The SLT2 (MPKI) MAP Kinase Homolog Is Involved in Polarized Cell Growth in Saccharomyces cerevisiae

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Abstract. Bud emergence, spindle pole body duplication and DNA replication are all dependent on the activation of the CDC28 protein kinase at the Start point in the G1 phase of the cell cycle. Bud emergence requires polarization of the cytoskeleton and secretory vesicles to a specific site on the cell surface. Cdc28p activated by G1-cyclins triggers polarization of actin to the site of bud emergence and favors apical bud growth (Lew, D. J., and S. I. Reed. 1993. J. Cell Biol. 120:1305-1320). We isolated slt2-1 as a mutation that enhances the division defect of cdc28 mutants with defects at Start. Slt2p(Mpklp) is a member of the MAP kinase family (Lee, K. S., K. Irie, Y. Gotoh, Y. Watanabe, H. Araki, E. Nishida, K. Matsumoto, and D. E. Levin. 1993. Mol. Cell. Biol. 13:3067-3075). We show that slt2 mutants exhibit phenotypes similar to those shown by mutants of the yeast actin cytoskeleton, including delocalization of chitin deposition and of actin cortical spots and the accumulation of secretory pathway membranes and vesicles. Furthermore, slt2::HIS3 act1-1 and slt2::HIS3 myo2-66 double mutants are inviable. We suggest that Slt2p functions downstream or in parallel with Cdc28p in promoting bud formation and apical growth.

Cell reproduction involves cell cycle-specific events occurring both in the nucleus and in the cytoplasm. In the budding cell cycle of Saccharomyces cerevisiae, a number of stage-specific events direct the position and the morphology of the bud. These events include the choice of a bud site, the localization of a specific group of proteins to the site from which the bud will emerge, and the polarization of cell surface growth to the new bud site. Assembly of the bud-site complex is required for the polarization of the yeast cytoskeleton and the targeting of secretory vesicles to the bud site (for reviews see references 9, 12, 32).

The initial growth of the bud is highly polarized. Actin cortical patches (22), calmodulin (6), and the SPA2 protein (53, 54) are localized to the tips of the bud during this period of apical growth. Actin localization and bud growth are less polarized later in the cell cycle. The timing of this shift and the degree of delocalization determine bud shape, with isotropic growth favoring spherical forms (28, 55).

The entry into a new cell cycle involves the coordinated but independent pathways of bud initiation, spindle pole body duplication, and nuclear DNA synthesis (43). The triggering of these pathways all require the activation of the CDC28 protein kinase in the G1 phase of the cell cycle (45). This regulatory point of the cell cycle has been baptized “Start” (18). The manner in which the CDC28 kinase promotes bud formation at Start is not well defined. Lew and Reed (28) have shown that actin polarization can be triggered by the kinase activity of Cdc28p complexed with G1 cyclins, and they mapped the time at which actin polarization occurs in their synchronized cells to be soon after Start. Localization of Cdc3p (23) and Spa2p (54) to the presumptive bud site occurs substantially before bud emergence, but the precise timing of these events relative to CDC28 kinase activation at Start was not examined. However, cdc28 mutants at their restrictive temperature and wild-type cells treated with mating pheromone can polarize their growth to form a projection in the apparent absence of CDC28 kinase activity (28, 31, 54). These observations suggest that a pathway for polarizing growth independent of Cdc28p also exists in S. cerevisiae.

We isolated a mutation of the SLT2 (MPKI) MAP kinase gene that augments the division defect of cells expressing a partially inactivated cdc28 mutant. We show that slt2 mutants exhibit defects in cell polarity and accumulate secretory vesicles. We suggest that Slt2p acts downstream or in parallel with Cdc28p to promote bud morphogenesis.

Materials and Methods

Yeast Strains and General Methods

The yeast strains used in this study are listed in Table I. The cdc28-109 allele was found amongst a collection of cdc28 mutants in the Department of Genetics (University of Washington, Seattle, WA). Its characterization will...
be described elsewhere. The CML6 and CML13 SLT2+ strains were obtained by transforming respectively CYM678 MATa sit2-1 and CML12 MATa sit2-1 strains with a YIp-URA3-SLT2 plasmid and targeting plasmid integration to the sit2-2 site. SLT2+ ura3-52 segregants were then selected by growth on a 5-fluoro-orotic acid (FOA)-containing medium. Media containing FOA for the counterselection of URA3+ plasmids were prepared as described (3). Standard genetic and molecular manipulations were performed as described (49, 50). The segregation in genetic crosses of cdc28-109, sit2-1, actl, and myo2-66 was followed by complementation testing of the temperature-sensitive mutations.

**Determination of Cell Volumes**

Cell volumes were determined by photographing cells at 250× magnification with a Zeiss Axioplan light microscope equipped with differential interference contrast optics. The negatives were projected onto a flat surface and the major and minor axes of unbudded cells and the mother cell portions of budded cells were measured. Cell volumes were estimated by assuming the cells to be prolate ellipsoids and using the formula \( V = 4/3 \pi ab^2/3 \), where \( a \) is the semi-major axis and \( b \) is the semi-minor axis. At least 250 cells were measured for each population examined.

**Time Lapse Photomicroscopy**

5 \( \mu \)l of a cell suspension containing approximately 6 \( \times \) 10^6 cells per ml were placed on a thin layer of YPD agar on a microscope slide. A cover slip was placed over the cells and cell proliferation was followed by differential interference contrast microscopy at a magnification of 250×.

