**

In this study, we have tested the hypothesis that the role of Hsl1p and Hsl7p in Swe1p degradation consists of tethering Swe1p to the septins at the bud neck. Our data indicate that even when Swe1p is artificially tethered to the neck by fusion to a septin, the degradation of the tethered septin-Swe1p still requires both Hsl1p and Hsl7p. Thus, Hsl1p and Hsl7p play roles in promoting Swe1p degradation beyond simply tethering it to the neck.

Cells with a septin-tethered Swe1p still required the Swe1p-Hsl7p interaction for degradation. As phosphorylation by Cdc5p is thought to be the trigger for Swe1p ubiquitination, one possibility would be that the Hsl7p-Swe1p interaction presents Swe1p as a substrate. Hsl7p has also been reported to enhance Cdc5p neck localization, consistent with the idea that Hsl7p may interact with Cdc5p as well, potentially bridging a Cdc5p-Swe1p interaction.

**FIG 3** Interaction between Hsl1p and Hsl7p is not required for degradation of septin-tethered Swe1p. (A) Schematic of Hsl1p. a.a., amino acids. (B) Swe1p is stabilized upon deletion of Hsl1p region 8, but septin-Swe1p is not. A single-cycle synchrony experiment, as described in the legend of Fig. 1C, was performed with hsl1Δ strain DLY15224. (C) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel B were scored for >200 cells per sample.

**FIG 4** Hsl1p kinase activity is required for degradation of septin-tethered Swe1p. (A) Schematic of Hsl1p. The K110R point mutation in the kinase domain is noted by an asterisk. (B) Septin-Swe1p is localized to the bud neck in hsl1K110R cells, but GFP-Swe1p is not. Cells of strains DLY12973 and DLY15597 were grown at 30°C and photographed. (C) Swe1p and septin-Swe1p are both stabilized in a strain harboring the kinase-dead hsl1K110R mutant. A single-cycle synchrony experiment, as described in the legend of Fig. 1C, was performed with hsl1K110R strain DLY12973. (D) Synchrony parameters (budding and nuclear division) for the experiment depicted in panel C were scored for >200 cells per sample.
Given that the interaction with Hsl7p is required for septin-Swe1p degradation, Hsl7p must presumably be present at the neck to promote degradation. How, then, is septin-Swe1p degraded in the \textit{hsl1}\textsuperscript{abs} strain, where Hsl1p cannot recruit Hsl7p to the neck? It seems likely that in this case, septin-Swe1p itself recruits Hsl7p to the neck, reversing the normal tethering order.

Cells with a septin-tethered Swe1p also required Hsl1p kinase activity for degradation to occur. Hsl1p is homologous to the Nim1 kinase in \textit{Schizosaccharomyces pombe}, which can directly phosphorylate the Swe1p homolog Wee1 \textit{in vitro} (9, 22, 30). However, the direct phosphorylation of Swe1p by Hsl1p has been difficult to detect \textit{in vitro} (7), and it remains unclear whether or not it occurs \textit{in vivo}. Hsl1p is thought to phosphorylate Hsl7p \textit{in vivo} (19), and it also catalyzes extensive autophosphorylation (3), but the functional relevance of these modifications is unknown. As Hsl1p kinase activity is required for optimal Hsl7p recruitment, it could be that the phosphorylation of either Hsl1p or Hsl7p enhances the interaction between those proteins. However, the degradation of septin-tethered Swe1p did not require an intact Hsl1p-Hsl7p interaction, so promoting that interaction cannot be the only role for the Hsl1p kinase activity.

The finding that the need for the Hsl1p-Hsl7p interaction in Swe1p degradation can be bypassed by tethering Swe1p to the septins provides strong evidence that tethering is indeed one important role for Hsl1p and Hsl7p. Thus, our findings in aggregate indicate that Hsl1p and Hsl7p play at least two separate roles in promoting Swe1p degradation: tethering Swe1p to the septins and a downstream role perhaps involving the presentation of Swe1p to other regulators for targeting to the degradation pathway.

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