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# Synthetic Lethal Analysis Implicates Ste20p, a p21-activated Protein Kinase, in Polarisome Activation<sup>□</sup>

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Submitted June 17, 2002; October 7, 2002; Accepted December 4, 2002  
Monitoring Editor: David Drubin

The p21-activated kinases Ste20p and Cla4p carry out undefined functions that are essential for viability during budding in *Saccharomyces cerevisiae*. To gain insight into the roles of Ste20p, we have used a synthetic lethal mutant screen to identify additional genes that are required in the absence of Cla4p. Altogether, we identified 65 genes, including genes with roles in cell polarity, mitosis, and cell wall maintenance. Herein, we focus on a set that defines a function carried out by Bni1p and several of its interacting proteins. We found that Bni1p and a group of proteins that complex with Bni1p (Bud6p, Spa2p, and Pea2p) are essential in a *cla4Δ* mutant background. Bni1p, Bud6p, Spa2p, and Pea2p are members of a group of polarity determining proteins referred to as the polarisome. Loss of polarisome proteins from a *cla4Δ* strain causes cells to form elongated buds that have mislocalized septin rings. In contrast, other proteins that interact with or functionally associate with Bni1p and have roles in nuclear migration and cytokinesis, including Num1p and Hof1p, are not essential in the absence of Cla4p. Finally, we have found that Bni1p is phosphorylated *in vivo*, and a substantial portion of this phosphorylation is dependent on STE20. Together, these results suggest that one function of Ste20p may be to activate the polarisome complex by phosphorylation of Bni1p.

## INTRODUCTION

Yeast cells undergo polarized growth during budding and other morphogenetic events in response to intracellular or extracellular cues (Drubin and Nelson, 1996). Polarized cell growth depends on assembly of a polarized actin cytoskeleton, which then directs transport of secretory vesicles containing cell wall and plasma membrane components to the site of growth (Novick and Botstein, 1985; Mulholland *et al.*, 1994; Ayscough *et al.*, 1997; Pruyne *et al.*, 1998). The p21 GTPase Cdc42p plays a critical role in the establishment of subcellular polarity and the execution of subsequent apical growth by regulating the actin cytoskeleton (Adams *et al.*,

1990; Johnson and Pringle, 1990; Ziman *et al.*, 1993; Li *et al.*, 1995; Richman and Johnson, 2000). Cdc42p is also required for septin ring function and for cytokinesis (Richman *et al.*, 1999; Toenjes *et al.*, 1999). How Cdc42p orchestrates these various activities is poorly understood, but some of its target effectors have been identified. For example, Gic1p and Gic2p can bind activated Cdc42p and are important for polarization of the actin cytoskeleton (Brown *et al.*, 1997; Chen *et al.*, 1997). Two other identified effectors for Cdc42p are the related p21-activated protein kinases, Cla4p and Ste20p, both of which interact with activated Cdc42p and localize to sites of polarized growth (Cvrckova *et al.*, 1995; Peter *et al.*, 1996; Leberer *et al.*, 1997; Holly and Blumer, 1999; Mosch *et al.*, 2001). Each kinase has unique roles in the cell. Ste20p functions in pheromone response and haploid invasive growth, whereas Cla4p promotes normal septin function (Ramer and Davis, 1993; Roberts and Fink, 1994; Benton *et al.*, 1997; Tjandra *et al.*, 1998; Gulli *et al.*, 2000; Bose *et al.*, 2001). A cell lacking both kinases is inviable (Cvrckova *et al.*, 1995), demonstrating that Ste20p is essential in the absence of Cla4p (and vice versa). One interpretation of this relationship is that these two kinases share a function that is essential, though other interpretations are possible. Currently, the

Article published online ahead of print. Mol. Biol. Cell 10.1091/mbc.E02-06-0348. Article and publication date are at [www.molbiolcell.org/cgi/doi/10.1091/mbc.E02-06-0348](http://www.molbiolcell.org/cgi/doi/10.1091/mbc.E02-06-0348).

<sup>□</sup> The online version of this article contains supplemental tabular material. The online version of the article is available at [www.molbiolcell.org](http://www.molbiolcell.org).

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only proposed targets of Ste20p and Cla4p are the two myosin I homologs Myo3p and Myo5p. Sites in Myo3p and Myo5p are phosphorylated *in vitro* by Ste20p and are required for *in vivo* function (Wu *et al.*, 1997). Myo3p and Myo5p are required for actin patch assembly (Evangelista *et al.*, 2000; Geli *et al.*, 2000; Lechler *et al.*, 2000). However, although an activated allele of Myo3p (*MYO3<sup>S357D</sup>*) can rescue the polarity defects of *myo3Δ myo5Δ* mutants, this mutant failed to compensate for the growth defects in *ste20Δ cla4Δ* mutants, indicating that there are other targets of Ste20p and Cla4p (Wu *et al.*, 1997). To identify the target(s) and function(s) regulated by Ste20p, we have carried out a search for mutations that are synthetically lethal in a *cla4Δ* mutant background with the expectation that the genes identified would suggest the nature of the physiological events that have been perturbed.

Herein, we present the results of two independent synthetic lethal mutant screens. One screen was based on random mutagenesis of the genome by using a red/white colony sectoring assay (Kranz and Holm, 1990; Bender and Pringle, 1991). The second screen used a yeast genome-wide deletion set and evaluated the viability of *cla4Δ* paired with 4672 different viable deletion strains (Tong *et al.*, 2001). From the collection of genes defined by these screens, we chose a subset for more detailed investigation. *BNI1* is at the center of this study and encodes a formin homology protein (Zahner *et al.*, 1996) that is required to assemble actin cables (Evangelista *et al.*, 2002; Sagot *et al.*, 2002). These cables seem to guide myosin motors that direct secretion, organelle and mRNA inheritance, and mitotic spindle orientation, thereby establishing cell polarity (Evangelista *et al.*, 2002). Bni1p has an intricate network of interactions involving a number of different groups of proteins. One such group of proteins, Bud6p, Spa2p, and Pea2p, complexes with Bni1p to form the "polarisome," which is involved in apical growth (Sheu *et al.*, 1998; Pruyne and Bretscher, 2000). We show that Bud6p, Spa2p, and Pea2p are essential in a *cla4Δ* mutant (Evangelista *et al.*, 1997; Fujiwara *et al.*, 1998; Sheu *et al.*, 2000). Conversely, other proteins that interact with Bni1p, including Hof1p and Num1p (involved in cytokinesis and nuclear migration, respectively), are not essential in cells lacking Cla4p (Kamei *et al.*, 1998; Heil-Chapdelaine *et al.*, 2000; Vallen *et al.*, 2000; Farkasovsky and Kuntzel, 2001). Finally, we further show that Bni1p is a Ste20p-dependent phosphoprotein, suggesting that Bni1p's function may be directly regulated by Ste20p. Together, these results imply that Ste20p may play an important role in activating the polarisome.

## MATERIALS AND METHODS

### Growth Conditions, Plasmids, and Strains

Yeast and bacterial strains were propagated using standard methods (Sambrook *et al.*, 1989; Rose *et al.*, 1990). YEPD and SD media were prepared as described previously (Rose *et al.*, 1990). Yeast transformations were performed using modifications of the LiOAc method (Chen *et al.*, 1992; Gietz *et al.*, 1995). Bacterial transformations, DNA preparations, and plasmid constructions were performed by standard methods (Sambrook *et al.*, 1989). The plasmids used in this study, YCp*HIS3cla4-75*, pY39tet1HA-*BNI1* (p925), *pcla4-75-td*, and pRS316*ADE8CLA4* have been described elsewhere (Cvrckova *et al.*, 1995; Evangelista *et al.*, 1997; Holly and Blumer, 1999; Mitchell and Sprague, 2001). To ensure that the version *cla4-*

*75-td* we were using was the same as the allele used in Holly and Blumer (1999), we sequenced the *pcla4-75-td* and the *cla4-75-td* alleles rescued from our strains. We found that the *cla4-75-td* alleles rescued from our strains were identical to that of the *pcla4-75-td* from the Blumer laboratory. Strains that were used in this study are listed in Table 1. Gene deletions were constructed by polymerase chain reaction (PCR) (Baudin *et al.*, 1993) by using either the pRS (Sikorski and Hieter, 1989) or pFA6a (Longtine *et al.*, 1998) plasmid series as templates. In all cases, the entire coding region was replaced with the indicated marker, and successful replacement was confirmed by PCR and phenotype when applicable. Single step gene deletion plasmids for *swe1::LEU2*, *spa2::URA3* (p210) and *pea2::URA3* (pNV44) were provided by I. Herskowitz and D. Lew (Booher *et al.*, 1993; Valtz and Herskowitz, 1996). We used *bni1::URA3* (p321), a single step gene deletion plasmid, to delete *BNI1* (Evangelista *et al.*, 1997). 5-Fluoroorotic acid (5-FOA) (Biovector, Oxford, CT) was used to select for uracil auxotrophs. The COOH-terminal deletion mutant *bni1-CTΔ1* lacks the coding sequence for amino acids 1749–1953 of Bni1p (Lee *et al.*, 1999). *bni1-CTΔ1* was created by amplification of the *kanMX6* cassette from pFA6a-*kanMX6* together with sequences immediately flanking base pairs 5247–5859 of *BNI1* by using the forward primer 5'-ATAAAATGAATACAAAAAGCTCAAGCGCAAATCTAGCTGAGGCGCGCCACTTCTAAA-3' and the reverse primer 5'-GTITTTGGTATTACTGTGGCATAATTTTTGGTTAATATGAAATTCGAGCTCGTTTAAAC-3' (the sequences flanking base pairs 5247–5859 of *BNI1* are underlined) (Longtine *et al.*, 1998). The amplified fragment was transformed into strains SY3357, SY3362, SY3380, and SY3764; the transformants were plated on YEPD medium and incubated overnight at 30°C. The lawn of cells was then replica-plated onto YEPD containing 200 μg/ml G418/geneticin (Invitrogen, Carlsbad, CA) to select stable G418-resistant transformants. The successful creation of the deletion strain was confirmed by PCR, and the protein function was verified in a *bni1Δ* strain, in which Bni1p is essential (our unpublished data) (Ozaki-Kuroda *et al.*, 2001).