**Fluorescence Microscopy**

Cells were fixed in 3.7% formaldehyde and stained with Calcofluor to detect chitin or with rhodamine-conjugated phalloidin to detect filamentous actin. Cells were fixed in 3.7% formaldehyde and stained with Calcofluor to detect chitin or with rhodamine-conjugated phalloidin to detect filamentous actin.

**Electron Microscopy**

Yeast cells were fixed by addition of glutaraldehyde directly to the YPD growth medium at a final concentration of 1% (vol/vol) for ~30 min. Cells were then centrifuged, resuspended in a buffer containing 2% glutaraldehyde, 0.1 M sodium cacodylate (pH 7.4), and 0.8 M sorbitol, and then incubated overnight. Cells were then postfixed for 1 h in reduced osmium (a 1:1 mixture of 2% aqueous osmium tetroxide and 3% aqueous potassium ferrocyanide), dehydrated in ethanol, and embedded in Epon. 80-um sections were cut with a Reichert ultramicrotome, stained for 2 min with lead citrate and examined at 80 kV with a Philips CM12 electron microscope.

**Results**

**Isolation of sit2-1**

As a cdc28-colethal Mutation

In the hope of identifying functions that are required either downstream or in parallel with the CDC28 protein kinase for cell-cycle progression, we are isolating mutants that are colethal (synthetically lethal) with cdc28. A cdc28-colethal mutant is one which in combination with a temperature-sensitive cdc28 mutant prevents growth at a normally permissive growth temperature.

**Table 1. Yeast Strains Used in This Study**

| Name | Genotype | Source |
|------|----------|--------|
| CYM678 | Mata, sit2-1, ura3-52, leu2, trpl1, ade2, lys2-801 | This study |
| CML6 | CYM678 SLT2+ isogenic strain | This study |
| CML12 | Mata, sit2-1, ura3-52, trpl1, ade2, lys2-801, leu2 | This study |
| CML13 | CML12 SLT2+ isogenic strain | This study |
| CML14 | Mata, sit2-1, ura3-52, trpl1, ade2, lys2-801, his3Δ200 | This study |
| CML15 | Mata/a, sit2-1/sit2-1, ura3-52/ura3-52, trpl1/trpl1, ade2/ade2, lys2-801/lys2-801, leu2/LEU2 his3Δ200/HIS3 | This study |
| SC55 | Mata/a, ura3-52/ura3-52, trpl1/trpl1, his3Δ200/his3Δ200, ade2-101/ade2-101, lys2-801/lys2-801, leu2/LEU2 can/CAN | This study |
| CML20 | Mata, sit2Δ::HIS3, ura3-52, trpl1Δ, ade2-101, lys2-801 | This study |
| CML22 | Mata, ura3-52, trpl1Δ, his3Δ200, ade2-101, lys2-801 | This study |
| CML712 | Mata, cdc28-4, ura3-52, trpl1Δ, leu2Δ1, his3Δ200, ade2-101, tyr1 | This study |
| CML27 | Mata/a, sit2Δ::HIS3, ura3-52, trpl1Δ, ade2-101, leu2Δ1 | This study |
| CML29 | Mata, sit2Δ::HIS3, ura3-52, trpl1Δ, ade2-101, leu2Δ1, lys2-801 | This study |
| CML776 | Mata, act1-1, ura3-52 | Novick and Botstein, 1985 and this study |
| CYM830 | Mata, myo2-66, ura3-52, leu2Δ1, ade2-101, his3Δ200, his6 | Johnston et al., 1991 and this study |
| YC190 | Mata, ura3-52, lys2-801, ade2-101, his3Δ200, trpl1Δ, leu2Δ90, slt1Δ::TRP1 | Costigan et al., 1991 |
| CML715 | Mata, cdc28-1N, ura3-52, leu2Δ1, lys2-801 | This study |
missive temperature for either single mutant. A conceptually similar scheme was used to search for possible downstream effectors of the Drosophila sevenless protein kinase (52). The results of this screen will be described in detail elsewhere. Briefly, a cdc28-109 temperature-sensitive mutant was transformed with an autonomously replicating plasmid containing the URA3 and CDC28 genes (YPEp24-CDC28). These cells were then mutagenized with ethylmethanesulfonylate and spread on YPD plates to have individual colonies at the permissive temperature (24°C) for the cdc28-109 mutant. The colonies were then replicated at the semi-permissive temperature of 28°C to plates containing media either selecting for (synthetic medium without uracil [-Ura]) or counterselecting (synthetic medium containing FOA) the YEp24-CDC28 plasmid. Colonies able to grow on the -Ura plates but not the FOA plates at 28°C were considered to harbor mutations coterhools with cdc28-109. We next screened amongst the coelothal mutations for those that caused temperature-sensitive growth at 37°C even in the presence of a wild-type CDC28 gene. One such mutation, slt2-1 (see below), was seen to give rise to a population of unusually small cells at the restrictive temperature and is the subject of this paper. The cdc28-109 slt2-1 YEp24-CDC28 mutant strain was backcrossed to a wild-type strain and tetrads were dissected after sporulation of the resulting diploid strain. The traits of coelothality with cdc28-109 at 28°C and temperature sensitivity for growth at 37°C were linked and segregated as a single mutation in 12 tetrads.

