Ste20-like protein kinases are required for normal localization of cell growth and for cytokinesis in budding yeast

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The yeast Ste20 protein kinase is involved in pheromone response. Mammalian homologs of Ste20 exist, but their function remains unknown. We identified a novel yeast STE20 homolog, CLA4, in a screen for mutations lethal in the absence of the G1, cyclins Cln1 and Cln2. Cla4 is involved in budding and cytokinesis and interacts with Cdc42, a GTPase required for polarized cell growth. Despite a cytokinesis defect, claa4 mutants are viable. However, double claa4 ste20 mutants cannot maintain septin rings at the bud neck and cannot undergo cytokinesis. Mutations in CDC12, which encodes one of the septins, were found in the same screen. Cla4 and Ste20 kinases apparently share a function in localizing cell growth with respect to the septin ring.

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Mechanisms of cytokinesis are extremely variable among eukaryotes. One variant of this theme is cell division by budding, typical for yeasts. The cleavage furrow in animal cells or the preprophase band (a cytoskeletal structure determining the future cell division plane) in plant cells forms only after most of the growth required to double the mass of the cell has occurred, whereas in budding, the formation of the "cleavage furrow" [bud neck] precedes the bulk of cell growth. Unlike cleavage or fission, budding is dependent on the ability to direct growth exclusively to one side of the division plane—to the bud. Despite differences in timing and morphology, at least some of the molecular mechanisms underlying budding and cytokinesis are conserved among eukaryotes (see e.g., Shinjo et al. 1990; Manser et al. 1994; Neufeld and Rubin 1994). Studies in yeast may be as useful for understanding cytokinesis and its regulation as they have been for elucidating the mechanisms of cell cycle control by cyclin-dependent protein kinases (for reviews, see Nasmyth 1993, Pines 1993).

Like other cell cycle processes, the timing of cytokinesis must be carefully regulated. In yeast, the first step toward cytokinesis is budding. The processes leading to bud emergence begin in late G1, concurrently with the onset of DNA replication and with the duplication of the spindle pole body. The simultaneous onset of these three processes [often referred to as "Start"] occurs as a consequence of the activation of a cyclin-dependent protein kinase, Cdc28, by at least one of the G1, cyclins Cln1, Cln2, and Cln3. Under normal conditions, Cln3 induces the accumulation of Cln1 and Cln2, which then trigger budding and DNA replication [Richardson et al. 1989; Dirick and Nasmyth 1991; Tyers et al. 1993; L. Dirick, pers. comm.].

Bud emergence results from several distinct events [for review, see Chant 1994]. The cell must choose a site on its surface (the bud site) and direct growth toward this site, ensuring at the same time that no other part of the cell exterior grows.

The first event necessary for budding—bud site establishment—requires the products of multiple genes including CDC42, which codes for a Rho-like GTPase that is localized at the bud site [Johnson and Pringle 1990; Ziman et al. 1993]. CDC24, coding for a GDP/GTP exchange factor [Sloat et al. 1981; Zheng et al. 1994], and CDC43/CAL1, encoding a prenyl transferase required to attach Cdc42 to membranes [Johnson et al. 1990; Ohya et al. 1995]. The nonessential genes RSRI/BUD1, BUD2, BUD3, BUD4, and BUD5 are responsible for nonrandom localization of the bud site with respect to the previous bud [see Chant 1994]. One of them, BUD2, is essential for budding in cells lacking Cln1 and Cln2, suggesting that bud emergence is controlled by Cln1 and Cln2 [Benton et al. 1993; Cvrčková and Nasmyth 1993].

A second prerequisite for budding is a change in cell growth mode that occurs at Start. Pre-Start cells grow in a nonpolar manner, increasing their size but maintaining their shape. After Start, growth becomes limited to a part of the cell—the nascent bud. Polarization of growth is
accompanied by translocation of cortical actin to the growing area, triggered by Cdc28 and Cln1 or Cln2. Overexpression of Cln1 or Cln2 causes "hyperpolarized" growth, suggesting that Cln1 and Cln2 have a role in directing growth toward the bud site (or toward the bud tip in budded cells). Later in the cell cycle, Cln cyclins are replaced by the mitotic B cyclins, at which point the bud begins to expand isometrically (Lew and Reed 1993).

The third prerequisite of normal bud development is the establishment of a "border" between the bud and the mother cell and restriction of surface growth to the bud. Unlike the previous steps, which are required not only for budding but also for the morphological changes ("shmooing") in haploid cells differentiating into gametes, border formation is specific for budding. The border is marked by a ring of 10-nm filaments attached to the cytoplasmic membrane. The ring forms around the bud site at the time of bud emergence and persists at the bud neck till cytokinesis (Byers and Goetsch 1976). Mutants in CDC3, CDC10, CDC11, and CDC12 fail to assemble the ring, produce malformed buds, and cannot undergo cytokinesis. CDC3, CDC10, CDC11 and CDC12 encode structural components of the ring (Haarer and Pringle 1987; Ford and Pringle 1991; Kim et al. 1991), which are proteins closely related to each other, to products of several anonymous mammalian cDNAs and to the Drosophila peanut (Pnut) protein required for cytokinesis (Neufeld and Rubin 1994). Members of this protein family were termed "septins" (Sander and Field 1994).