### Synthetic Lethal Mutant Screens

Two independent methods were used to search for mutations that are lethal in a *cla4Δ* background. Previously, we described the details of the NCS screen by using the colony sectoring assay (Mitchell and Sprague, 2001). Synthetic genetic array analysis (SGA) was also used to identify genes that were essential in a *cla4Δ* background as described in Tong *et al.* (2001). Y2928 (*MATα cla4Δ::natR mfa1Δ::MFA1pr-HIS3 can1Δ ura3Δ leu2Δ his3Δ1 lys2Δ*) was created in two steps. First, *CLA4* was deleted from Y2454 by using PCR-based integration with primers (5'-TTTG-GTGTAATAAAATCGAACA GTGAAACTGAAACATAAAAGAAAT-AGTGCAAAATGGAAACAGCTATG ACCATG-3' and 5'-AGAAAT-ACATAAGATTGTAGTATGTATGATATGCTTATAGAAATAGTTGT-GTGCTGTGTGTAACGACGGCCAGT-3'), which annealed to *URA3* and contained *CLA4* sequences (underlined), to generate Y2851 (*MATα cla4Δ::URA3 mfa1Δ::MFA1pr-HIS3 can1Δ ura3Δ leu2Δ his3Δ1 lys2Δ*). Second, *cla4Δ::URA3* was switched to *cla4Δ::natR* by PCR-based integration with primers (5'-AGTATTCTTAACCAACTGCACAGAACAAA-AACCTGCAGGAAACGAAGATAAATCATGACCCTCTTGACGA-CACGG-3' and 5'-TTGAAGCTCTAATTTGTGAGTTTGTATATCATGCATTTACITATAAATACAGTTTCTAGGGCGAGGGCATGCTCAT-3'), which anneal to *natMX4* DNA (Goldstein *et al.*, 1999) and contain *URA3* sequences (underlined). We performed SGA on *cla4Δ::natR* four times. A total of 100 potential positives were identified and 62 were confirmed by tetrad analysis.

### Isolation of BNI1, BUD6, and Other NCS Genes

Wild-type *NCS8* and *NCS5* were identified as *BNI1* and *BUD6* by complementation of *ncs8-1* (SY3372) and *ncs5-1* (SY3369) mutants, respectively. For *NCS8* isolation, 20,000 library transformants yielded six complementing clones from a yeast genomic library (ATCC no. 77162). An 8.6-kb region shared by all of them was

**Table 1.** Yeast strains used in this study

Strain <sup>a</sup>	Genotype	Source
SY3357	<i>MATa leu2-Δ1 ura3-52 his3-Δ200 trp1-Δ63 ade8Δ ade2-101 mfa2-Δ1::FUS1-lacZ</i>	Mitchell and Sprague (2001)
SY3358	<i>MATα leu2-Δ1 ura3-52 his3-Δ200 lys2-801 trp1-Δ63 ade8Δ ade2-101 mfa2-Δ1::FUS1 lacZ</i>	Mitchell and Sprague (2001)
SY3362	SY3357 except <i>cla4Δ::TRP1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3364	SY3357 except <i>cla4Δ::TRP1 ncs1Δ::LEU2</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3366	SY3357 except <i>cla4Δ::TRP1 ncs2-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3367	SY3357 except <i>cla4Δ::TRP1 ncs3-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3368	SY3357 except <i>cla4Δ::TRP1 ncs4-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3369	SY3357 except <i>cla4Δ::TRP1 ncs5-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3370	SY3357 except <i>cla4Δ::TRP1 ncs6-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3371	SY3357 except <i>cla4Δ::TRP1 ncs7-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3372	SY3357 except <i>cla4Δ::TRP1 ncs8-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3373	SY3357 except <i>cla4Δ::TRP1 ncs10-1</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3380	SY3357 except <i>cla4Δ::TRP1</i> (YCpHIS3 <i>cla4-75</i> )	Mitchell and Sprague (2001)
SY3403	SY3357 except <i>cla4Δ::TRP1 swe1::LEU2 ncs1Δ::HIS3</i> (pRS316ADE8CLA4)	Mitchell and Sprague (2001)
SY3756	SY3357 except <i>cla4Δ::TRP1 bni1Δ::HIS3</i> (pRS316ADE8CLA4)	This study
SY3757	SY3357 except <i>cla4Δ::TRP1 bud6Δ::HIS3</i> (pRS316ADE8CLA4)	This study
SY3758	SY3358 except <i>cla4Δ::TRP1 spa2::ura3</i> (pRS316ADE8CLA4)	This study
SY3759	SY3357 except <i>cla4Δ::TRP1 pea2::ura3</i> (pRS316ADE8CLA4)	This study
SY3760	SY3357 except <i>cla4Δ::TRP1 bni1::URA3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3761	SY3357 except <i>cla4Δ::TRP1 bud6Δ::URA3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3762	SY3357 except <i>cla4Δ::TRP1 spa2::URA3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3763	SY3357 except <i>cla4Δ::TRP1 pea2::URA3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3764	SY3357 except <i>cla4Δ::TRP1 ste20Δ::URA3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3766	SY3357 except <i>cla4Δ::TRP1 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3767	SY3357 except <i>cla4Δ::TRP1 ste20Δ::TRP1 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3768	SY3357 except <i>cla4Δ::TRP1 bni1::ura3 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3769	SY3357 except <i>cla4Δ::TRP1 bud6Δ::his3 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3770	SY3358 except <i>cla4Δ::TRP1 spa2::ura3 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3771	SY3357 except <i>cla4Δ::TRP1 pea2::ura3 swe1::LEU2</i> (pRS316ADE8CLA4)	This study
SY3772	SY3357 except <i>cla4Δ::TRP1 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3773	SY3357 except <i>cla4Δ::TRP1 ste20Δ::URA3 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3774	SY3357 except <i>cla4Δ::TRP1 bni1::ura3 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3775	SY3357 except <i>cla4Δ::TRP1 bud6Δ::ura3 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3776	SY3357 except <i>cla4Δ::TRP1 spa2::ura3 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3777	SY3357 except <i>cla4Δ::TRP1 pea2::ura3 swe1::LEU2</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3778	SY3358 except <i>bni1::HIS3</i>	This study
SY3779	SY3358 except <i>bni1::HIS3</i> (pY39tet1HA-BNI1)	This study
SY3780	SY3358 except <i>ste20Δ::TRP1 bni1::HIS3</i> (pY39tet1HA-BNI1)	This study
SY3781	SY3357 except <i>cla4Δ::TRP1 ste20Δ::TRP1 bni1::ura3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3782	SY3357 except <i>cla4Δ::TRP1 ste20Δ::kanMX6 bud6Δ::ura3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3783	SY3357 except <i>cla4Δ::TRP1 ste20Δ::kanMX6 spa2::ura3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3784	SY3357 except <i>cla4Δ::TRP1 ste20Δ::kanMX6 pea2::ura3</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3785	SY3358 except <i>bni1-CTΔ1::kanMX6</i>	This study
SY3786	SY3357 except <i>cla4Δ::TRP1 bni1-CTΔ1::kanMX6</i> (pRS316ADE8CLA4)	This study
SY3787	SY3357 except <i>cla4Δ::TRP1 ste20Δ::TRP1 bni1-CTΔ1::kanMX6</i> (pRS316ADE8CLA4)	This study
SY3788	SY3357 except <i>cla4Δ::TRP1 bni1-CTΔ1::kanMX6</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3789	SY3357 except <i>cla4Δ::TRP1 ste20Δ::TRP1 bni1-CTΔ1::kanMX6</i> (YCpHIS3 <i>cla4-75</i> )	This study
SY3790	SY3357 except <i>cla4Δ::TRP1 URA3::cla4-75-td</i>	This study
SY3791	SY3357 except <i>cla4Δ::TRP1 bni1::HIS3 URA3::cla4-75-td</i>	This study
SY3792	SY3357 except <i>cla4Δ::TRP1 bud6Δ::ura3 URA3::cla4-75-td</i>	This study
SY3793	SY3357 except <i>cla4Δ::TRP1 spa2::ura3 URA3::cla4-75-td</i>	This study
SY3794	SY3357 except <i>cla4Δ::TRP1 pea2::ura3 URA3::cla4-75-td</i>	This study
SY3795	SY3357 except <i>cla4Δ::TRP1 ste20Δ::TRP1 URA3::cla4-75-td</i>	This study
DY759	<i>cry1 ade2-101(am) his3-11 leu2-3,112 ura3-1</i>	Weiss <i>et al.</i> (2000)
DY2060	DY759 except <i>cla4Δ::LEU2 ste20Δ::KanMX URA3::cla4-75-td</i>	Weiss <i>et al.</i> (2000)
Y2454	<i>MATα mfa1Δ::MFA1pr-HIS3 can1Δ ura3Δ leu2Δ his3Δ1 lys2Δ</i>	This study
Y2851	Y2454 except <i>cla4Δ::URA3</i>	This study
Y2928	Y2851 except <i>cla4Δ::natR</i>	This study

<sup>a</sup> All are derivatives S288C except DY759 and DY2060 (W303 derivatives).

sequenced and found to include *BNI1*. *NCS8* was shown to be *BNI1* by deletion and linkage analysis (see below). For *BUD6*, two clones complementing *ncs5-1* were found among 8000 library transformants. An 8-kb fragment shared by both complementing plasmids was sequenced. Deletion and subcloning analysis identified *BUD6* as the complementing gene. To isolate *NCS2*, strain SY3366, which harbors an *ncs2* mutation, was transformed with a high copy YEp13 based library (ATCC no. 37323), yielding six complementing clones in 6000 transformants. A 2-kb fragment containing two overlapping open reading frames (ORFs) shared by all complementing plasmids was sequenced. Deletion analysis identified *YNL119w/YNL120c* as the complementing ORF(s). For *NCS3* isolation, 7000 library transformants yielded six complementing clones from a high copy YEp13-based library. A 3.6-kb fragment containing three ORFs shared by all complementing plasmids was sequenced. Deletion and subcloning analysis identified *UBA4* as the complementing gene. Because *ncs4* mutants had a strong mating defect, it seemed reasonable that members of this complementation group could contain mutations in *STE20*. Indeed, we found that *STE20* on a plasmid complemented these mutants. In the case of *NCS6*, transformation of SY3370 with yeast genomic library (ATCC no. 77162) yielded two complementing clones from 9000 transformants. A 6.8-kb fragment containing six ORFs shared by all complementing plasmids was sequenced. Deletion and subcloning analysis identified *YGL211w* as the ORF containing the complementing gene. In the case of *NCS10*, 16,000 transformants of a yeast genomic library yielded 16 complementing clones. A 4.6-kb fragment containing three ORFs shared by all complementing plasmids was sequenced. Deletion and subcloning analysis identified *ELP2* as the complementing gene.

