Progression through and completion of mitosis require the actions of the evolutionarily conserved Polo kinase. We have determined that the levels of Cdc5p, a Saccharomyces cerevisiae member of the Polo family of mitotic kinases, are cell cycle regulated. Cdc5p accumulates in the nuclei of G2/M-phase cells, and its levels decline dramatically as cells progress through anaphase and begin telophase. We report that Cdc5p levels are sensitive to mutations in key components of the anaphase-promoting complex (APC). We have determined that Cdc5p-associated kinase activity is restricted to G2/M and that this activity is posttranslationally regulated. These results further link the actions of the APC to the completion of mitosis and suggest possible roles for Cdc5p during progression through and completion of mitosis.

**MATERIALS AND METHODS**

**Plasmids and strains.** Saccharomyces cerevisiae strains used in this study are listed in Table 1. Plasmid DNA was transformed into yeast by the lithium acetate method as described previously (15). Yeast strains bearing plasmids were grown in YPD. Yeast strains bearing plasmids were grown in selective synthetic medium (SC) with 2% sugar (galactose or raffinose as indicated). Strains with plasmids to be induced with galactose were first grown in synthetic medium with 2% raffinose to an A600 of 0.2 to 0.4. pCH740 carries the CDC5 open reading frame ligated into the BamHI site of pCH765 (pRS424–GAL1-HA, where HA is the hemagglutinin epitope). Wild-type strains were grown at 30°C unless noted otherwise in the text.

**ProA tagging of Cdc5p.** The chromosomal copy of the CDC5 gene was tagged by a C-terminus, in-frame integration of a DNA fragment encoding the immunoglobulin G (IgG) binding domains of protein A (ProA) (40). The protein A gene and adjacent HIS3 and URA3 markers were amplified by PCR using pProA–HIS3–URA3 (a gift from Mike Rout and John Atlison) (1). The following primers were used for the PCR: CDC5 sense primer (5′-GAG AAA CTA ACT TTG ATA AAG GAA GGT TTG AAG CAG AAG TCA ATT GTT ACC GTA GAT GGA GAA GGT ACC AT TTT ATG TAG TTG TAG TTA TTA AGG CCC AAT CAA ATT AAT TA CTT ATT GTC ACC AT TTT ATG TAG TTG TAG-3′) and β-tubulin and microtubule-associated proteins (36).

Recently, Polo kinases have been implicated in budding yeast (3, 32), mammalian (22), and Xenopus (5) cells in the late mitotic mechanism, which activates the cyclin-specific APC activity. In order to understand the role Cdc5p plays in this key cell cycle event, we need to know more about the regulation of its protein levels and activity during the cell cycle. We report here that Cdc5p accumulates in the nuclei of G2/M-phase but not G1 cells. We show that the levels of Cdc5p drops dramatically as cells complete anaphase and that degradation of Cdc5p in G2 is sensitive to mutations in a key APC component, Cdc23p. We also report that Cdc5p-associated kinase activity is restricted to G2/M and is posttranslationally regulated. Taken together, these results provide more evidence that Cdc5p specifically and Polo kinases in general play key regulatory roles in the pathway leading to the completion of mitosis.
TABLE 1. Yeast strains used in this study