Other proteins involved in budding are also evolutionarily conserved. Homologs of Cdc42 have been found in higher eukaryotes (Shinjo et al. 1990). Their biological function remains largely unknown, the expression of mutationaly activated Cdc42 in fibroblasts may cause a cytokinesis defect (T. Leung, E.M. Lim, and L. Lim, unpubl.). Biochemical partners of Cdc42 in mammalian brain tissue were found (Manser et al. 1993, 1994). One of these partners, the Pak protein kinase, is related to a yeast protein—Ste20—necessary for the response of haploid cells to sexual pheromones. Ste20 transduces the signal elicited by the binding of pheromone to a membrane receptor toward a protein kinase cascade consisting of Ste11, Ste7, and Fus3/Kss1, one of several yeast homologs of the mammalian MAPK system. Induction of the kinase cascade leads to transcriptional activation of pheromone-regulated genes and to cell cycle arrest in G1, as a prelude to conjugation. Loss of Ste20 or other components of the pheromone response pathway does not affect budding or the establishment of cell polarity in haploids (Leberer et al. 1992; for review, see Ammerer 1994). The interaction between the mammalian Ste20 and Cdc42 homologs was therefore surprising. So far, there is only one observation suggesting a role for Ste20 in morphogenesis: Ste20 is, among other components of the pheromone response pathway, required for the formation of pseudomycelia in nitrogen-starved diploid cells (Liu et al. 1993). It is, however, unclear whether Ste20 functions in cell morphogenesis or in perceiving nutrient conditions during pseudomycelial differentiation.

How is budding linked to the rest of the cell cycle? We described previously the isolation of mutants defective in bud emergence only in the absence of Cln1 and Cln2 and proposed that Cln1 and Cln2 are involved in bud site establishment (Cvréková and Nasmyth 1993). The same genetic screen yielded mutants unable to undergo cytokinesis because of a defect in bud neck morphogenesis, suggesting that Cln1 and Cln2 have a role also in determining the border between the mother cell and its bud. One of the genes required for cytokinesis in cln1 cln2 cells, CLA10, is allelic to CDC12, suggesting that septin ring assembly or function may be controlled by Cln1 and Cln2. Other mutants with a similar phenotype define a novel gene, CLA4, that codes for a putative protein kinase related to Ste20 and Pak. Similar to Pak, Cla4 binds Cdc42. CLA4 is a nonessential gene, but Cla4 and Ste20 share an essential function required for the restriction of cell growth to the portion of cell surface enclosed by the septin ring. The phenotype of cla4 ste20 double mutants identifies not only a role for Ste20-like protein kinases in morphogenesis, but also a novel step in bud development.

Results

Mutations in CLA4 and CLA10 cause a cytokinesis defect in cells lacking Cln1 and Cln2

To find additional regulators of Start, we set up a screen for yeast mutants unable to survive in the absence of the two G1 cyclins, Cln1 and Cln2 [cln mutants—Cln activity dependent]. Unexpectedly, no novel genes involved in all aspects of Start were found. We described previously the bud emergence mutants obtained in this screen and proposed that Cln1 and Cln2 have a role in the control of bud emergence (Cvréková and Nasmyth 1993).

Other mutants, especially cla4 (two alleles characterized, allelic to erc10 described by Benton et al. 1993), clal6 (two alleles), and clal0 (live alleles), suffer from defects in cytokinesis, that is, at a very late stage of the cell cycle. This was surprising because G1 cyclins are thought to act early. In the absence of Cln1 and Cln2, cla4, clal6, and clal0 mutant cells divide their nuclei and bud, but they cannot undergo cytokinesis [Fig. 1]. Subsequent observations revealed that even cultures of the viable cln1 cln2 mutant contain cells with broad bud necks and cell clusters resulting from delayed cytokinesis. This further supports the idea that Cln1 and Cln2 have a role in bud development and cytokinesis in addition to the previously reported function in bud emergence; the apparent cytokinesis failure in cla4, clal6, and clal0 mutants could perhaps be a late consequence of an early defect.

CLA10 is allelic to CDC12

The wild-type CLA10 gene was cloned by complementation of cla10l mutation and sequenced (EMBL/GenBank accession no. X82498). The cloned DNA fragment was shown to be linked to the CLA10 locus. Sequence comparison revealed that CLA10 is allelic to CDC12.
A possible explanation for the phenotypic defects caused by a mutation in one of the septins is that phosphorylation of some protein associated with the septin ring by Cln1- or Cln2-coupled mitotic kinases becomes essential if a component of the septin ring is damaged by mutation.

We cloned the minimal complementing fragment for the cla4-1 allele (see Materials and methods). Diploid cells heterozygous for cla4::LEU2 and Cla4+ alleles were seen in liquid media (Figs. 3 and 4, below). Homozygous cla4::LEU2 diploid cells grew more slowly than wild-type or heterozygous controls. We conclude that Cla4 is a nonessential gene.