Linkage analysis was performed to verify that the cloned genes represented wild-type versions of the mutant alleles. A *HIS3* marker was introduced at the locus of interest in a diploid homozygous for the *cla4* mutation and heterozygous for the *NCS* gene of interest. The strain carried plasmid-borne *CLA4* so that segregation of the *ncs* mutation in tetrads could be scored. After sporulation of the marked strains, the *Ncs*<sup>+</sup> phenotype cosegregated with the *HIS3* marker in at least 22 tetrads.

### Microscopy

Standard microscopic techniques were used, and cells were examined using an Axioplan 2 fluorescence microscope (Carl Zeiss, Thornwood, NY) fitted with an Orca 100 digital camera (Hamamatsu, Bridgewater, NJ). Methods for staining with rhodamine-phalloidin (Molecular Probes, Eugene, OR) to visualize F-actin was performed essentially as described previously (Pringle *et al.*, 1989). All assays were performed in triplicate. Indirect immunofluorescence was performed to visualize the septins by using an  $\alpha$ -Cdc3p antibody (a generous gift from John Pringle) (Roberts *et al.*, 1991). Cells were grown in YEPD at 30°C to 0.7 OD<sub>600</sub>/ml before fixation. Strains containing plasmids were grown first in selective medium, transferred to YEPD, and then grown for 3–4 h at 37°C. Cells were fixed by adding a final concentration of 3.7% formaldehyde to the culture medium for 1 h. The cells were pelleted by centrifugation, resuspended in 4% paraformaldehyde (final concentration) in 50 mM KPO<sub>4</sub>, pH 6.5, and incubated for 18 h at room temperature. The fixed cells were then spheroplasted and permeabilized with 5% SDS for 5 min. Antibody incubations were carried out for 1 h at 22°C. The secondary antibody used was Alexa (A594)-conjugated goat anti-rabbit antibody (Molecular Probes).

### Mating and Invasive Growth Assays

Quantitative mating assays were done using strain 227 as a tester (Sprague, 1991). For invasive growth assays, cells were transformed with YEplac181-*FLO8* to allow the S288c background to manifest the switch from the yeast form to the filamentous form (Gagiano *et al.*, 1999). YEplac181-*FLO8* was a generous gift from Isak S. Pretorius (Stellenbosch University, Stellenbosch, South Africa). The plate

washing assay was performed as previously described (Roberts and Fink, 1994).

### In Vivo Labeling

To label with <sup>32</sup>P<sub>i</sub>, cultures of yeast strains SY3778, SY3779, and SY3780 were pregrown overnight in synthetic medium lacking leucine. Cells were washed once with phosphate-depleted medium (Rubin, 1975) and transferred into phosphate-depleted medium at an OD<sub>600</sub> of 0.2 and grown to an OD<sub>600</sub> of 0.5. For labeling with <sup>32</sup>P<sub>i</sub>, 50 ml of culture was harvested, the pellet was suspended in 15 ml of phosphate-depleted medium, and 1 mCi of <sup>32</sup>PO<sub>4</sub> (ICN Pharmaceuticals, Costa Mesa, CA) was added. After labeling for 45 min, 10 mM sodium azide was added to the cultures and cells were harvested. Cells were spheroplasted as described previously (Graham *et al.*, 1998). Frozen spheroplasts were resuspended in lysis buffer (50 mM Tris, pH 8.0, 1% NP-40, 50 mM NaCl, 1 mM EDTA) containing a mixture of protease inhibitors (1 mM phenylmethanesulfonyl fluoride, 1  $\mu$ g/ml leupeptin, 1  $\mu$ g/ml pepstatin A, 1  $\mu$ g/ml mg/ml aprotinin [all from Sigma-Aldrich, St. Louis, MO] and one tablet of Roche Diagnostics protease inhibitor mixture Complete/25 ml) and phosphatase inhibitors (5 mM sodium pyrophosphate, 0.1 mM sodium metavanadate, 50 mM NaF; all from Sigma-Aldrich). The spheroplasts were incubated in lysis buffer for 15 min at 4°C and unlysed cells were removed by centrifugation at 13,000  $\times$  g for 2 min. A total of 0.5 ml of lysate was precleared with 50  $\mu$ l of protein A-Sepharose beads and bovine serum albumin (1 mg/ml). Bni1p was immunoprecipitated from these extracts with 3  $\mu$ l of rabbit anti-hemagglutinin (HA) antiserum (a kind gift from T. Stevens, Institute of Molecular Biology, University of Oregon) for 30 min at 4°C, after which 50  $\mu$ l of protein A-Sepharose beads was added and incubation continued for another 30 min at 4°C. These pellets were then washed four times with lysis buffer, boiled in 1% SDS before the addition of 9 volumes of lysis buffer, and the immunoprecipitation was repeated. These pellets were washed four times in lysis buffer, resuspended in 30  $\mu$ l of sample buffer, and the entire sample was run on a 6% SDS-PAGE gel, transferred to nitrocellulose, and visualized on a Storm PhosphorImager. Immunoprecipitated Bni1p was also detected by Western blot analysis with an anti-HA monoclonal antibody followed by a horseradish peroxidase-labeled anti-mouse secondary antibody. Blots were developed with ECLplus (Amersham Biosciences, Piscataway, NJ), visualized by chemiluminescence, and quantified by chemifluorescence on a Storm PhosphorImager (Invitrogen, Sunnyvale, CA) with a wavelength of 450 nm.

## RESULTS

### Identification of *BNI1* and *BUD6* by Synthetic Lethal Interactions with a *cla4* Null Mutation

In an effort to identify activators and targets of Ste20p, we screened for mutations that are lethal in combination with a *cla4* null mutation by using two independent methods. The first method used a random mutagenesis of the genome and identified synthetic lethal mutations via a red/white colony sectoring assay (Table 2). Such a screen can be expected to identify two classes of genes: 1) genes that encode upstream activators and downstream effectors of *STE20*, and 2) genes that share a function with *CLA4* that is independent of *STE20*. Previously, we described the isolation of *NCS1* (Needs *CLA4* to Survive), which falls into the second class (Mitchell and Sprague, 2001). *NCS1/RRD1* was unique among *NCS* genes in that *ncs1 $\Delta$*  mutants were not defective for any known *STE20* function. *NCS1* is a phosphotyrosyl phosphatase activator that may share a function with *CLA4* required at the G2/M phase transition (Mitchell and Sprague, 2001).

**Table 2.** NCS mutants

Complementation group	Gene or ORF	Isolates	<i>ste1Δ</i> Overcomes synthetic lethality	Invasive growth	Mating competency
NCS1	<i>RRD1</i>	1	Yes	No defect	WT
NCS2	YNL119w/YNL120c	6	Yes	Defect	WT
NCS3	<i>UBA4</i>	6	Yes	Defect	WT
NCS4	<i>STE20</i>	4	No	Defect	1000 fold lower than wild type
NCS5	<i>BUD6</i>	2	No	Defect	N/D
NCS6	YGL211w	3	N/D	Defect	N/D
NCS7	N/D	2	Yes	Defect	WT
NCS8	<i>BNI1</i>	5	No	Defect	~3-fold lower than wild type
NCS10	<i>ELP2</i>	2	Yes	Defect	WT

N/D, not determined; WT, wild type.

The screen also yielded complementation groups that exhibited some *ste20Δ* phenotypes; in many cases, a defect in haploid invasive growth and in some cases a partial defect in mating as well. As described in MATERIALS AND METHODS, molecular cloning identified the genes corresponding to these complementation groups (Table 2). In this study, we chose to concentrate on *NCS8/BNI1*, both because it is required for efficient mating and because a second complementation group, *NCS5/BUD6*, encodes a protein known to interact with Bni1p (Evangelista *et al.*, 1997). Bni1p (Bud neck involved protein) is a formin homology protein that interacts with a large number of proteins and has many functions attributed to it. These functions include roles in bipolar bud site selection in diploids, cell polarity, cytokinesis, and spindle alignment during nuclear migration and may all stem from its role in actin cable assembly (Zahner *et al.*, 1996; Lee *et al.*, 1999; Miller *et al.*, 1999; Sheu *et al.*, 2000; Vallen *et al.*, 2000; Evangelista *et al.*, 2002). We therefore sought to establish which Bni1p functions were essential in a *cla4Δ* background and whether Bni1p has a specific link to Ste20p. To this end, we asked whether the loss of other proteins that interact with Bni1p or function in the same processes as Bni1p was lethal in a *cla4Δ* background. Some of these new double mutants were constructed by transformation with the appropriate gene disruption followed by a genetic cross. Other double mutants were created in the second synthetic lethal screen, which used a systematic method, known as SGA analysis, to construct double mutants (Tong *et al.*, 2001). This screen used a genome-wide deletion set and evaluated the viability of *cla4Δ* paired with deletion of 4672 open reading frames. The results of this screen can be viewed in Figure 1 and in Supplementary Table 1. The two approaches, random mutagenesis coupled with subsequent directed double mutant construction and the use of the genome-wide deletion set, were complementary. Each identified some unique genes and in cases where the same gene was investigated, the two approaches gave congruent results. The SGA analysis provides a global view of the proteins that become essential in the absence of Cla4p. Such proteins include ones involved in apical growth, bud emergence, cytokinesis, mitosis, and cell wall maintenance. However, although the SGA method enabled us to perform a more complete synthetic lethal analysis, three of the nine