The cdc28-109 mutation leads to an arrest of cell division in both the G1 and G2 phases at the restrictive temperature of 37°C (unpublished data). To see if the incompatibility between the slt2 and cdc28 mutations could be assigned to a G1 or G2 specific function of cdc28, we crossed the slt2-1 mutant with cdc28-4 and cdc28-IN mutants. The cdc28-4 mutation preferentially inactivates the G1 function of Cdc28p (46), whereas the cdc28-IN allele preferentially inactivates a G2/mitotic function of the kinase (40, 57). slt2-1 cdc28-IN double mutants were readily obtained and grew as well as the individual single mutants at the permissive temperature for these mutants. In contrast, the slt2-1 cdc28-4 double mutants, although viable at 24°C, grew more slowly and exhibited phenotypes at 24°C that were only seen at higher temperatures in the single mutants. Furthermore, cdc28-4 and slt2-1 mutants can each grow at the semi-permissive temperature of 30°C, whereas the double mutant was inviable at this temperature. These genetic interactions suggest that CDC28 and SLT2 cooperate in performing a G1-specific function.

A DNA fragment complementing the slt2-1 temperature-sensitive growth defect was obtained from a YCP50 yeast genomic DNA library (47). DNA sequencing of the complementing region revealed the presence of a gene identical to the previously described SLT2 (suppressor of lyt2) gene (60), except for three nucleotide differences (see Materials and Methods). The SLT2 gene encodes a protein that is similar to the CDC28, FUS3, and KSS1 protein kinases (26, 60). Cdc28p is a member of the CDC2 family of protein kinases whereas Fus3p and Ksslp are members of the MAP/ERK family of kinases. A sequence comparison of Slt2p with all known CDC2-like and MAP kinases shows that Slt2p clearly belongs to the MAP kinase family (Fig. 1). Slt2p has a carboxyl-terminal extension relative to most of the CDC2/ CDC28 and MAP kinases. If only the shared catalytic domains are considered, Slt2p is about 28% identical to the CDC2/CDC28 kinases compared to about 42% identity with the MAP kinases. The sequence PSTAIRE is present in most of the CDC2-like kinases and has been implicated in the binding of mitotic cyclins (13). The PSTA portion of PSTAIRE is not conserved in SLT2 (aa 66-72) and the MAP kinases. The sequence GTYG is present near the aminoterminus of all the CDC2/CDC28 kinases. An inhibitory phosphorylation of this tyrosine residue, and sometimes the adjacent threonine residue, regulates the timing of activation of CDC2 and the entry into mitosis (37). In SLT2 (amino acids 32-35) and the MAP kinases, this sequence is replaced by GAYG and there is no evidence so far for the phosphorylation of the tyrosine residue in this sequence. Furthermore, we have identified an additional 8-amino acid residues (Fig. 1) that are conserved in all MAP kinase homologs examined to date, including SLT2, but are not found in any of the CDC2-like kinases. Amongst these is a tyrosine residue (aa 192 of SLT2) whose phosphorylation by distinct MAP kinase kinases is required for the activation of the MAP kinases (24). This tyrosine residue is contained within the sequence TETY found in all MAP kinases except the recently identified HOG1 kinase (5), where it is TGY. The SLT2 gene has also been re-isolated by Lee et al. as a multicopy suppressor of a bck1(slkl) mutant (26). They have shown that a Xenopus MAP kinase can complement an slt2 mutation, that mutation of threonine-190 to alanine or tyrosine-192 to phenylalanine inactivates SLT2, and they have renamed the gene MPKI.

A multicopy plasmid containing an SLT2 gene deleted for the sequence encoding the last 158 amino acids of the protein was obtained while isolating multicopy suppressors of a cdc28-IN mutation (Marc Blondel, personal communication). This deletion removes the carboxy-terminal amino acid sequence specific to slt2p as well as two residues (glu327 and pro328) that are found in all MAP kinases examined to date (see Fig. 1). Further examination showed that this plasmid could partially suppress the temperature-sensitive phenotype of the cdc28-IN mutation, but was incapable of complementing the slt2-1 mutation. The cdc28-IN allele gives a G2/mitosis-specific arrest at 37°C (40, 57). We found that both the truncated SLT2 gene and the intact SLT2 gene on multicopy plasmids were capable of partially suppressing the cdc28-IN temperature-sensitive phenotype. We interpret these results as indicating that the slt2p truncation deletes residues that are required for the function of slt2p, but not necessarily for the basic kinase activity of the protein. In this model, the partial suppression of the cdc28-IN mutation might be explained by the ability of the overexpressed SLT2 kinase, in its intact or truncated form, to phosphorylate certain critical substrates of CDC28 in G2/mitosis. This possibility is rendered plausible by the similarity in structure and substrate specificity of the CDC2 and MAP kinase families (39). Wild-type cells are transiently inhibited in their growth after transformation with multicopy plasmids containing the intact or truncated SLT2 gene, but then seem to adapt to this overexpression and grow normally.