CLA4 codes for a novel protein kinase

We cloned the CLA4 gene by complementation of the cla4-1 mutation, confirmed that the cloned DNA is linked to CLA4, and sequenced the gene. The minimal complementing fragment contained a single open reading frame. Northern analysis of RNA from a synchronous culture revealed that CLA4 is transcribed throughout the cell cycle (data not shown). CLA4 (EMBL/GenBank accession no. X82499) encodes a protein of 842 amino acids, with a putative protein serine/threonine kinase domain on its carboxyl terminus and two conserved motifs in its amino-terminal part (Fig. 2). One of these motifs is the Pleckstrin homology [PH] domain, shared by several proteins interacting with GTP-binding proteins and other components of signaling pathways (see Mustacchio et al. 1993). The PH domain can bind phosphatidylinositol-4,5-bisphosphate in vitro (Harlan et al. 1994). The other motif, flanking the carboxy-terminal end of the PH domain, is shared by two kinases whose overall structure is similar to that of Cla4: the mammalian brain-specific kinase Pak [Manser et al. 1994] and the yeast Ste20 kinase [Leberer et al. 1992, Ramer and Davis 1993]. Neither Ste20 nor Pak contains a PH domain. In Cla4, the Pak/Ste20 homology region contains a good match to the sequence motif SHXPHxxH shared by several kinases associated with Rho-related small GTPases. In Pak, this region binds the GTP-bound form of two Rho-like proteins, Cdc42Hs and Rac1 (Manser et al. 1994). Cdc42Hs is a human homolog of the yeast Cdc42 GTPase required for bud emergence.

To study the role of CLA4 in wild-type cells, we replaced the Cla4 kinase domain by the LEU2 gene, producing the cla4::LEU2 allele (see Materials and methods). Diploid cells heterozygous for cla4::LEU2 were indistinguishable from the wild type with respect to colony growth and cell morphology. Haploid cla4::LEU2 cells derived by sporulation of this diploid were viable and grew at a rate comparable to wild-type haploids (data not shown), but they had grossly abnormal morphology. They produced rough colonies containing clusters of deformed cells apparently defective in cytokinesis; similar defects were seen in liquid media (Figs. 3 and 4, below). Homozygous cla4::LEU2 diploid cells grew more slowly than wild-type or heterozygous controls. We conclude that CLA4 is required for normal cell morphology and possibly for proper timing of cytokinesis.

Deletion of CLA4 in cells lacking any single CLN gene did not cause any additional phenotype. Triple cln1 cln2 cla4::LEU2 or cln2 cln3 cla4::LEU2 mutants are viable, though they grow more slowly and have a more severe cytokinesis defect than cla4 mutants. One could argue that the lethality of cln1 cln2 cla4 mutants results from a decrease in fitness owing to the combination of three mildly deleterious mutations. However, loss of Clb5 and Clb6, the pair of B-type cyclins involved in S-phase regulation, has as great an effect on growth rate and cell size as deletion of CLN1 and CLN2 (Schwob and Nasmyth 1993), but clb5 clb6 cla4::LEU2 mutants are viable. This indicates that CLA4 is not required for S-phase entry.
is consistent with Cln1 and Cln2 having a specific role in bud morphogenesis and cytokinesis.

**Physical and genetical interactions between Cla4 and Cdc42**

The presence of a putative GTP–Cdc42-binding site in Cla4 suggests that in spite of the different mutant phenotypes, Cla4 and Cdc42 may be involved in the same process. Binding between Cdc42 and Cla4 could, for instance, be needed for localization or spatially restricted activation of the Cla4 kinase at the bud site, which is marked by Cdc42, and local activity of Cla4 may be required for proper function of the septin ring. We therefore decided to test whether Cla4 could bind to GTP–Cdc42 in vitro.

We could not express full-length Cla4 as a glutathione S-transferase (GST) fusion protein in *Escherichia coli*, but two fragments of Cla4, containing the putative Cdc42-binding site, were successfully purified. Both fragments bind GTP-loaded yeast and human Cdc42 efficiently in overlay assays. The related yeast Rho1 GTPase is not bound (Fig. 3). Yeast GTP–Cdc42 binds better to the Cla4 fragments than to the mammalian Pak kinase, whereas the converse is true for human GTP–Cdc42. No such difference was observed with another mammalian Cdc42-binding kinase, Ack (Manser et al. 1993).

Interaction between Cla4 and Cdc42 was also detected using the two-hybrid system in vivo (Table 1). No interaction was seen between full-length, normal Cdc42 and either full-length Cla4 or a fragment of Cla4 that bound Cdc42 in vitro (line 1). However, an interaction was found with a mutant Cdc42 that should be unprenylatable and hence less tightly (if at all) membrane associated (line 2). The interaction was stronger when Cdc42 also contained mutations inactivating its GTPase activity and thus locking it in the GTP-bound state (lines 3 and 4) and disappeared when Cdc42 contained a mutation believed to lock it in the GDP-bound state (line 5). To avoid high backgrounds observed with the unprenylated Cdc42, Ack was not used as a control in these experiments. The results demonstrate that Cla4 and Cdc42 can interact both in vitro and in vivo.
latable Cdc42 (lines 2–4), the tests were repeated with the vectors reversed [lines 16–19]. In this case, background values were low, and a weak but significant interaction was detected between full-length Cla4 and the GTPase-defective, unprenylatable Cdc42 [line 16]. More striking results were obtained with the truncated Cla4 [line 17], where a very weak interaction was detected even with normal Cdc42. The interaction was ~30-fold stronger for the unprenylatable Cdc42 and another 2.6-fold stronger for Cdc42 locked in the GTP-bound form. No interactions were detected between either full-length or truncated Cla4 and a variety of other small GTPases [lines 6–15].

We conclude that Cla4 directly interacts with Cdc42, that this interaction is specific for Cdc42 among the family of small GTPases, that the interaction is much stronger (and perhaps only occurs) with GTP-bound Cdc42, and that the interaction is mediated by a site within amino acids 174–294 of Cla4.