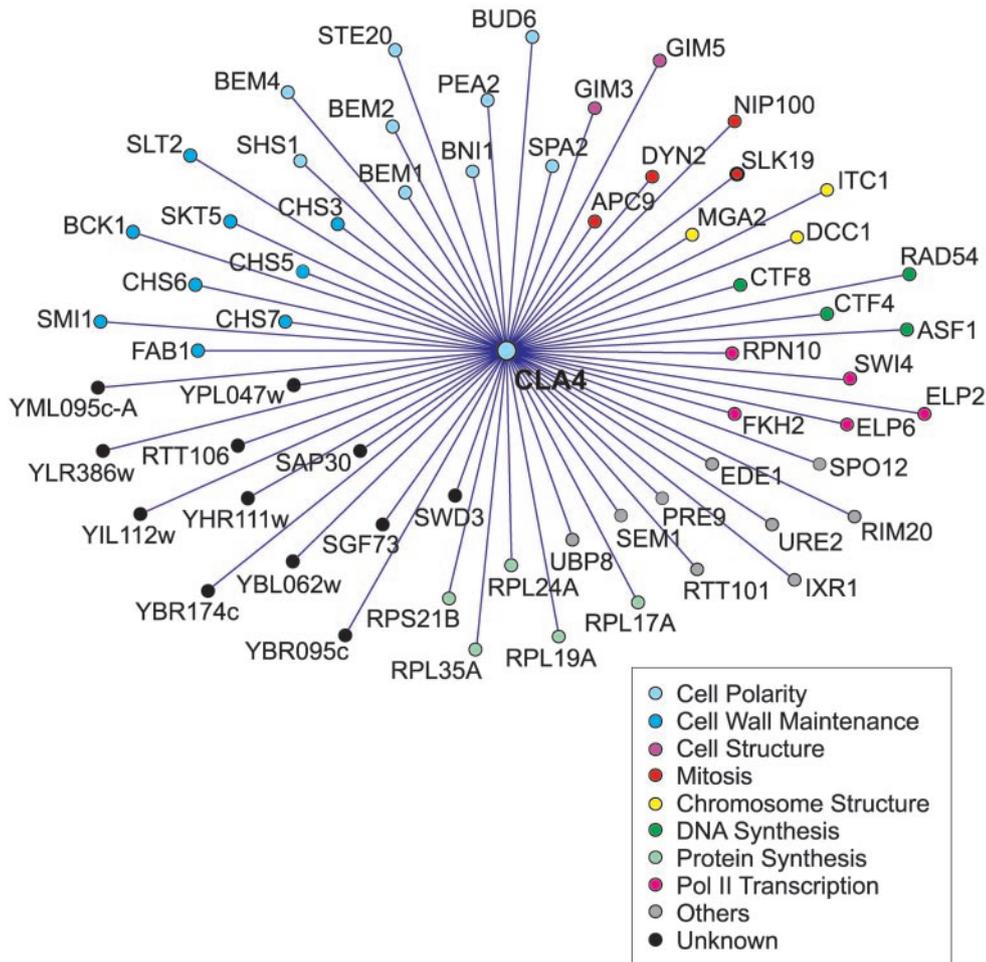
complementation groups identified in the random mutagenesis were not identified by the SGA method.

### ***Polarisome Components Are Essential in a *cla4Δ* Background***

As noted above, loss of either *BNI1* or *BUD6* is lethal in a *cla4Δ* strain. Each gene is involved in cell polarity establishment and in bipolar budding (Zahner *et al.*, 1996; Evangelista *et al.*, 1997; Sheu *et al.*, 2000). Moreover, Bni1p and Bud6p have been shown to interact by two-hybrid analysis (Evangelista *et al.*, 1997). We also found that loss of Spa2p, another protein that interacts with Bni1p (Fujiwara *et al.*, 1998), is essential in a *cla4Δ* mutant background. These three proteins, together with a fourth protein Pea2p, form a 12S complex termed the polarisome that has been suggested to promote polarized morphogenesis (Sheu *et al.*, 1998; Pruyne and Bretscher, 2000). We found that Pea2p is likewise essential in a *cla4Δ* strain.

To examine the terminal phenotype of *cla4Δ* mutants lacking polarisome function, we used a plasmid-borne thermo-sensitive allele of *CLA4* (YCpHIS3*cla4-75*). A striking phenotype of *ste20Δ cla4Δ* YCpHIS3*cla4-75* mutants is the mislocalization of the septin ring (Cvrckova *et al.*, 1995). We therefore examined septin localization in *bni1Δ cla4Δ* YCpHIS3*cla4-75* and in other polarisome *cla4Δ* double mutants. The septin phenotype of *bni1Δ cla4Δ* YCpHIS3*cla4-75* mutants at the restrictive temperature resembled that of *ste20Δ cla4Δ* YCpHIS3*cla4-75* mutants. The septin ring was formed at the proper time and location. However, as the bud began to grow, the septin ring frequently localized at the tip of the misshapened bud rather than remaining at the bud neck (Figure 2). These results imply that new growth is on the mother side of the neck rather than the bud side. Similar phenotypes were observed with *bud6Δ cla4Δ* YCpHIS3*cla4-75*, *spa2Δ cla4Δ* YCpHIS3*cla4-75*, and *pea2Δ cla4Δ* YCpHIS3*cla4-75* mutants (Figure 2). Other aspects of the polarisome *cla4* double mutants will be discussed below.

To corroborate the results observed using strains harboring YCpHIS3*cla4-75*, we also used strains expressing an integrated *cla4-75-ts* degon construct (*cla4-75-td*). This version of Cla4p is reported to be degraded rapidly following a shift to the restrictive temperature (Holly and Blumer, 1999);



**Figure 1.** Genetic interaction network of the synthetic lethal interactions identified by the SGA analysis. The genes that are essential or required for normal growth rate in a *cla4Δ* are represented as nodes. Each node is color coordinated according to the functional classification of the gene according to YPD (Hodges *et al.*, 1999; Costanzo *et al.*, 2001). For genes that have multiple roles assigned to them, we chose the function that we considered most probable based on a review of published abstracts for the gene of interest.

note, however, that wild-type Cla4p and Cla4-75p are also degraded rapidly at 37°C (Figure 3B). The phenotype of *bni1Δ cla4Δ cla4-75-td* mutants at the restrictive temperature recapitulated that of cells carrying YCp*HIS3cla4-75*: the septin ring was mislocalized to the tip of the bud (Figure 3A). Together, these results indicate that the polarisome is essential in the absence of Cla4p and further suggest that Ste20p may activate the polarisome.

### *Bni1p Is Phosphorylated In Vivo and Phosphorylation Is Partially Dependent on STE20*

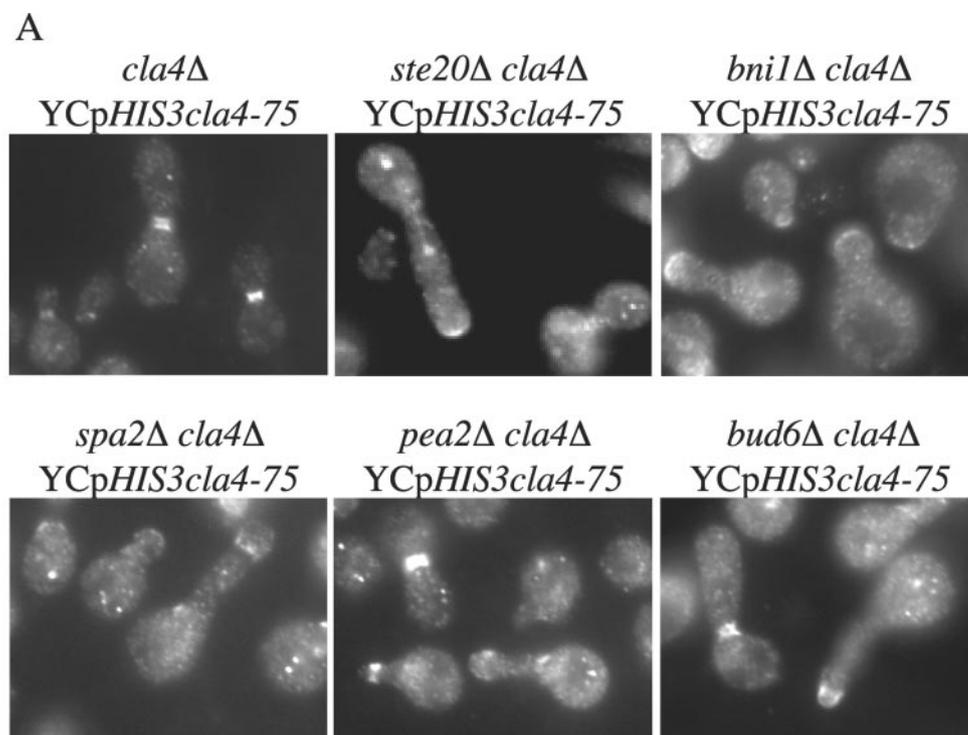
Because Bni1p and Ste20p colocalize to the bud tip in a Cdc42p-dependent manner (Peter *et al.*, 1996; Leberer *et al.*, 1997; Ozaki-Kuroda *et al.*, 2001) and show similar genetic interactions with *CLA4*, it seemed plausible that Bni1p and Ste20p might physically interact and that this interaction would be necessary for the essential activity that Bni1p has in the absence of Cla4p. To investigate this possibility, we created two-hybrid constructs of full-length, N-terminal, and C-terminal fusions of Ste20p and Bni1p but were unable to detect an interaction. We also failed to detect an interaction using coimmunoprecipitation under a variety of assay conditions. Moreover, the

proper localization of Bni1p or Ste20p to the bud tip did not require the presence of the other protein (our unpublished data).

Although we were unable to detect a physical interaction between Ste20p and Bni1p by using the methods described, we considered the possibility that the interaction is transient. In particular, because Ste20p is a protein kinase we asked whether Bni1p is a Ste20p-dependent phosphoprotein. A culture of cells expressing HA-tagged Bni1p was labeled with  $^{32}\text{P}_i$ , Bni1p was immunoprecipitated from the labeled extracts with the HA antibody, and radiolabeled proteins in the immune complexes were visualized by a Phosphor-Imager and subsequent immunoblot analysis. Bni1p is indeed a phosphoprotein (Figure 4). Moreover, in cells lacking Ste20p, the amount of phosphorylated Bni1p was twofold less than that found in wild-type cells, suggesting that a substantial portion of Bni1p phosphorylation is dependent on Ste20p *in vivo* (Figure 4).