**Phenotypic Characterization of the slt2-1 Mutant**

Disruption of the SLT2 (MPKI) gene caused a temperature-sensitive cell lysis phenotype (26, 60). We observed a similar
phenotype for the EMS-induced sit2-1 mutation. Transfer of sit2-1 cells from 24°C to 37°C leads to a rapid loss of cell viability (Fig. 2 A) associated with a large degree of cell lysis as seen by phase contrast microscopy. The cell division arrest was not homogeneous, although the percentage of unbudded cells increased at 37°C from 49 to 71% before death as seen by phase contrast microscopy. The cell division arrest phenotype for the EMS-induced sit2-1 cells in the culture was observed at 37°C as monitored by cytofluorimetry (data not shown). Continuous incubation at 30°C, there is a significant level of cell death of the entire population (Fig. 2 A). An isogenic control strain (Fig. 2, B and C) that was accompanied at this temperature by the death of most cells in the culture after 4 h of incubation. A proportional increase in the fraction of cells with a G1 DNA content was observed at 37°C as monitored by cytofluorimetry (data not shown). Cell lysis was observed for both budded and unbudded cells in the culture.

We additionally noticed a novel sit2 phenotype: mutant cells exhibit a strikingly heterogeneous range of sizes including unusually small cells with a characteristic pear-shaped morphology and abnormally large, round cells. This size heterogeneity was characterized by measuring cell volumes from photomicrographs. At the permissive temperature of 24°C, exponentially growing sit2-1 cells were highly viable with little or no signs of cell lysis, but a subpopulation of small cells was manifest that was never seen in the wild-type control strain (Fig. 2, B and C). Transferring the cells to the semi-permissive temperature of 30°C increased dramatically the cell size heterogeneity and increased the percentage of unbudded cells in the population from 49 to 66%. After 4 h at 30°C, about 20% of the cells were unusually small. At the same time, a subpopulation of unusually large, round cells was observed. Although the sit2-1 cells can be grown continuously at 30°C, there is a significant level of cell death observed at this temperature (Fig. 2 A). Transferring cells from 24°C to 37°C also induced a cell size heterogeneity (Fig. 2, B and C) that was accompanied at this temperature by the death of the entire population (Fig. 2 A). An isogenic SLT2 strain showed no change in their size distribution when transferred from 24°C to 30°C. Transferring the SLT2 strain from 24°C to 37°C for 4 h caused an overall shift to larger cell sizes compared to 24°C (Fig. 2 C) that may be attributed to a transitory G1 arrest associated with the 37°C heat shock (21). Note, however, that the cell size distribution remains clustered and no small cells are generated.

We never observed a bud associated with the subpopulation of small, pear-shaped sit2 cells. Time-lapse photomicrography (Fig. 3) demonstrated that these cells were in fact stillborn. The small cells are stillborn even at the permissive temperature of 24°C where the vast majority of sit2-1 cells are viable (Fig. 2 A). We do not know why these cells are dead. Staining of DNA with the fluorochrome DAPI showed that the small, pear-shaped cells had inherited nuclei.

Another phenotype presented by the sit2 mutants is an increased mortality in stationary phase. An example of this sensitivity is shown in Fig. 4. Congenic sit2-1 and SLT2+...
Figure 2. Cell viability and cell size heterogeneity in the slt2-1 mutant. (A) Cell viability of CMY678 slt2-1 cells growing in exponential phase in YPD at 24°C (○) and after transfer to 30°C (●) or 37°C (△). Cell viability was determined by spotting cells on YPD plates and observing microscopically the percentage of cells capable of forming a microcolony after 12–24 h of growth at 24°C. (B) Histogram of the individual cell volumes of CMY678 slt2-1 cells grown in exponential phase in YPD at 24°C (□) or after transfer to 30°C for 4 h (□) or to 37°C for 4 h (●). (C) Histogram of the individual cell volumes of the CML6 SLT2 isogenic control strain grown in YPD at 24°C (□) and after transfer to 37°C for 4 h (●). Cells were partitioned into volume classes differing by 12 μm as per Johnston et al. (20).

The slt2::HIS3 Disruption Has a Phenotype Similar to That of the slt2-1 Mutant

A disruption of the SLT2 (MPKI) gene was reported to cause a temperature-sensitive cell lysis defect (25, 60). We repeated this analysis in order to determine whether the same result is true in our strain background and to compare the phenotypes of our EMS-induced slt2-1 mutant with that of an SLT2 gene deletion. A 1.3-kb HindIII fragment internal to the SLT2 gene was replaced with the HIS3 gene. This substitution deletes sequences encoding all but the first 74 amino acids of Slt2p. The slt2::HIS3 fragment was used to disrupt one of the SLT2 genes in a diploid yeast strain. After sporulation and dissection of tetrads at 24°C, two slow growing slt2::HIS3 segregants and two faster-growing SLT2 segregants were generally observed. The slt2::HIS3 segregants were temperature sensitive for growth at 37°C with the expected cell-lysis phenotype. We observed that the slt2::HIS3 strains at 24°C grew more slowly than the slt2-1 mutant, with slt2::HIS3 cells showing at 24°C a cell-size heterogeneity and a degree of cell lysis resembling that seen for slt2-1 at 30°C.

The Heterogeneous Size of the slt2 Mutants Recalls the Phenotype of Actin Cytoskeleton and Polarized Growth Mutants

The presence of both abnormally small and abnormally large cells in the slt2 cultures could be due to a defect in polarized cell growth. During budding, cell surface growth is directed primarily to the bud via secretory vesicles that are targeted to this portion of the plasma membrane. If this process of directed secretion is partially inhibited without otherwise affecting cell growth and division, then we would expect buds to be smaller than usual and mother cells to grow abnormally large (assuming that untargeted secretory vesicles can fuse with the plasma membrane of the mother cell). Just such a phenotype has been observed for several mutants of the yeast cytoskeleton, including act1 (34), myo2 (20), and rpm1 (29), defective in actin, an unconventional myosin, and tropomyosin, respectively. Because of these similarities, we examined our slt2 mutants for other phenotypes associated with defects in polarized growth.