In Pak, the GTP–Cdc42-binding domain regulates autophosphorylation-mediated kinase activation. Cla4 could also be regulated by its interaction with GTP–Cdc42, that is, Cdc42 might have another role in cytokinesis in addition to its known function required for polarized cell growth. This second function might have escaped detection, because polarized cell growth is itself required for budding. Perturbation of the GTPase cycle of Cdc42 can cause a phenotype reminiscent of that caused by loss of Cla4. In our strain background, cdc42-1 mutant cells often produce elongated buds at the permissive temperature [Fig. 3]. Some mutant alleles of CDC24, a gene encoding the GDP/GTP exchange factor for Cdc42, produce elongated buds at semipermissive temperature [Sloat et al. 1981]. Consistent with an interaction between Cla4 and Cdc42 in vivo is the finding that double cla4 cdc42-1 mutants are inviable at the permissive temperature for cdc42-1. We did not obtain any double mutants from 22 tetrads of a diploid strain heterozygous for cla4::LEU2 and cdc42-1. All 10 spores predicted to have the cla4 cdc42-1 genotype germinated and underwent one to four cell divisions, producing finally a microcolony of misshapen dead cells (Fig. 3).

Cla4 and Ste20 share an essential function

The similarity between Cla4 and Ste20 suggests that the two kinases might have overlapping functions. Because the loss of STE20 in one of our strain backgrounds (K1107) leads only to a decrease in mating efficiency but not to sterility [B. Stevenson, pers. comm.], we could obtain a diploid strain heterozygous for cla4::LEU2 and cdc42-1. All 10 spores predicted to have the cla4 cdc42-1 genotype germinated and underwent one to four cell divisions, producing finally a microcolony of misshapen dead cells (Fig. 3).

Figure 3. (a) Cla4 binds Cdc42–GTP. [Left] Recombinant GST fusion proteins separated on a SDS–polyacrylamide gel and stained with Coomassie blue. CLA4-A corresponds to Cla4 residues 174–279, CLA4-B to Cla4 residues 174–222, PAK to Pak residues 69–150, and ACK to Ack residues 499–570. CLA4-B corresponds to the minimal p21-binding domain described for Ack [Manser et al. 1993]. [Middle and right] p21 overlay assays on parallel gels with the indicated recombinant p21 proteins. (b) cla4 and cdc42 mutations are synthetic lethal. Microcolonies grown from spores of a diploid (K5057) heterozygous for cdc42-1 and cla4::LEU2 after 24 hr on YEPD at 25°C.
that Clα4 and Ste20 share a function essential for vegetative growth.

Clα4 does not act through the pheromone signaling or protein kinase C (PKC1) pathway

Ste20 transduces the signal elicited by the binding of the mating pheromone to its receptor toward a protein kinase cascade related to the mammalian mitogen-activated protein kinase [MAPK] pathway [for review, see Ammerer 1994]. The pheromone response system consists of kinases encoded by STE11, STE7, and FUS3 or KSS1; the Ste5 protein [which is not a kinase] is also required for signal transduction, possibly at multiple steps. One explanation for the lethality of clα4 ste20 double mutants is that the Clα4 and Ste20 protein kinases share substrates whose phosphorylation is needed for cytokinesis. Alternatively, Clα4 might share a direct or indirect target with an element of the pheromone response system downstream of Ste20. If the latter were true, constitutive, Ste20-independent activation of the pheromone response pathway should abolish the requirement for Ste20 in clα4 mutants, and disruption of the pheromone response pathway downstream of Ste20 may be as deleterious for the clα4 mutants as the loss of Ste20 itself.

Low-level ectopic expression of the dominant hyper-activated STE11-279s allele constitutively activates pheromone response and is sufficient to restore mating to ste20 mutants in a strain background where these are sterile [Stevenson et al. 1992]. However, it does not affect the phenotype of MET::C1α4 ste20 mutants on media containing methionine. Furthermore, double clα4 ste5 and clα4 ste11 mutants are viable. The essential function of Ste20 in clα4 cells does not therefore require known components of the pheromone response pathway immediately downstream of Ste20.

The functional redundancy between Clα4 and Ste20 might be attributable to the ability of Ste20 to activate another MAPK-like kinase cascade. The essential protein kinase C [PKC1]-dependent system, consisting of the Bck1, Mkk1/Mkk2, and Mpk1 kinases, is required for the maintenance of cell wall integrity. Constitutive activation of the PKC1 pathway by the dominant BCK1-20 allele [Lee and Levin 1992] does not affect the phenotype of MET::C1α4 ste20 mutants on media containing methionine [data not shown]; therefore, this system is unlikely to be involved in the essential function common to Clα4 and Ste20.
Activation of the high osmolarity response (HOG) pathway may compensate for loss of Cla4

Another MAPK-like kinase cascade known to function in vegetative yeast cells—the HOG [high osmolarity glycerol] response system—is required for proliferation on high osmolarity media. Mutants lacking either of the bottom two kinases of this cascade—Pbs2 or Hog1—cannot accumulate intracellular glycerol nor transcribe some osmolarity-regulated genes nor undergo cytokinesis on high salt media [Brewster et al. 1993]. The phenotype of cla4 mutants is reminiscent of the described defect of hog1 and pbs2 mutants on high salt media, suggesting that a Cla4-dependent mechanism and the HOG system might have a similar role, the former being functional only on low salt media, the later in a high osmolarity environment.