### *Septin Ring Mislocalization Is Not the Cause of Lethality in a *bni1Δ cla4Δ* Cells*

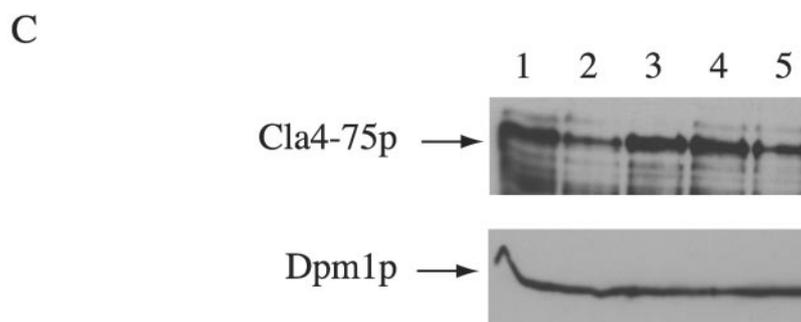
Cells lacking *CLA4* exhibit a bud morphology that suggests a defect in the apical-to-isotrophic bud transition that occurs



**Figure 2.** Morphological phenotypes of *bni1Δ cla4Δ*, *spa2Δ cla4Δ*, *pea2Δ cla4Δ*, and *bud6Δ cla4Δ* carrying YCpHIS3*cla4-75*. (A) Exponential cultures of haploid strains SY3380 (*cla4Δ*), SY3760 (*bni1Δ cla4Δ*), SY3761 (*bud6Δ cla4Δ*), SY3762 (*spa2Δ cla4Δ*), SY3763 (*pea2Δ cla4Δ*), and SY3764 (*ste20Δ cla4Δ*) carrying YCpHIS3*cla4-75* were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for Cdc3p. (B) Quantitation of Cdc3p mislocalization. For each strain, 250 cells were counted in three independent experiments. (C) Immunoblot analysis by using anti-Cla4p antibodies. Lysates of haploid strains SY3357 (WT; lane 1), SY3380 (*cla4Δ*; lane 2), SY3764 (*ste20Δ cla4Δ*; lane 3), SY3760 (*bni1Δ cla4Δ*; lane 4), and SY3761 (*bud6Δ cla4Δ*; lane 5) carrying YCpHIS3*cla4-75* were analyzed by SDS-PAGE and immunoblot analysis by using affinity purified polyclonal anti-Cla4p antibodies (provided by D. Kellogg, Department of Molecular, Cellular, and Developmental Biology, University of California, Santa Cruz) and monoclonal antibodies to Dpm1p (to confirm equal protein loading). The band corresponding to wild-type Cla4p and Cla4-75p are indicated. All strains expressed similar amounts of Cla4-75p.

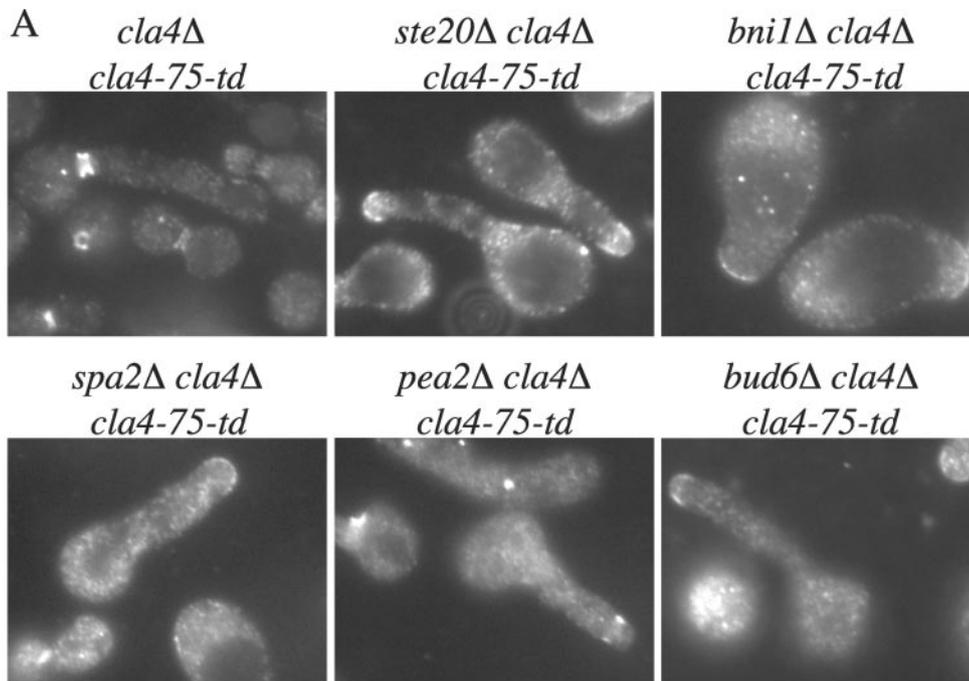
**B**

Cdc3p mislocalization	
genotype	%mislocalization
<i>cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	2
<i>ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	46
<i>bni1Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	66
<i>spa2Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	49
<i>pea2Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	48
<i>bud6Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	37

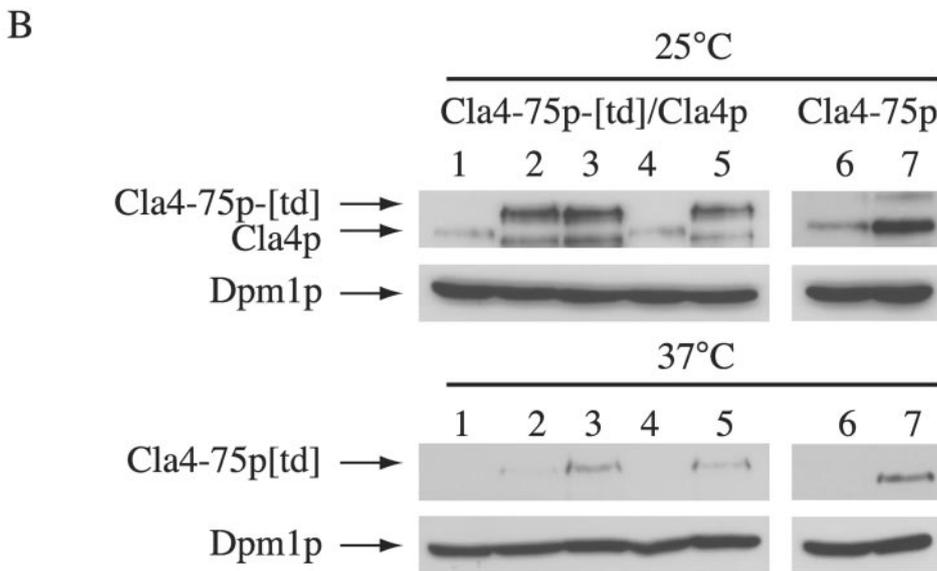


in the G2 phase of the cell cycle. This phenotype is reminiscent of the phenotype conferred by misregulation of Cdc28p kinase activity (Lew and Reed, 1995a,b; McMillan *et al.*, 1998;

Richman *et al.*, 1999). Indeed, we showed previously that deletion of *SWE1*, which encodes a protein kinase thought to be part of a morphogenetic checkpoint that negatively reg-

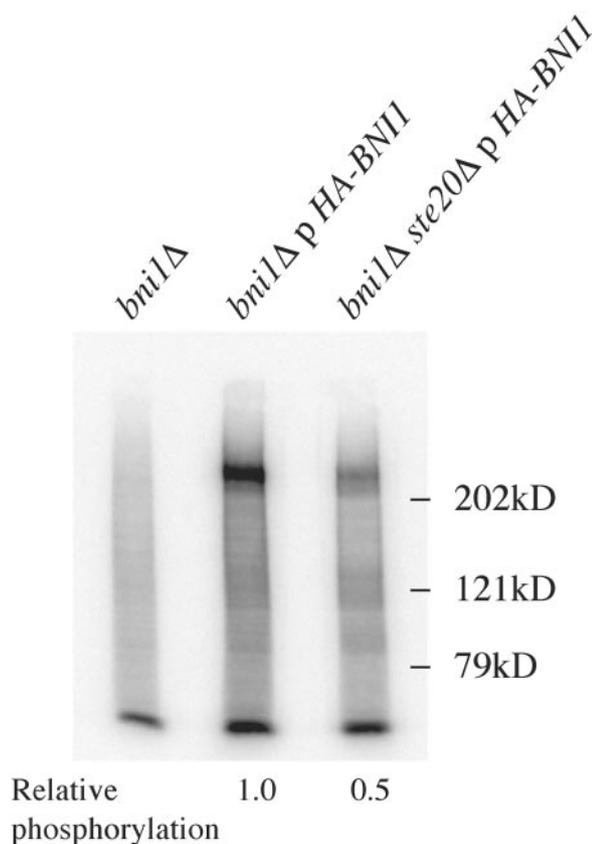


**Figure 3.** Morphological phenotypes of *bni1Δ cla4Δ cla4-75-td*, *spa2Δ cla4Δ cla4-75-td*, *pea2Δ cla4Δ cla4-75-td*, and *bud6Δ cla4Δ cla4-75-td*. (A) Exponential cultures of haploid strains SY3790 (*cla4Δ cla4-75-td*), SY3791 (*bni1Δ cla4Δ cla4-75-td*), SY3792 (*bud6Δ cla4Δ cla4-75-td*), SY3793 (*spa2Δ cla4Δ cla4-75-td*), SY3794 (*pea2Δ cla4Δ cla4-75-td*), and SY3795 (*ste20Δ cla4Δ cla4-75-td*) were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for Cdc3p. (B) Immunoblot analysis using anti-Cla4p antibodies. Lysates of haploid strains SY3357 (WT; lane 1), SY3790 (*cla4Δ cla4-75-td*; lane 2), SY3795 (*ste20Δ cla4Δ cla4-75-td*; lane 3), DY759 (WT; lane 4) and DY2060 (*ste20Δ cla4Δ cla4-75-td*; lane 5), SY3380 (*cla4Δ YCpHIS3cla4-75*; lane 6), and SY3764 (*ste20Δ cla4Δ YCpHIS3cla4-75*; lane 7), were grown at 25°C in YEPD, shifted to 37°C for 1 h, and analyzed by SDS-PAGE and immunoblot analysis by using affinity purified polyclonal anti-Cla4p antibodies (provided by D. Kellogg) and monoclonal antibodies to Dpm1p (to confirm equal protein loading). The position of wild-type Cla4p, Cla4-75p, and Cla4-75p[td] are indicated. In lanes 2, 3 and 5, a degradation product of Cla4-75p[td] is visible at 25°C. Lanes 6 and 7 are a darker exposure than lanes 1–5. As previously reported, at 37°C, there is residual Cla4-75p[td] protein present (Holly and Blumer, 1999; Weiss *et al.*, 2000).



ulates Clb1,2p-Cdc28p activity, restores normal bud morphology in *cla4Δ* mutants (Longtine *et al.*, 2000; Weiss *et al.*, 2000; Mitchell and Sprague, 2001). By bypassing this morphogenetic checkpoint with *swe1Δ*, we were able to class the NCS genes into two groups, an NCS1-related group and an STE20-related group. Loss of *SWE1* restores normal bud morphology and overcomes the synthetic lethality of *ncs1Δ cla4Δ YCpHIS3cla4-75* cells. In the case of *ste20Δ cla4Δ YCpHIS3cla4-75* cells, however, loss of *SWE1* restores the localization of the septin ring to the mother bud junction but does not restore viability (Mitchell and Sprague, 2001). To test whether *BNI1* is in the NCS1 group or the STE20 group,

we deleted *SWE1* in *bni1Δ cla4Δ YCpHIS3cla4-75* cells. The loss of *SWE1* from *bni1Δ cla4Δ YCpHIS3cla4-75* yielded a phenotype similar to that of *swe1Δ ste20Δ cla4Δ YCpHIS3cla4-75* cells (Figure 5), suggesting that septin ring mislocalization is not the cause of lethality. Furthermore, this result suggests that *BNI1* may facilitate or orchestrate some STE20 functions. Similar results were obtained when *SWE1* was deleted from *pea2Δ cla4Δ*, *spa2Δ cla4Δ*, and *bud6Δ cla4Δ* cells carrying *YCpHIS3cla4-75* (Figure 5). Moreover, these polarisome genes constitute a group distinct from other NCS genes based on their genetic interactions with *SWE1* (Table 2).