<table>
<thead>
<tr>
<th>Strain</th>
<th>Description (source)</th>
</tr>
</thead>
<tbody>
<tr>
<td>W303-1A</td>
<td>MAT a ade2-1 his3-11,15 leu2-3,112 trpl-1 ura3-1 can1-100 LYS2 (R. Rothstein)</td>
</tr>
<tr>
<td>W303-1B</td>
<td>MAT a ade2-1 his3-11,15 leu2-3,112 trpl-1 ura3-1 can1-100 (R. Rothstein)</td>
</tr>
<tr>
<td>YCH3</td>
<td>CDC5-ProA-HIS3-URA3 bar1 1 spc42::LEU2 TRP1::SPC42-GFP (segregant of YCH199 × YJC1203)</td>
</tr>
<tr>
<td>K193</td>
<td>CDC5-proA-URA3 bar1 1 ade2 ura3 his3 (backcrossed 4 times to W303) (K. Nasmyth)</td>
</tr>
<tr>
<td>YCH212</td>
<td>CDC5-ProA-HIS3-URA3 (ProA-tagged CDC5) in K1983</td>
</tr>
<tr>
<td>YCH30</td>
<td>CDC5-ProA-HIS3-URA3 bar1 1 spc42::LEU2 TRP1::SPC42-GFP (segregant of YCH121 × YJC1203)</td>
</tr>
<tr>
<td>K2034-CH-a</td>
<td>CDC5-ProA-HIS3-URA3 bar1 1 ade2 ura3 his3 (backcrossed 5 times to W303) (K. Nasmyth backcrossed 3 times and then we backcrossed it an additional 3 times to W303)</td>
</tr>
</tbody>
</table>

Cdc5p was epitope tagged with the five IgG binding sites of ProA (for protein A) in both wild-type and cdc mutant cells (40). The Cdc5-ProA-tagged strains exhibited no growth defects, and therefore, the Cdc5-ProA fusions must be performing all the essential functions of Cdc5p. We initially examined

**RESULTS**

Cdc5p protein levels decrease as cells exit mitosis. **CDC5-ProA cdc15-2** (YHC306) cells were synchronized in telophase by growth at 37°C for 3 h prior to release into fresh medium at 23°C. Samples for immunoblot analysis of Cdc5-ProA and actin, FACS analysis of DNA content, and detection of anaphase spindles by immunofluorescence staining of tubulin were taken at the times (in minutes) indicated. The percentage of cells with anaphase spindles is shown on the left side of the FACS analysis profile. Spindle morphology in cells was determined by indirect immunofluorescence staining with an anti-tubulin antibody. Actin detection was used as an internal loading control. Cells released from a cdc15-2 block are delayed in cytokinesis which results in a 2n and 4s shift in DNA content after replication.
the fluctuations of Cdc5p levels in a population of cdc15-2 cells synchronously released from a temperature-induced (37°C) telophase arrest state. Progression through the cell cycle was followed by FACS analysis to determine DNA content and by indirect immunofluorescence using tubulin staining to determine the percentage of cells with anaphase spindles. These results (Fig. 1) show that Cdc5p levels are high in cells blocked in telophase and that the levels decline dramatically 40 min after release from the temperature block, when virtually all the cells have entered G1. Cdc5p begins to reaccumulate after 80 min to 90 min when the cells have finished or are just finishing S phase but have not yet entered anaphase.

**Cdc5p degradation is APC dependent.** The pattern of CDC5 message and of Cdc5p levels is reminiscent of the message and protein levels of mitotic cyclins which decline sharply as cells complete anaphase (12, 19, 21). Mitotic cyclins are targeted for degradation by the APC as cells complete anaphase (19). During G1, the APC is active in Pds1p, Clb2p, and Ase1p degradation (2, 4, 18, 45). Moreover, the activity of the APC in G1 cells can be specifically inhibited by mutation of CDC23 encoding a subunit of the APC (4). To determine whether the cell cycle-regulated pattern of Cdc5p loss was a result of APC-mediated degradation, the stability of Cdc5p was examined in wild-type versus cdc23-1 G1-arrested cells. CDC23 and cdc23-1 cells containing plasmids expressing HA-Cdc5p under control of the GAL1 promoter were grown in raffinose at 23°C, synchronized in G1, with α-factor, and then shifted to 37°C to inactivate the cdc23-1 gene product. At this point, HA-Cdc5p expression was induced by the addition of galactose for 15 min, followed by the addition of glucose to turn off its expression. The restrictive temperature (37°C) was maintained while performing these steps. As shown in the immunoblot in Fig. 2, the majority of HA-Cdc5p was degraded after 60 min in the wild-type cells (CDC23) but was still present after 2 h in the cdc23-1 cells (Fig. 2). These results provided evidence that Cdc5p proteolysis might be APC mediated.