If this were the case, increased salt concentration in the media might rescue the morphological defect caused by the loss of Cla4. It is therefore remarkable that cla4::LEU2 mutants acquire an almost wild-type shape in high salt media [Fig. 4]. Disruption of either HOG1 or PBS2 aggravates the morphological phenotype of cla4 on standard media and abolishes the rescue by salt [Fig. 4]. However, the latter result should be interpreted cautiously, because single hog1 or pbs2 mutants are already impaired under the same conditions. Nevertheless, our observations are consistent with the idea that the HOG pathway and Cla4 have common functions.

Cla4 or Ste20 is required for proper function of the septin ring

The phenotype of cla4 mutants and the isolation of cdc12 mutants in the same genetic screen suggest that Cla4 may be needed for proper function of the septin ring. Additional support for this hypothesis comes from the finding that double cla4::LEU2 cdc12-1 mutants are inviable or at least severely impaired even at permissive temperature for the cdc12-1 mutant: no viable double mutants were obtained in 19 tetrads from a diploid heterozygous for the two mutations. We thus decided to study the behavior of septins in cla4 mutants.

To examine the phenotype caused by the loss of the essential function of Cla4, we generated a thermosensitive allele of CLA4 [cla4-75] by PCR mutagenesis of the cloned gene and introduced it into a strain lacking CLA4 and STE20. The resulting cla4-75 ste20 strain was thermosensitive for growth, exhibiting a uniform first cell cycle arrest with a phenotype similar to the original cln1 cla2 cla4 mutant. To characterize this phenotype more thoroughly, we compared the progression of the cell cycle of CLA4 ste20 and cla4-75 ste20 cells. G1 cells were isolated by centrifugal elutriation and grown at 37°C, the nonpermissive temperature for cla4-75. Inactivation of Cla4 in cla4-75 ste20 cells did not affect the timing of DNA replication, mitotic spindle formation or budding compared with the CLA4 ste20 strain [Fig. 5]. However, buds formed at 37°C were malformed, resembling the protuberances or “shmoos” of yeast gametes preparing to mate [Fig. 6a]. Immunofluorescent staining of the Cdc11 septin revealed that a septin ring forms in mutant cells at the same time and place [i.e., at the site of “bud/shmoo” emergence] as in ste20 cells that do not have any cytokinesis defect. However, at later stages the ring appears in most cases at the tip of the malformed bud rather than at a defined bud neck and often becomes diffuse [Fig. 6b]. Whereas wild-type cells deposit new cell wall only on the bud side of the septin ring, cla4 ste20 mutant cells grow new wall mostly if not exclusively on the mother side of the ring [Fig. 6c]. Cytokinesis is required for proper budding at this site. The clamp-like activity of Cdc42 [i.e., toward the bud tip] in cla4 ste20 cells, suggesting that at least some Cdc42-dependent events remain undisturbed in the double mutant [Fig. 7]. In addition to the

Figure 4. (a) Loss of Cla4 causes a morphological defect. [Left] cla4::LEU2 [K3591], [right] wild-type [K699] cells grown in YEPD. (b) This defect is compensated for under conditions inducing the HOG pathway. [Left] cla4::LEU2 [K3591], [right] wild-type [K699] cells grown in YEPD + 0.4 M NaCl. (c) This rescue requires the HOG pathway. [Left] cla4::LEU2 hog1::TRP1 [K4329] or cla4::LEU2 pbs2::URA3 [K4333] cells, [right] hog1::TRP1 [K4327] or pbs2::URA3 [K4331] cells grown in YEPD or YEPD + 0.4 M NaCl.
Figure 5. CLA4 ste20 and CLA4 ste20 cells (K3621) were synchronized by centrifugal elutriation and released at 37°C (the restrictive temperature for cla4-75). DNA replication was followed by FACS analysis, fractions of unbudded cells, cells with mitotic spindles of various stages, and Cdc11-containing structures were determined microscopically. (a) cla4-75 ste20 and CLA4 ste20 cells undergo DNA replication and acquire polarized shape at the same time (arrows denote the time when half of the culture budded). (b) Initiation of DNA replication, local accumulation of Cdc11, budding, and mitotic spindle formation occur at a comparable cell size in mutant and control cells, whereas spindle elongation is delayed in the cla4-75 ste20 mutant.

described phenotype, cla4 ste20 mutants have a delay in mitotic spindle elongation: only about a third of the cells undergoes anaphase within 4 hr, and many of the others die and lyse as uninucleate cells (Fig. 5b; data not shown), suggesting that bud neck formation may not be the only event dependent on Cla4/Ste20. More likely, proper establishment of bud neck may be required for normal timing of nuclear division, as in the case of bud emergence (Lew and Reed 1995).

It is remarkable that cla4 ste20 cells assemble the septin ring but fail to develop the constriction ("neck") at the base of the bud. Septin ring assembly, although required, is therefore not sufficient for the establishment of a normal border between the mother cell and the bud. The phenotype of cla4-75 ste20 mutants seems to define a novel step in bud neck development: Cla4 and Ste20 share a function necessary for proper localization of cell surface growth with respect to the septin ring (or vice versa).