Chitin Deposition Is Delocalized in the slt2-1 Mutant

Chitin is deposited in the cell wall to form a ring at the base of the bud in wild-type cells (7). In haploid cells, the site of bud emergence is adjacent to the previous bud, whereas in diploid cells the pattern of budding is bipolar (8). Chitin can
be stained specifically with the dye Calcofluor (42, 44). Localization of chitin in the haploid slt2-1 mutant at the permissive temperature of 24°C (Fig. 5 A) was quite similar to that of the wild type (Fig. 5 E), with fluorescence restricted to one extremity of the cell in the form of clearly discernible bud scars. After incubation at 37°C for 3 h, wild-type cells showed a slight and transient delocalization of chitin synthesis (Fig. 5 F) that we attribute to a transitory G1 arrest induced by the heat shock (Fig. 2 C and reference 21). The mutant cells at 37°C showed a much greater chitin delocalization (Fig. 5 B). Large, round cells showed the brightest staining with the greatest degree of delocalization. A similar result was seen with the act1 (34), myo2 (20), and tpm1 (29) mutants. We also observed that slt2/slt2 homozygous diploid mutants have more severe phenotypes than the corresponding slt2 haploid cells. For example, a heterogeneous cell size

Figure 3. The abnormally small, pear-shaped slt2-1 cells are stillborn. CMY678 slt2-1 cells growing at 24°C in YPD, or after a 2-h incubation at 30°C to increase the frequency of the small, pear-shaped cells, were spotted onto a thin layer of YPD agar on a microscope slide and cell proliferation was monitored at room temperature over a 24-h period by time-lapse photomicroscopy. Arrows indicate the positions of the small stillborn cells. Cells of this highly characteristic size and morphology never gave rise to a bud. The arrowhead in the 24°C 24-h panel points to a mother cell that has grown anomalously large. Bar, 10 μm.

Figure 4. slt2-1 mutants lose viability rapidly in stationary phase. CML12 slt2-1 (●) and isogenic CML13 SLT2+ cells (○) were grown at 24°C in YPD medium over a period of 8 d. Cell viability (A) and the percentage of budded cells (B) were monitored as a function of time.
Figure 5. *slt2*/*slt2* mutants show delocalized deposition of cell surface chitin. *slt2*/*slt2* cells growing in YPD were fixed in 3.7% formaldehyde and stained for chitin with Calcofluor. (A) CMY678 *slt2*/*slt2* haploid cells grown at 24°C and (B) after transfer to 37°C for 3 h. (C) CML15 *slt2*/*slt2* homozygous diploid cells grown at 24°C and (D) after transfer to 37°C for 3 h. (E) CML6 SLT2+ isogenic strain at 24°C and (F) after transfer to 37°C for 3 h. All cells are shown at the same magnification. Bar, 10 μm.

and delocalized chitin staining was observed for *slt2*/*slt2* diploid cells at the permissive temperature of 24°C (Fig. 5 C). The intensity of chitin staining of the *slt2*/*slt2* diploid cells increased after incubation at 37°C for 3 h (Fig. 5 D). Interestingly, diploid cells deleted for the *TPM1* gene encoding tropomyosin (29) and *myo2-66* homozygous diploids (20) were also reported to have more severe phenotypes than the corresponding haploid cells.

**Actin Cortical Patches Are Delocalized in *slt2* Mutants**

The lysis observed in the *slt2*::*HIS3* strains at 24°C makes their characterization difficult. The addition of osmotic
stabilizers has been reported to suppress the temperaturesensitive lysis of slt2 (mpkl) disruption mutants (26, 60). Hence, to overcome the technical difficulties due to cell lysis, we tested the growth of slt2::HIS3 and SLT2+ cells in YPD media containing sorbitol at concentrations of 0.5-1 M. We found 0.8 M sorbitol to give the best suppression of the slt2::HIS3 mutant phenotypes with the least inhibition of growth of our isogenic SLT2+ cells. However, although 0.8 M sorbitol suppressed the temperature sensitivity of the slt2-I and slt2::HIS3 strains, this suppression was incomplete. Some cell lysis and unusually small cells were still observed at 37°C. We looked at the distribution of filamentous actin in slt2::HIS3 and SLT2+ strains at 37°C in YPD + 0.8 M sorbitol using rhodamine-conjugated phallolidin. In S. cerevisiae, actin localization reflects the sites of cell surface growth (22).

slt2::HIS3 and wild-type cells growing in YPD medium containing 0.8 M sorbitol at 24°C had a normal actin distribution (data not shown). After transfer to 37°C for 2 h, the slt2::HIS3 cells growing in YPD medium containing 0.8 M sorbitol showed a clear delocalization in their distribution of cortical actin spots (Fig. 6 B) when compared to the SLT2+ control strain (Fig. 6 A). Numerous actin spots were observed in the mother cell portion of mutant cells with small buds, a situation which is virtually never seen in wild-type cells. There remained a significant residual polarization of the actin spots in buds in these cells that is consistent with the partial suppression of the mutant phenotypes by the sorbitol. Thus, the SLT2 kinase is required at high temperatures for the normal distribution of the actin cytoskeletal network.