**Figure 4.** Bni1p is a phosphoprotein. Cultures of SY3779 (*bni1Δ*) and SY3780 (*bni1Δ ste20Δ*) carrying pY39tet1HA-BNI1 and SY3778 (*bni1Δ*) were incubated with 1 mCi of  $^{32}\text{PO}_4$ . Rabbit anti-HA antibody was used for immunoprecipitation and an anti-HA monoclonal antibody was used for immunoblot analysis. The relative amount of phospho-Bni1p, normalized to total Bni1p, is indicated below each lane.

#### Not All Proteins Involved in Bipolar Budding Pattern Are Essential in *cla4Δ* Cells

Recent studies have shown that some mutants defective for bipolar budding pattern selection also show a defect in apical growth and that lengthening the apical growth phase enhances the accuracy of bud site selection (Sheu *et al.*, 2000). Ste20p, Bni1p, Pea2p, Spa2p, and Bud6p have been implicated in apical growth and also have roles in bipolar bud site selection (Snyder, 1989; Valtz and Herskowitz, 1996; Zahner *et al.*, 1996; Evangelista *et al.*, 1997; Sheu *et al.*, 2000). To ascertain whether there is a specific connection between their roles in bipolar budding and the essential function they have in the absence of Cla4p, we looked for genetic interactions between *CLA4* and other components of the bipolar bud site machinery. Loss of *BUD8* was not synthetically lethal with *cla4Δ* (Figure 1 and Supplementary Table 1). In addition, the SGA screen showed that the loss of other genes that affect the budding pattern in diploids, such as *BUD9*, *BUD14*, *BUD16-32* was not synthetically lethal with *cla4Δ*. These results suggest the roles of Bni1p, Spa2p, Pea2p, and Bud6p in bipolar budding pattern are not essential in the absence of Cla4p.

#### Roles of Bni1p in Spindle Alignment during Nuclear Migration and Cytokinesis Are Not Essential in *cla4Δ* Cells

In addition to its roles in bipolar budding pattern and apical growth, Bni1p has a role in cytokinesis. Other proteins important for cytokinesis that function with Bni1p are Myo1p, Hof1p, and Bnr1p, a formin homology protein related to Bni1p (Bi *et al.*, 2000). Based on synthetic lethal interactions, *BNI1* and *MYO1* are believed to be in one functional pathway, whereas *HOF1* and *BNR1* are in another. Loss of *BNI1* is synthetically lethal with *bnr1Δ* and with *hof1Δ*. However, none of these genes (except *BNI1*) is essential in a *cla4Δ* background (our unpublished data). Based on these results, the role of Bni1p in actomyosin contraction during cytokinesis is not essential in the absence of Cla4p.

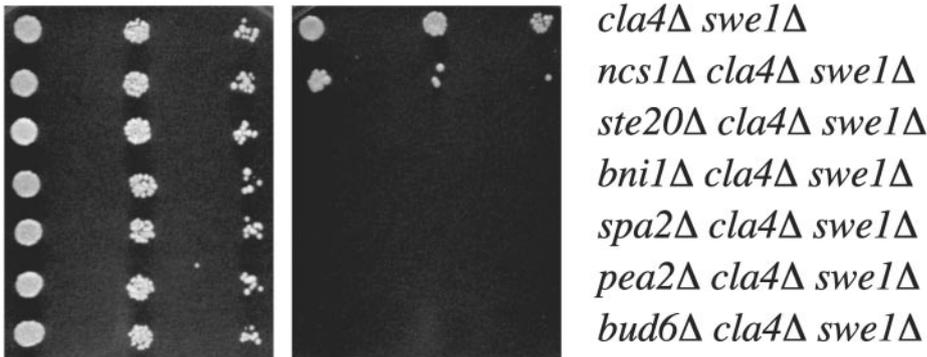
Bni1p has also been shown to play a role in positioning the mitotic spindle during nuclear migration (Lee *et al.*, 1999; Miller *et al.*, 1999). We investigated whether the genetic interactions between *BNI1* and *CLA4* are related to the role of *BNI1* in nuclear migration by looking for genetic interactions with *KIP3*. Kip3p is a kinesin-related protein hypothesized to function with Bni1p to organize and position the mitotic spindle. Loss of *KIP3* was not synthetically lethal with *cla4Δ* (our unpublished data). Furthermore, the loss of *NUM1*, which encodes a protein that controls interaction of bud-neck cytoskeleton with the nucleus in G2 and also interacts with Bni1p, was not synthetically lethal with *cla4Δ* (our unpublished data).

#### Bni1p Has Roles Distinct from Spa2p, Pea2p, Bud6p, and Ste20p in *cla4Δ* Cells

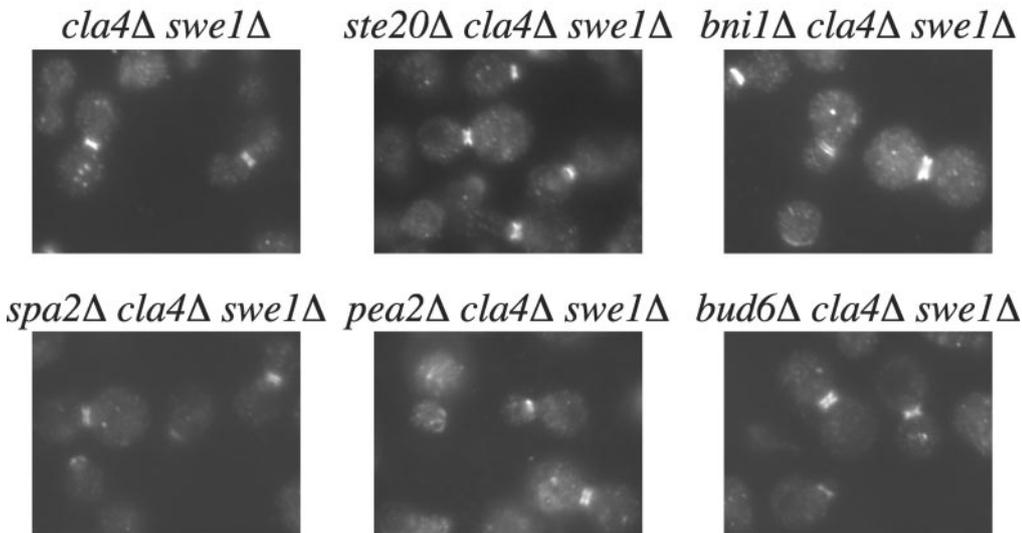
To gain more insight into the role of Bni1p in *cla4Δ* cells, we examined the terminal phenotype of *bni1Δ cla4Δ* YCpHIS3*cla4-75* mutants in more detail. With respect to the septin ring localization, the terminal phenotype of *bni1Δ cla4Δ* YCpHIS3*cla4-75* cells was similar to *ste20Δ cla4Δ* YCpHIS3*cla4-75* cells, but with respect to other phenotypes, the two phenotypes were distinct. *bni1Δ cla4Δ* YCpHIS3*cla4-75* cells have both wider bud necks and defects in actin localization compared with *ste20Δ cla4Δ* YCpHIS3*cla4-75* at the restrictive temperature (Figure 6). In particular, *bni1Δ cla4Δ* YCpHIS3*cla4-75* mutants had no visible actin cables and only 6% of the cells had organized patches of actin at the tip of the bud. In contrast, *cla4Δ* single mutants had no observable defects in actin polarization and only 38% of *bni1Δ* single mutant cells had defects in actin polarization (Figure 6). Thus, it seems that Bni1p is more critical for actin organization in the absence of Cla4p than in wild-type cells, suggesting that Cla4p may also participate in actin organization but that its role in this process is functionally redundant with that of Bni1p.

As previously reported, actin polarization in *ste20Δ cla4Δ* YCpHIS3*cla4-75* cells was indistinguishable from that in *cla4Δ* mutants, with actin cables traversing from mother to bud and actin patches localized toward the bud tip (Cvrckova *et al.*, 1995). Likewise, *spa2Δ cla4Δ* and *pea2Δ cla4Δ* mutants carrying the YCpHIS3*cla4-75* construct did not seem to have defects in actin organization compared with wild-type cells or *ste20Δ cla4Δ* YCpHIS3*cla4-75* mutants (Figure 6). *bud6Δ cla4Δ* YCpHIS3*cla4-75* mutants had some noticeable actin defects with fewer actin cables and polarized

A



B

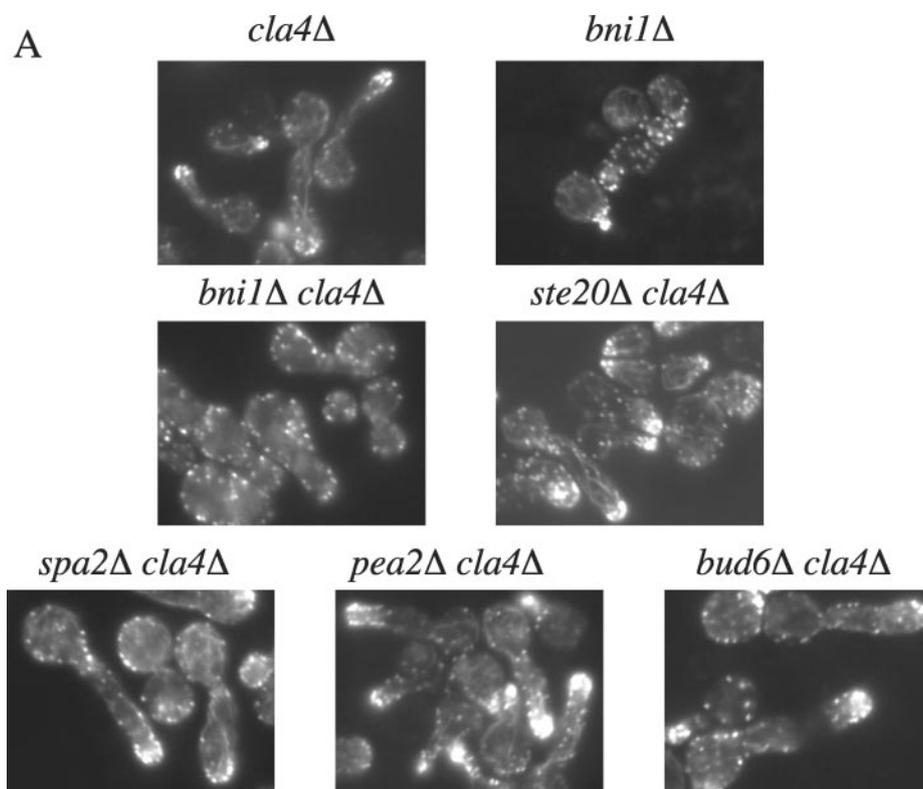


C

Cdc3p mislocalization

genotype	%mislocalization
<i>cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	2
<i>ste20Δ cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	5
<i>bni1Δ cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	13
<i>spa2Δ cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	7
<i>pea2Δ cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	6
<i>bud6Δ cla4Δ swe1Δ</i> (YCpHIS3 <i>cla4-75</i> )	5