Discussion

\section*{G1 cyclins have a role in cytokinesis}

Cyclin-dependent protein kinases (Cdks) are responsible for the timing of cell cycle events in eukaryotic cells. Cytokinesis is one of the processes likely to be controlled by Cdks. The first step toward cytokinesis in yeast—budding—occurs in late G1, simultaneously with the onset of DNA replication and spindle pole body duplication. These three events (often referred to as Start) are all regulated by a Cdk, Cdc28, associated with the G1 cyclins Cln1, Cln2, and Cln3. Mutants lacking one or two of the G1 cyclins are viable, whereas the triple cln1 cln2 cln3 mutant cannot undergo Start (Richardson et al. 1989). This, however, does not mean that the roles of the three G1 cyclins are equivalent. The concentration of Cln3 remains constant throughout the cell cycle, while the abundance of Cln1 and Cln2 and the associated kinase activity oscillate during the cell cycle with a maximum around Start. Cln3 is probably involved in the transcriptional activation of Cln1 and Cln2 in late G1 (Cross and Tinkelenberg 1991; Dirick and Nasmyth 1991; Tyers et al. 1993; L. Dirick and K. Nasmyth, unpubl.) and in the activation of the B-type cyclins Clb5 and Clb6 that trigger DNA replication. The observed functional overlap of the G1 cyclins results from the capability of Cln1 and Cln2 to induce their own—delayed—accumulation and the transcription of Clb5 and Clb6 in the absence of Cln3 and from the ability of Clb5 and Clb6 to provide the function of Cln1 and Cln2, albeit with a decreased efficiency (Epstein and Cross 1992; Schwob and Nasmyth 1993).
The isolation of two genes involved in cytokinesis (CLA10/CDC12 and CLA4) in a screen for genes required for proliferation in the absence of Cln1 and Cln2 suggests that some proteins required for an early step toward cell division may be a direct substrate for the Cdc28/Cln complex. The regulated step could be the establishment of the bud neck—an event that involves the assembly of a septin ring on the border between the mother cell and its bud, one of the components of the ring is encoded by CDC12. In the absence of Cln1 and Cln2, their function can be provided by another cyclin present at the same stage of the cell cycle (e.g., by Cib5 and Cib6), as long as all other proteins required for bud neck establishment and septin ring assembly are intact. If a component of the ring is damaged (as in cdc12 mutants) or a regulatory event (e.g., phosphorylation of some protein by the protein kinase encoded by CLA4) is missing, Cln1 and Cln2 may become the only cyclins able to induce proper bud neck development. We proposed previously a similar model suggesting a direct role for Cln1 and Cln2 in the first step toward budding—in bud site establishment and cell polarization (Cvrčková and Nasmyth 1993). This model can now be extended—also the next step in bud development, the formation of the border between the bud and its mother cell, appears to be controlled by G1 cyclins. The apparent evolutionary conservation of cytokinesis mechanisms indicates that a similar control [perhaps involving different Cdk species] may be at work also in higher eukaryotes.

Figure 7. Cla4 and Ste20 are not required for normal distribution of cortical actin. Exponential cultures of cla4-75 ste20 (K4580) or CLA4 ste20 (K3621) cells grown at 25°C in YEPD were shifted to 37°C for 3 hr and fixed, and actin was stained with rhodamin-labeled phallidin.

Figure 6. cla4-75 ste20 cells produce deformed buds with mis-localized septins at restrictive temperature and deposit new cell wall proximally to the septins. [a] Exponential cultures of cla4-75 ste20 or CLA4 ste20 cells grown at 25°C in YEPD were shifted to 37°C for 3 hr and observed by phase-contrast microscopy. [b] Typical cells from the synchronous cultures from Fig. 5., representing successive stages of the cell cycle (denoted by numbers). [Blue] DNA, [yellow] Cdc11. Because the staining requires the removal of cell wall, small buds may become difficult to see. (c) Exponential cultures of cla4-75 ste20 or CLA4 ste20 cells grown at 25°C in YEPD were shifted to 37°C for 1 hr, the cell wall was labeled with fluorescein-conjugated Concanavaline A, and cells were grown for an additional 30 min at 37°C in the absence of the label and observed by fluorescence and phase-contrast microscopy. [Left] cla4-75 ste20 (K4580), [right] CLA4 ste20 (K3621).

**Ste20-like kinases and the Cdc42 GTPase are required for cytokinesis**

Eukaryotes have evolved several ways to divide one cell into two. Most animal cells divide by cleavage, that is, by cytoskeleton-driven contraction of a part of the cell surface. Cells of higher plants form their septa “from within”, by fusion of membrane vesicles carrying cell wall material to a location determined by cytoskeletal structures, producing the cell plate, which develops into a septum. Cytokinesis by budding, typical for yeasts, is a somewhat unusual variation of this theme. Nevertheless, similar mechanisms seem to be involved in budding and cleavage. Genes required for various steps of cytokinesis in the yeast *Saccharomyces cerevisiae* have been
isolated and often shown to have relatives in higher eukaryotes.

One of the central molecules involved in the early steps toward yeast cytokinesis is the small GTPase Cdc42. In *S. cerevisiae*, Cdc42 is required for polarized cell growth both in the vegetative cell cycle (budding) and in the process of mating (asymmetric growth of gametes—shmooing). Cdc42 cycles between the GTP- and GDP-bound form; the putative guanine nucleotide exchange factor for Cdc42—Cdc24—is also required for directional growth [Zheng et al. 1994; for review, see Hall 1992].