Secretory Vesicles Accumulate in the slt2::HIS3 Mutant

Cytoskeletal mutants with defects in polarized growth, such as actl (34), myo2 (20), and tpml (29), have been shown to accumulate secretory vesicles. We examined slt2::HIS3 cells growing at 24°C or at 37°C in YPD + 0.8 M sorbitol by electron microscopy for a similar phenotype. Sections of glutaraldehyde-fixed cells were postfixed in a 1:1 mixture of 2% osmium tetroxide and 3% potassium ferrocyanide before staining with lead citrate. This treatment results in a strong staining of reducing sugars that allows an excellent visualization of elements of the secretory pathway and the yeast cell wall (Fig. 7). The slt2::HIS3 cells growing at 24°C in YPD + 0.8 M sorbitol (Fig. 7 A) were indistinguishable from wild-type cells. These cells contained a small number of tubules near the inner surface of the plasma membrane or extending from the nuclear membrane and some strongly stained cis-ternae in the cytoplasm (data not shown). The tubules resemble elements of the endoplasmic reticulum and the cis-ternae resemble discrete Golgi compartments that have been previously identified by immunoelectron microscopy (41). A small number of secretory vesicles was also observed. In budded cells, the secretory vesicles were found preferentially near the tip of the bud, as shown in Fig. 7 A and as seen in wild-type cells (Fig. 7, WT). In contrast, two types of aberrant internal morphology were observed in slt2::HIS3 cells transferred to 37°C for 1.5 h. Both budded and unbudded cells with dramatic accumulations of secretory vesicles distributed throughout the cytoplasm were observed (Fig. 7 B).
Fragmented vacuoles were seen in many of these cells. In some sections, a proliferation of tubular networks and unusual doughnut-shaped structures was also observed (Fig. 7C). Similar doughnut-shaped forms have been seen in sec mutants blocked late in the secretory pathway (35). Isogenic wild-type cells growing in YPD + 0.8 M sorbitol at 37°C did not show any unusual accumulation of secretory vesicles or internal membranes (Fig. 7, W7). We saw no signs of obvious cell wall defects in the sit2 mutant. A similar analysis was carried out for the sit2-1 mutant transferred from 24°C to 37°C for 1.5 h in YPD without sorbitol. An accumulation of secretory vesicles within the sit2-1 mutant was also observed, although to a lesser degree than that seen in the sit2::HIS3 mutant at 37°C.

We have examined thin sections of the myo2-66 and act1-l mutants at the restrictive temperature of 37°C with the same electron microscope strain used to study the sit2 mutant. We found that the myo2 (Fig. 7, D and E) and act1 (Fig. 7, F and G) mutants greatly resemble the sit2 mutant in their accumulation of secretory vesicles and membranous structures. This result is in complete agreement with the previous descriptions of myo2 (20) and act1 (36) and suggests that the sit2 mutant is similarly defective in targeting secretory vesicles to the bud.

**Genetic Interactions Implicating SLT2 in Cytoskeletal/Polarized Growth Functions**

Characterization of sit2 mutants reveals defects in polarized growth similar to those seen for the cytoskeletal mutants act1 and myo2. To test for genetic interactions between these mutations, we crossed the sit2-1 mutant with act1-l and myo2-66 mutants and sporulated the resulting heterozygous diploids. Tetrad analysis showed that the sit2-1, act1-l, and myo2-66 single mutant spores germinated and grew well at the permissive temperature of 24°C, whereas the sit2-1 act1-l and sit2-1 myo2-66 double-mutant spores were generally inviable. Approximately 90% of double mutant spores were inviable in a sample of at least 20 tetrads from each cross (the genotype of the dead segregants was deduced from that of the remaining viable spores in each tetrad). The sit2, act1, and myo2 mutants are not in isogenic strain backgrounds. We suspect that the existence of a small number of viable double mutants in these crosses is due to the segregation of polymorphic alleles in the different strain backgrounds that suppress partially the mutant phenotypes. In support of this idea, we found that one-half of the sit2 segregants in these crosses showed greater residual growth at 37°C than did the sit2 parental strain. Strain background effects on the severity of cytoskeletal mutant phenotypes have been frequently reported (1, 2, 29).

A more definitive demonstration of colethality between sit2 and act1 mutants was obtained by crossing CML29 (sit2::HIS3/pFL44-SLT2) with CMY776(act1-l). The sit2::HIS3 mutation of strain CML29 is complemented by the SLT2 gene on the multicopy pFL44(URA3) vector (4). Transmission through meiosis of pFL44-SLT2 occurs with high efficiency. After sporulation and asci dissection of a CML29 by CMY776 diploid, we analyzed the segregation of the sit2::HIS3 and the act1-l mutations in 10 tetrads in which the pFL44-SLT2 plasmid was transmitted to all spores. We next examined whether meiotic segregants were able to lose the pFL44-SLT2 plasmid after non-selective growth on YPD plates at the permissive temperature of 24°C by replica plating to an FOA-containing agar medium at 24°C. FOA (3) inhibits the growth of those cells that contain the pFL44(URA3) plasmid. All sit2::HIS3 act1-l/pFL44-SLT2 segregants were incapable of growing on the FOA-containing medium at 24°C, whereas all other segregants grew on this medium. A representative tetratype ascus is shown in Fig. 8. Thus, sit2::HIS3 act1-l cells are inviable in the absence of pFL44-SLT2. An sit2::HIS3/pFL44-SLT2 strain (CML27) was also crossed to a myo2-66 mutant (CMY830). All sit2::HIS3 myo2-66/pFL44-SLT2 segregants were unable to grow on FOA plates at 24°C, thereby showing that these two mutations are also incompatible. We conclude that the inactivation of SLT2 can be lethal in backgrounds in which the actin/Myo2p cytoskeleton is perturbed, thus suggesting a role for the SLT2 kinase in the optimal functioning of the yeast cytoskeleton.