**Figure 5.** Loss of *SWE1* in *bni1Δ cla4Δ* restores septin ring localization to the mother-bud junction. (A) Strains SY3766 (*cla4Δ swe1Δ*), SY3403 (*ncs1Δ cla4Δ swe1Δ*), SY3767 (*ste20Δ cla4Δ swe1Δ*), SY3768 (*bni1Δ cla4Δ swe1Δ*), SY3769 (*bud6Δ cla4Δ swe1Δ*), SY3770 (*spa2Δ cla4Δ swe1Δ*), and SY3771 (*pea2Δ cla4Δ swe1Δ*) carrying pRS316*ADE8CLA4* were grown to midlog in YEPD at 30°C. A serial dilution (1/10) was performed starting with 10,000 cells. Cells were spotted onto either YEPD (left) or 5-FOA (right) and grown 3 d at 30°C. (B) Exponential cultures of haploid strains SY3772 (*cla4Δ swe1Δ*), SY3773 (*ste20Δ cla4Δ swe1Δ*), SY3774 (*bni1Δ cla4Δ swe1Δ*), SY3775 (*bud6Δ cla4Δ swe1Δ*), SY3776 (*spa2Δ cla4Δ swe1Δ*), and SY3777 (*pea2Δ cla4Δ swe1Δ*) carrying YCpHIS3*cla4-75* were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for Cdc3p. (C) Quantitation of Cdc3p mislocalization. For each strain 250 cells were counted in three independent experiments.

**B**

## Actin polarization

genotype	%polarization
<i>cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	87
<i>bni1Δ</i> (YCpHIS3 <i>cla4-75</i> )	62
<i>ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	87
<i>bni1Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	6
<i>spa2Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	86
<i>pea2Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	89
<i>bud6Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	67

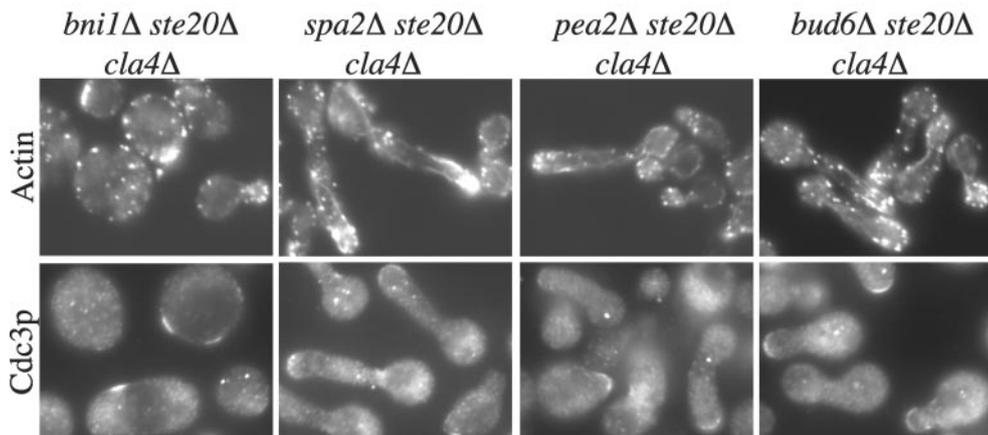
**Figure 6.** *bni1Δ cla4Δ* mutants have a severe actin polarization defect. (A) Strains SY3380 (*cla4Δ*), SY3778 (*bni1Δ*), SY3764 (*ste20Δ cla4Δ*), SY3760 (*bni1Δ cla4Δ*), SY3761 (*bud6Δ cla4Δ*), SY3762 (*spa2Δ cla4Δ*), and SY3763 (*pea2Δ cla4Δ*) carrying YCpHIS3*cla4-75* were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for F-actin (by using rhodamine-phalloidin). (B) Quantitation of actin polarization. For each strain, 250 cells were counted in three independent experiments.

actin patches, yet the defects were not as severe as those of *bni1Δ cla4Δ* YCpHIS3*cla4-75* mutants (Figure 6). Thus, deletion of polarisome genes in *cla4Δ* cells leads to two broad phenotypic classes, one that includes *spa2Δ*, *pea2Δ*, and *bud6Δ*, with a terminal phenotype resembling that associated with *ste20Δ*, and another class whose sole member is *bni1Δ*, associated with a more severe terminal phenotype.

To further test for a functional relationship between Ste20p and the polarisome components, we examined the terminal phenotype of triple mutants containing *spa2Δ*, *pea2Δ*, *bud6Δ*, or *bni1Δ* in combination with *ste20Δ cla4Δ*

YCpHIS3*cla4-75*. The terminal phenotypes of *spa2Δ*, *pea2Δ*, and *bud6Δ* triple mutants were similar to that of the *ste20Δ cla4Δ* YCpHIS3*cla4-75* double mutant (Figure 7). Moreover, no synthetic growth defects were observed in strains where the polarisome genes were deleted in combination with *ste20Δ*. In contrast, the terminal morphology of the *bni1Δ* triple mutant was more severe than that associated with either the *bni1Δ cla4Δ* YCpHIS3*cla4-75* or the *ste20Δ cla4Δ* YCpHIS3*cla4-75* double mutants. The *bni1Δ ste20Δ cla4Δ* YCpHIS3*cla4-75* cells were large and unbudded with no visible actin cables and mostly unpolarized patches of actin

A



B

Actin polarization	
genotype	%polarization
<i>bni1Δ ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	24
<i>spa2Δ ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	71
<i>pea2Δ ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	82
<i>bud6Δ ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	70

(Figure 7). In these unbudded cells, components of the septin ring still localized to the presumptive bud site, but in 2% of cells, Cdc3p (a septin component) localized to more than one site on the cell (Figure 7). In contrast, the actin defects in *ste20Δ bni1Δ* mutants were no more severe than that of *bni1Δ* single mutant cells (62 vs. 59% polarized actin, respectively). Together, these results suggest that Ste20p functions within a Spa2p/Pea2p/Bud6p pathway, but that is also participates in Bni1p-independent functions.

#### Disruption of Bni1p-Bud6p Interaction Leads to Synthetic Lethality with *cla4Δ*

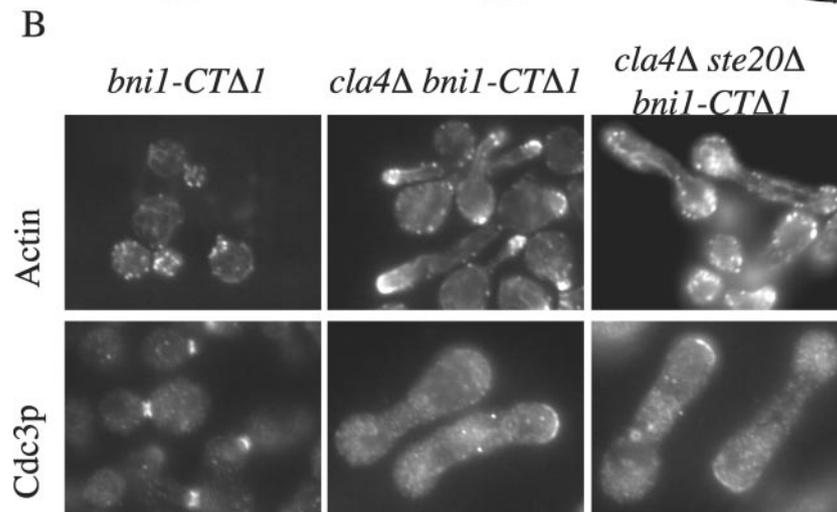
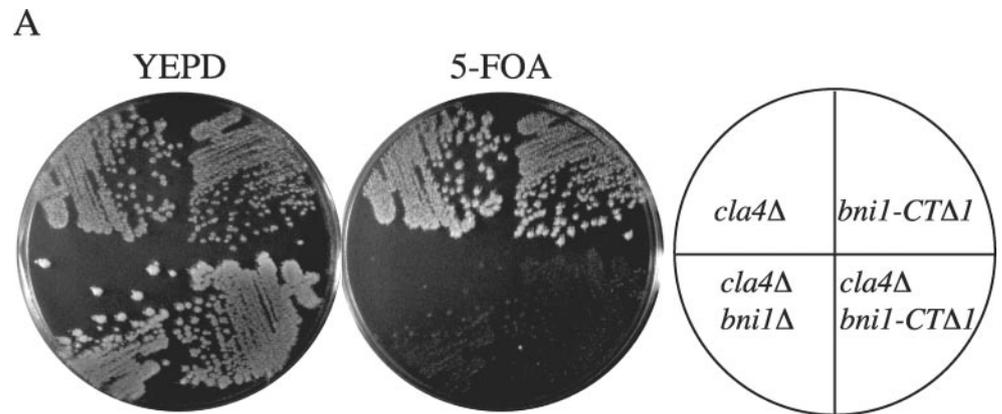
The finding that Bni1p and other polarisome components are essential in the absence of Cla4p, leads to the natural inference the disruption of the interactions between these proteins would lead to synthetic lethality with *cla4Δ*. To explore this possibility, we examined the phenotype conferred by a version of Bni1p that is lacking the Bud6p binding region but still competent to carry out other Bni1p functions. Amino acid residues 1749–1953 of Bni1p contain the Bud6p binding domain (Evangelista *et al.*, 1997). Accordingly, we constructed an allele of *BNI1* that encodes a version of Bni1p truncated at the C terminus (*bni1-CTΔ1*) (Lee *et al.*, 1999). *bni1-CTΔ1* retains some Bni1p functions as it is able to complement the synthetic lethality of a *bni1Δ bnr1Δ* (Ozaki-Kuroda *et al.*, 2001). In contrast, *bni1-CTΔ1* was lethal in a *cla4Δ* background (Figure 7A). Moreover, the terminal phenotype of *bni1-CTΔ1 cla4Δ* YCpHIS3*cla4-75* was