Cdc42 is required for at least two events in bud development: One leads to actin polarization and subsequently to bud growth; the other leads to the deposition of a ring of 10-nm filaments (the septin ring) under the cytoplasmic membrane at the site of the future bud. Proteins related to components of the ring were found in animals, and a fruit fly homolog is required for cytokinesis [Neufeld and Rubin 1994].

Cdc42 homologs have also been found in higher eukaryotes. Biochemical studies identified a mammalian brain-specific protein kinase (Pak) that selectively binds to the GTP-bound form of Cdc42. Interestingly, Pak is related to the yeast Ste20 kinase, a component of the signal transduction mechanism required for differentiation of haploid cells into gametes in the presence of sexual pheromones and for pseudomycelial differentiation of diploid cells starved for nitrogen [Leberer et al. 1992; Liu et al. 1993; Ramer and Davis 1993]. Until now, there was no indication that Ste20 might be involved in budding. It is therefore somewhat surprising that Ste20 also can, like Pak, bind Cdc42-GTP [Manser et al. 1994; Z.-S. Zhao, T. Leung, E. Manser, and L. Lim, in prep.; C. De Virgilio and J.R. Pringle, unpubl.].

We have isolated a novel yeast gene, *CLA4*, whose product is closely related to Ste20 and Pak and specifically binds Cdc42-GTP both in vitro and in a two-hybrid in vivo assay. The interaction between Cla4 and Cdc42 might be regulated in vivo, because truncated forms of Cla4 bind Cdc42-GTP better than a full-length protein. The observation that *cla4 cdc42-1* double mutants are lethal under conditions permissive for single mutants indicates that the interaction between Cdc42 and Cla4 may be significant in vivo. Binding to GTP—Cdc42 activates the mammalian Pak kinase [Manser et al. 1994]; similarly, in yeast the interaction between Cla4 and GTP—Cdc42 might be needed for spatially restricted activation of the Cla4 kinase at the bud site, which is marked by Cdc42.

We found that Cla4 and Ste20 share an essential role in bud development. The previously observed lack of cell cycle-related phenotypes in *ste20* mutants was apparently a result of a functional overlap between Ste20 and Cla4. Ste20 interacts with the pheromone receptor-associated G protein and acts through a signaling pathway involving three protein kinases corresponding to the mammalian MAPK module. Unlike Ste20, other components of this pathway downstream of Ste20 are not required for cytokinesis in cells lacking Cla4. Ste20 therefore does not exert its essential function in *cla4* cells by activating the pheromone response pathway. However, some other MAPK-like kinase cascade activated by either Cla4 or Ste20 might be required for bud neck development. It is interesting that the nonlethal morphogenetic defect of *cla4* mutants is compensated under conditions that induce another MAPK-like system, the *HOG* pathway, and this rescue seems to require the bottom two kinases of the *HOG* kinase cascade. Alternatively, some protein associated with the bud neck may have to be directly phosphorylated by either Cla4 or Ste20 to become functional.

**Ste20-like kinases are required for proper localization of cell growth with respect to the division plane**

Double *cla4 ste20* mutants have a severe defect at an early stage of bud morphogenesis and cannot undergo cytokinesis. Normal bud neck development and cytokinesis require the establishment in late G1 of a ring of septin-containing filaments defining the site of future cell division. Later on, all cell growth occurs on the bud side of the ring, whereas the mother cell ceases growing. The shape of *cla4 ste20* mutants superficially resembles that of mutants defective in components of the septin ring. Moreover, mutants in *CDC12*, which codes for one of the septins, were found in the same genetic screen. Cla4 or Ste20 may thus be required for proper localization or function of the septins. However, upon closer inspection, *cla4 ste20* double mutants differ from septin-defective mutants. *cdc12-1* cells under restrictive conditions maintain a border between the mother cell and the bud, despite gross morphological abnormalities and despite the absence of a detectable septin ring [Haarer and Pringle 1987]. On the other hand, the septin ring assembles at the correct time and place in *cla4 ste20* mutant cells. However, it fails to stay at the bud neck and remains on the tip of the growing misshapen “bud” instead and usually becomes diffuse. Contrary to wild-type cells, new cell wall deposits proximally to the septin structure (i.e., on the mother cell side) in *cla4 ste20* mutants.

These observations suggest that the assembly of the septin ring, albeit essential for cytokinesis, is not the only prerequisite for development of a normal bud neck. A separate event that requires a Ste20-like protein kinase (possibly regulated by Cdc42 and the Cdc28/Cln complex) is needed to direct growth to a distinct part of the cell surface (Fig. 8). The growing part of the surface is normally enclosed by the septin ring, but the border between the growing and nongrowing surface apparently can form even in mutants defective in components of the ring itself, as documented by the phenotype of previously described cytokinesis mutants (*cdc3, cdc10, cdc11, and cdc12*) that are defective in the septin ring. Mutants lacking Cla4 and Ste20 thus define a novel event in cell morphogenesis: They are impaired in the formation of a normal border separating the bud from the mother cell.

Septins, Ste20-like protein kinases, and Cdc42-like GTPases are evolutionarily conserved. Interestingly,
mammalian homologs of these proteins are abundantly expressed in the brain, thus, in a tissue that contains rather few dividing cells. Could these proteins function in another process besides cytokinesis? The role of Cla4 and Ste20 in yeast morphogenesis suggests the possibility that a similar mechanism might be involved also in spatial restriction of cell surface growth in other situations, including, perhaps, neural cell development.