Deletion of the SLK1(BCK1) kinase gene leads to a constellation of mutant phenotypes that greatly resembles those associated with the sit2 mutants, including a temperature-sensitive cell lysis defect, starvation sensitivity, and heterogeneous cell sizes with a population of unusually small cells (11, 25). We found that the temperature-sensitive growth and lysis phenotype of an slkl-D2 mutant (11) could be partially suppressed by transforming the strain with a multicopy plasmid containing the SLT2 kinase. Lee et al. (26) and Irie et al. (19) also observed this suppression and have carried out an extensive series of epistasis experiments suggesting that the Stl2p(Mpklp) kinase is a downstream activation target of the Slk1p(Bcklp) kinase (see Discussion section).

The SSDI-v allele is capable of suppressing a wide range of mutations that were isolated in ssdl-d backgrounds (56, 58, 61). Since we knew our lab strains to contain an ssdl-d allele (56), and because Costigan et al. had previously shown that slk1 is suppressed by SSDI (11), we tested for suppression of the sit2-1 mutation by SSDI-v. A YCP50-SSDI-v plasmid (58) was capable of partially suppressing the temperature-sensitive growth and lysis of our sit2-1 ssdl-d strain. Thus, SSDI represents at least one polymorphic allele in dif-
ferent laboratory strain backgrounds that can modify the penetrance of the slt2 phenotypes. Sslp (also known as Srklp) has been suggested to play a role in protein phosphatase function, but no biochemical studies of this protein have yet been reported (58, 61).

**Discussion**

**slt2 Mutants Have Phenotypes Reminiscent of Actin Cytoskeletal Mutants and Are Defective in Polarized Growth**

slt2 mutants exhibit a number of phenotypes that recall those of yeast cytoskeletal mutants, including cell-size heterogeneity and morphological defects, delocalization of both chitin deposition and actin cortical spots, and the accumulation of secretory vesicles. Similar phenotypes have been reported for mutants of actin (act1) (34), an unconventional myosin (myo2) (20), tropomyosin (tpm1) (29), and sac1, sac2, and sac3 (cold-sensitive suppressors of an actin mutant) (36). These phenotypes may all be manifestations of defects in polarized growth in these mutants due to perturbations of the actin cytoskeleton. Hence, the actin cytoskeleton seems to be required for directing secretory growth to the chosen site of bud emergence in unbudded cells, and then, for restricting growth to the bud after bud emergence. If this process is disturbed, then secretory vesicles (and possibly Golgi compartments as well) will not be efficiently targeted to the bud, and the buds will be smaller than usual. If the non-targeted secretory vesicles can fuse to the plasma membrane of mother cells, then these cells should grow abnormally large at the expense of the buds. The fact that some cytoskeletal mutant cells do get larger and show a delocalized deposition of chitin in their cell walls indicates that secretory vesicles can fuse to the mother cell plasma membrane. However, the observation that secretory vesicles and aberrant cisternal membranes accumulate within the act1, myo2, and slt2 mutants suggests that the nontargeted secretory vesicles do not fuse efficiently to the cell membrane.

Cell size heterogeneity accompanied by abnormal actin distributions and delocalized chitin deposition has also been observed in profilin (pfy) (17), fimbrin (sac6) (1), actin capping protein (cap1 and cap2) (2), rho3 and rho4 (33), and bem1 (10) mutants. These mutants have not yet been examined by electron microscopy for accumulation of secretory vesicles. Profilin, fimbrin, and actin capping protein all interact directly with actin, but the relationship of Rhp3p, Rhp4p, and Bemlp with the actin cytoskeleton has not yet been determined. The similarity of the slt2 and cytoskeleton mutant phenotypes suggests that the SLT2 kinase phosphorylates one or more of the cytoskeletal components and/or the bud-site complex proteins in such a way that polarized growth is stimulated. Less direct mechanisms are also possible. The SLK1(sck1) kinase is a putative upstream activator of the SLT2(MPK1) kinase (see below). Costigan et al. have shown that slt1 spa2 double mutants are inviable, whereas either single mutant is viable at 24°C (11). Since Spa2p is specifically localized to sites of polarized growth (53), this synthetic lethal interaction supports the idea that SLK1/SLT2 pathway functions in regulating polarized growth.

The starvation sensitivity of the slt2 mutants (Fig. 4) is one phenotype that has not been reported for the cytoskeletal mutants. It is not clear whether the necessity of SLT2 for normal stationary phase viability is related to its postulated role in the actin cytoskeleton and cell polarity, or whether it represents a completely independent function of the kinase. The slt1 mutant also loses viability rapidly upon nutrient starvation (11).