**Figure 7.** Terminal phenotype of *bni1Δ ste20Δ cla4Δ*. Exponential cultures of SY3781 (*bni1Δ cla4Δ ste20Δ*), SY3782 (*bud6Δ cla4Δ ste20Δ*), SY3783 (*spa2Δ cla4Δ ste20Δ*), and SY3884 (*pea2Δ cla4Δ ste20Δ*) carrying YCpHIS3*cla4-75* were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for F-actin (by using rhodamine-phalloidin) or for Cdc3p. (B) Quantitation of actin polarization. For each strain, 250 cells were counted in three independent experiments.

similar to the *ste20Δ cla4Δ* YCpHIS3*cla4-75* terminal phenotype but not the *bni1Δ cla4Δ* YCpHIS3*cla4-75* phenotype (Figure 8B). The *bni1-CTΔ1 cla4Δ* YCpHIS3*cla4-75* cells have polarized actin, whereas *bni1Δ cla4Δ* YCpHIS3*cla4-75* cells have no visible cables and unpolarized patches of actin. In addition, the *ste20Δ cla4Δ bni1-CTΔ1* YCpHIS3*cla4-75* triple mutant phenotype resembled the *ste20Δ cla4Δ* YCpHIS3*cla4-75* and the *bni1-CTΔ1 cla4Δ* YCpHIS3*cla4-75* double mutant phenotypes (Figure 8B). The cells were elongated, with mostly polarized actin, but the septin ring was mislocalized. These results support the idea that the Bni1p C terminus, which interacts with Bud6p, carries out one function that is essential in the absence of Cla4p, whereas the remainder of Bni1p is critical for actin organization.

## DISCUSSION

The related protein kinases Ste20p and Cla4p have unique activities, an inference made from the distinct phenotypes of strains lacking an individual kinase. However, a strain lacking both kinases is inviable, implying that there is a physiological connection between their activities. One possibility is that Cla4p and Ste20p share an essential activity. A less constrained interpretation is simply that Ste20p carries out a function that is essential in cells lacking Cla4p cells (and vice versa). In an effort to shed light on Ste20p function, we carried out two independent screens for mutations that are lethal in a *cla4Δ* mutant background. This effort identified a



**Figure 8.** *bni1-CTΔ1 cla4Δ* mutants have the same phenotype as *ste20Δ cla4Δ*. (A) *bni1-CTΔ1* is synthetically lethal with *cla4Δ*. Strains SY3362 (*cla4Δ*), SY3756 (*bni1Δ cla4Δ*), SY3785 (*bni1-CTΔ1*), and SY3786 (*bni1-CTΔ1 cla4Δ*) harboring pRS316ADE8CLA4 were streaked on YEPD plates and incubated at 30°C for 2 d. Plates were then replica-plated to 5-FOA and incubated at 30°C for 3 d. (B) Exponential cultures of SY3785 (*bni1-CTΔ1*), SY3788 (*bni1-CTΔ1 cla4Δ*) and SY3789 (*bni1-CTΔ1 ste20Δ cla4Δ*) carrying YCpHIS3*cla4-75* were grown at 25°C in YEPD, shifted to 37°C for 4 h, fixed, and stained for F-actin (by using rhodamine-phalloidin) or for Cdc3p. (C) Quantitation of actin polarization. For each strain 250 cells were counted in three independent experiments.

**C**

Actin polarization	
genotype	%polarization
<i>bni1-CTΔ1</i> (YCpHIS3 <i>cla4-75</i> )	89
<i>bni1-CTΔ1 cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	81
<i>bni1-CTΔ1 ste20Δ cla4Δ</i> (YCpHIS3 <i>cla4-75</i> )	77

surprisingly large number of genes. Herein, we focus on a group of genes whose products are known to interact. Each of the proteins that form this group, Bni1p, Bud6p, Spa2p, and Pea2p, has been implicated in several different facets of cell biology. For example, Bni1p, a formin homology protein, has been implicated in actin polarization, cytokinesis, nuclear migration, and apical growth. Similarly, Bud6p, Spa2p, and Pea2p are involved in budding pattern determination and apical growth. The common denominator among the functions attributed to this set of proteins is involvement in apical growth, and we therefore suggest that proper regulation of this growth is the essential in the absence of Cla4p. We further suggest that the connection between these proteins and the Ste20p/Cla4p

essential function is likely to be direct because Bni1p is a Ste20p-dependent phosphoprotein.

#### *Bni1p* Has Roles Distinct from That of *Spa2p*, *Pea2p*, *Bud6p*, or *Ste20p* in *cla4Δ* Cells

The lethality of *bni1Δ cla4Δ* strains may have more than one cause. As summarized above, we think one aspect of Bni1p function that is required in a *cla4Δ* mutant background is its participation in apical growth. This interpretation is supported by the finding that deletion of the Bni1p C-terminal 200 amino acids, the region of Bni1p that interacts with Bud6p, is lethal in a *cla4Δ* mutant background. Indeed, the terminal phenotype of *bni1-CTΔ1 cla4Δ* YCpHIS3*cla4-75* is

similar to the *bud6Δ cla4Δ* YCpHIS3*cla4-75* terminal phenotype. The C-terminal region is not only the Bud6p interaction domain on Bni1p, but it is also believed to be an autoinhibitory domain. In the case of other formin homology proteins, this autoinhibitory domain has been shown to interact with the Cdc42p binding domain on the same molecule (Alberts, 2001). Perhaps binding of activated Cdc42p releases the autoinhibitory domain and enables Bni1p to interact with Bud6p and the 12S complex. It will be interesting to determine whether the Ste20p-dependent phosphorylation of Bni1p influences interaction of it with Bud6p or other proteins.

Our results suggest that Bni1p has at least one other function that is important in a *cla4Δ* mutant background. This possibility emerges from the observation that *bni1Δ cla4Δ* YCpHIS3*cla4-75* double mutants have additional phenotypes beyond those seen for the *ste20Δ cla4Δ*, *bni1-CTΔ1 cla4Δ* YCpHIS3*cla4-75*, and *bud6Δ cla4Δ* YCpHIS3*cla4-75* mutant strains. What is this additional important function? In addition to its role in apical growth, Bni1p also has roles in bud site selection, nuclear migration, cytokinesis, and actin polarization. Genetic tests, coupled with careful examination of the terminal phenotype of the *bni1Δ cla4Δ* YCpHIS3*cla4-75* double mutant, point to actin polarization as the likely function. In particular, loss of Hof1p (required for cytokinesis), or Num1p (required for nuclear migration) is not lethal in the absence of Cla4p. However, diminution of Cdc42p activity is lethal in a *cla4Δ* mutant background (Cvrckova *et al.*, 1995). Cdc42p interacts with Bni1p and is required for polarization of the actin cytoskeleton. Moreover, *bni1Δ cla4Δ* YCpHIS3*cla4-75* mutants contain very few actin cables, whereas *ste20Δ cla4Δ*, *bud6Δ cla4Δ*, *pea2Δ cla4Δ*, and *spa2Δ cla4Δ* mutants carrying YCpHIS3*cla4-75* contain abundant cables. Together, these results support the idea that a Bni1p role in actin polarization is critical in the absence of Cla4p.

### CLA4 Synthetic Lethal Universe

The two screens for mutations that are synthetically lethal in a *cla4Δ* mutant background identified a large number of genes. The number of genes is large in absolute terms, but it is surprisingly large compared with the number of genes identified in a complementary synthetically lethal screen. Specifically, in a preliminary effort to identify mutations synthetically lethal with the absence of STE20 by using the colony-sectoring assay, only the CLA4 gene was identified (Mitchell, Goehring, and Sprague, unpublished data). The functions identified in the CLA4 synthetic screens reported herein cover a wide spectrum of cell biological processes and include bud emergence (*BEM1*, *BEM2*, *BEM4*), cytokinesis (*SHS1*), nuclear migration (*DYN2*, *NIP100*, *APC9*, *SLK19*), and cell wall maintenance (*GIM5*, *BCK1*, *CHS3*, *SKT5/CHS4*, *CHS5*, *CHS6*, *CHS7*, *FAB1*, *SLT2*, *SMI1*). Not all of the proteins, or even a majority of the proteins, involved in a particular process were identified. This finding implies that lethality does not result because an entire process has become essential in the *cla4Δ* mutant background, but rather implies that a particular activity or role of the protein has become essential.

It is often difficult to identify the targets of signaling proteins. Synthetic lethal screens for genes required in the absence of specific signaling molecules may provide a gen-

eral means to identify potential downstream targets of the signaling molecule. In this study, we screened for genes that, like STE20, were synthetically lethal with CLA4 and identified the polarisome as a potential target of Ste20p. By extension of this logic, potential downstream targets of Cla4p may be identified in synthetic lethal screens that use query mutations in STE20 or any one of the other genes identified in the CLA4 synthetic lethal universe. For example, because BNI1 is synthetically lethal with CLA4, the set of ~50 genes that are synthetically lethal with BNI1 (Tong *et al.*, 2001), which includes genes involved in bud emergence, chitin synthase III activity, and the dynein/dynactin spindle orientation pathway, may be Cla4p targets. Candidate targets are then identified by determining which single mutants exhibit phenotypes that resemble facets of the *cla4Δ* mutant phenotype. Thus, global synthetic lethal networks should be useful for large-scale mapping of functional relationships between signaling molecules and their downstream targets.

### ACKNOWLEDGMENTS

We thank J. Pringle, I. Herskowitz, I. Pretorius, and D. Lew for providing plasmids. We also thank David Rivers, Hilary Kemp, Greg Smith, and Paul Cullen for helpful comments and suggestions. This work was supported by grant GM-30027 (to G.F.S.) from the National Institutes of Health, National Research Service Award GM-18002-03 (to D.A.M.), training grant 5 T32 GM-07759 (to A.S.G.) from the National Institutes of Health, Canadian Institute of Health Research (to C.B.), and operating grant from the National Cancer Institute of Canada (C.B.).

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