Materials and methods
Yeast strains and genetic techniques
Genotypes of the strains not fully described in the text are listed in Table 2. The cln1, cln2, and ade1 alleles were described previously (Dirick and Nasmyth 1991; Cvrčková and Nasmyth 1993), bgl1::URA3 and pbs2::LEU2 were provided by Ch. Schüller (University of Vienna, Austria), ste20::URA3 (Leberer et al. 1992) and the plasmid carrying the STE11-279S allele under the GAL1–GAL10 promoter (Stevenson et al. 1992) were provided by B. Stevenson. BCK1-20 on a centromeric plasmid was a gift from D. Levin (Johns Hopkins University, Baltimore, MD). Standard genetic techniques (Mortimer and Hawthorne 1969) and standard methods of DNA manipulation (Ausubel et al. 1987) were used throughout. Yeast cells were transformed by the spheroplast or LiAc methods. Integrative transformations were verified by Southern blots. Yeast cells were grown aerobically at 30°C on YE media (see Nasmyth and Dirick 1991) supplemented with 20 g/liter of the indicated sugars (D-glucose, Gal–galactose, Raft–raffinose), unless otherwise stated. Standard synthetic media supplemented with appropriate amino acids and nucleotides were used for experiments involving the methionine-regulated constructs; methionine was added to final concentration 2 mM to repress the MET3 promoter. Plates containing 5-FOA (Boeke et al. 1987) were made as described by Nasmyth and Dirick (1991).

Table 2. Genotypes of yeast strains

<table>
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<tr>
<th>Strain</th>
<th>Genotype</th>
<th>Origin</th>
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<td>K699</td>
<td>MATa, ade2-1, trp1-1, can1-100, leu2-3,112, his3-11,15, ura3, GAL::pss1, ssd1-d2(cla1)</td>
<td>W303 IA</td>
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<td>Dirick and Nasmyth (1991)</td>
</tr>
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<td>MATa, trp1, leu2, ura3, his3, ade2-1, ade3::HIS3, cln1::His3, cln2::del, SDL1, [YCp URA3 ADE3 spADH::CLN2]</td>
<td>Cvrčková and Nasmyth (1993)</td>
</tr>
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Plasmid-borne markers are enclosed in square brackets. [bkg] The strain was obtained by transformation of the given parent. [mut] it was derived from the given genetic background and carries the same genetic markers unless stated otherwise by multiple transformations and crosses of isogenic strains; [mut] an EMS-induced mutant.
Isolation of CLA10 and CLA4

Wild type CLA10/CDC12 and CLA4 were cloned by complementation of cla10-1 and cla4-1 mutants, respectively. The cloning strategy, the genomic library used, and the sequencing procedure were as described [Cvrčková and Nasmyth 1993]. For CLA10 isolation, 24,000 library transformants yielded 10 complementing clones. A 2.94-kb region shared by all of them was sequenced. For CLA4, six clones complementing cla-1 were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were found among 20,000 library transformants; additional clones were...
mutagenic PCR (Amon et al. 1993). A library of PCR products in YCplac22 (Gietz and Sugino 1988) was screened for plasmids that suppress the lethality of MET::CLN4 ste20 (K4220) cells in the presence of methionine at 25°C but not at 37°C. Four such plasmids were isolated and among 18,000 transformants were shown to be thermosensitive for the ability to rescue growth of cla1 cla2 cla4-75 GAL::CLN1 (K3546) cells on glucose. One of the mutant alleles—cla4-75—was chosen for further study. To obtain a temperature-sensitive cla4-75 ste20 strain, a plasmid carrying cla4-75 was introduced into a diploid (K3622) heterozygous for CLA4 and STE20, and haploids were isolated by tetrad dissection.

Synchronous cultures

Cells were grown at 25°C in YEPD and synchronized by centrifugal elutriation as described (Schwob and Nasmyth 1993). Elutriated G1 cells were inoculated into YEPD medium at 37°C and followed over the period of 4 hr, determining periodically the fraction of budded cells [microscopically] and cell size (using the CASY1 apparatus, Schärfe Systems GmbH) and taking samples for immunofluorescence staining and for the analysis of DNA content. The DNA content was measured on a Becon-Dickinson FACScan analyzer as described (Epstein and Cross 1992, Lew et al. 1992). For quantitative evaluation of microcopy data, at least 100 cells were counted for each sample and characteristic.

Fluorescence microscopy

For in situ immunofluorescence, cells were fixed with 0.05 ml of the same medium with 0.12 formaldehyde for 30 min, converted to spheroplasts, and stained for in situ immunofluorescence, cells were fixed with 3.7% formaldehyde (Ford and Pringle 1991). For staining of Cdcll, a rabbit polyclonal antibody (Ford and Pringle 1991) and a FITC-labeled DAPI. For in situ immunofluorescence, cells were fixed with 3.7% formaldehyde (Ford and Pringle 1991) and a FITC-labeled DAPI. For staining of Cdcll, a rabbit polyclonal antibody (Ford and Pringle 1991) and a FITC-labeled DAPI. For staining of Cdcll, a rabbit polyclonal antibody (Ford and Pringle 1991) and a FITC-labeled DAPI.

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