Deletion of the SLT2 gene is not lethal. The slt2::HIS3 cells grow slowly at 24°C, have defects in cell polarity, and are prone to lysis. At 37°C, cells die rapidly with lots of lysis. Many cytoskeletal components have similarly been shown to be necessary for normal cell polarity and cytoskeletal structure at 22°-30°C without being necessary for cell viability at these temperatures. Most of these proteins, including fimbrin (1), profilin (17), tropomyosin (29), Sac3p (36), Sac7p (14), and Bemlp (10) are required for viability at low (14°-16°C) and/or high (36°-38°C) temperatures. The relative dispensability of these proteins may mean that there are functionally redundant equivalents still to be found within the cell. Alternatively, it may mean that the actin cytoskeleton can function, albeit poorly, in the absence of these proteins at 22°-30°C, but not under conditions of stress, such as growth at low or high temperatures.

Torres et al. (60) have suggested that SLT2 is involved in cell wall biosynthesis. This suggestion was based largely on the fact that osmotic stabilizers can suppress (partially in our strain background) the temperature-sensitive cell lysis phenotype of slt2 mutants. However, mutants of the cortical cytoskeleton in yeast (1, 34) and in animal cells (30) can be fragile and prone to lysis. As discussed by Costigan et al. (11), mutants with defective cytoskeletons can sometimes be suppressed by osmotic stabilizers. It is also possible that osmotic stabilizers suppress slt2 phenotypes by a completely distinct mechanism. Slt2p is clearly a member of the MAP kinase family (see Fig. 1 and reference 26). Recently, another MAP kinase isoform in S. cerevisiae, Hoglp, was shown to be necessary for cell growth in media of high osmolality (5). Hoglp is tyrosine phosphorylated, and presumably activated, upon addition of 0.7 M sorbitol to the growth medium. Given the extensive sequence similarity between Slt2p and Hoglp (see Fig. 1), it is conceivable that the partial suppression of slt2 phenotypes could occur by Hoglp activation allowing it to phosphorylate some of the substrates that are normally phosphorylated by Slt2p. In summary, suppression of slt2 cell lysis by osmotic agents may be explained in several different ways. We suggest that any cell wall defects that exist in slt2 mutants could be secondary consequences of defects in polarized growth.

Two other MAP kinase isoforms of S. cerevisiae, FUS3 and KSS1, are activated by the mating pheromone pathway (16). We have observed that slt2 mutants are slightly defective in projection formation upon addition of alpha factor or in mating mixtures of yeast cells (data not shown). slt2::HIS3 mutants also exhibit a projection formation defect (11). The formation of a mating projection is another manifestation of polarized cell growth in the yeast life cycle. The fact that slt2 mutants are only slightly defective in projection formation may be due to the activation of the FUS3 and KSS1 kinases allowing them to phosphorylate and activate substrates critical for polarized growth that are normally activated by the SLT2 kinase during mitotic growth.
Activation of the SLT2(MPK1) MAP Kinase Homolog

Recent genetic evidence of Lee et al. (26) and Irie et al. (19) suggests that the SLT2(MPK1) kinase is activated in a cascade in which the BCK1(SLK1) kinase activates the redundant pair of kinases MKKI and MKK2, which in turn activate the SLT2(MPK1) kinase. Remarkably, the MKK kinases are most similar in sequence to a class of MAP kinase-kinases including STE7, PKS2, bpl1+, and MAPKK (15). Furthermore, Bck1p(Slk1p) is most similar in its kinase catalytic domain to the STEII kinase which is thought to phosphorylate and activate the STE7 MAPKK. A dominant, activated mutant of the BCK1(SLK1) gene has been isolated as a bypass suppressor of protein kinase C mutants (pkcl). It was thus proposed that Pkclp is an activator of the Bck1p(Slk1p)–Mkklp2p–Slt2p(Mpklp) cascade. pkcl mutants have an osmotic–remedial cell cycle phenotype that has been suggested to be due to cell wall defects (27, 38).

We isolated our slt2 allele in a screen for mutants that enhance the division defect of partially inactivated cdc28 mutants. This phenotypic enhancement seems to concern a G1-specific function of Cdc28p, since it was observed with the cdc28-4 allele that is mainly defective in G1 functions of Cdc28p, and it was not seen with the cdc28-2 allele that is mainly defective in G2/mitotic functions of Cdc28p. Our phenotypic analysis has suggested that Slt2p promotes polarization of cell growth to the bud. Cdc28p activation at Start in G1-phase also promotes bud emergence. We thus propose that the SLT2 pathway is either a downstream activation target of the CDC28 kinase or that Slt2p and Cdc28p function in parallel in promoting bud emergence and subsequent growth. A Xenopus 42-kD MAP kinase can be activated by the addition of CDC2-MPF to interphase oocyte extracts. CDC2-MPF functions upstream of a MAPKK kinase in this system (24). 42- and 44-kD MAP kinases are also activated biphasically in the GI/S and G2/M phases of Chinese hamster ovary cells (39). Ultimately, direct biochemical assays will determine whether Slt2p(Mplp) activation is dependent on Cdc28p and/or Pkclp and whether its activation varies during the cell division cycle.

We thank Corinne Le Moal for technical assistance in preparing specimens for electron microscopy and we thank Christine Costigan and Mike Snyder for strains and discussions. Thanks as well to John Cooper, Mike Gustin and John Pringle for discussions and to Catherine Jackson for reviewing the manuscript.

Cristina Mazzoni was supported by a fellowship from the Istituto Pasteur Fondazione Cenci Bolognetti in Italy and Patrick Zarrow by a fellowship from the Istituto Pasteur vielle-Sebastia, and F. Lacroute. 1991. A family of low and high copy replicative, Integrative and single-stranded S. cerevisiae/E. coli shuttle vectors. Yeast. 7:609–615.

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