**Cdc5p accumulates in the nuclei of G2- and M-phase cycling cells.** To determine the spatial and temporal dynamics of Cdc5p in cells, we performed indirect immunofluorescence on Cdc5-ProA Spc42-GFP cells (YCH301). These experiments showed that in an asynchronous culture Cdc5-ProA was detected in only a subset of cells and that in these cells Cdc5-ProA was predominantly localized in the nucleus (Fig. 3). Multiple fields of such asynchronous cells were quantitated for the presence or absence of Cdc5-ProA staining (Fig. 4). Spc42p, a component of the spindle pole body (SPB) was tagged with the green fluorescent protein Spc42-GFP, which allowed detection of SPBs in this (YCH301) and other strains used in this report (6). Cdc5-ProA was rarely detected in unbudded cells. Of 75 unbudded cells in compiled asynchronous fields, 70 of the cells had the cytological phenotype shown in Fig. 4 (I), exhibiting no detection of Cdc5-ProA signal. Cdc5-ProA was also rarely detected in late mitotic cells, with only 6 of 97 large budded cells IVa having signal or 91 of 97 in class IVb with no signal (Fig. 4). These late mitotic cells were distinguished by two distinct nuclei and SPBs localized to the poles. Cdc5-ProA was detected in approximately one half of the budded cells containing a single nucleus with duplicated SPBs localized exclusively to the mother cell (Fig. 4, Ia and Ib). In contrast, the large fraction (47 of 50) of cells undergoing anaphase exhibited an intense Cdc5p staining pattern, where the nucleus was in the mother and daughter neck and SPBs were widely separated (Fig. 4, III). These results, combined with our previous immunoblot analysis of Cdc5p levels during the cell cycle, suggested that Cdc5p was not present during G1, did not begin to accumulate until at least late S, and disappeared as cells finish anaphase but before they completed mitosis or began cytokinesis.
Cdc5p accumulates in the nucleus of S- as well as M-phase-arrested cells. To more fully determine the subcellular pattern of Cdc5p localization during the cell cycle, the CDC5-ProA epitope-tagged gene was expressed in a panel of cdc mutant strains and immunofluorescence was performed on the temperature-arrested cells. Cdc5-ProA was not present in cells synchronized and immunofluorescence was performed on the temperature-to-our immunofluorescence results with the same cdc4-1 a-facor-factor-arrested bar1 cells and cdc4-1 cells grown at the restrictive temperature as shown in Fig. 5. Cdc5-ProA was observed only in the nucleus, and its signal was present in cells synchronized in S phase (cells arrested with HU) and in cells synchronized early in M, during metaphase (cdc13-1, nocodazole, and cdc23-1) (Fig. 5). Cdc5-ProA was also detected in the nuclei of cdc15-2 cells synchronized late in mitosis during telophase (Fig. 6). The arrest state of these cells was determined by FACS analysis (see Fig. 7g).

Immunoblot analysis was also performed using extracts derived from these synchronized cells. As shown in Fig. 7a, Cdc5-ProA was not detected in extracts derived from cells arrested in late G1 with α-factor. However, a low level of Cdc5-ProA was detected in cdc4-1 late-G1-arrested cells. This is in contrast to our immunofluorescence results with the same cdc4-1 arrested cells in which Cdc5-ProA staining was not detected (Fig. 4). This apparent discrepancy might be explained if the level of Cdc5p in cdc4-1 arrested cells is below the threshold level required for Cdc5p staining. Cdc5-ProA was also detected by immunoblotting in cells blocked in very early S. Cells synchronized in G1 with α-factor were released into medium containing HU at 23°C (Fig. 7a, right blot). In agreement with our immunofluorescence results, Cdc5-ProA was detected by immunoblot analysis in extracts derived from cells synchronized in S phase (with HU), G1, (cdc13-1, cdc23-1, and nocodazole), and those synchronized late in M (cdc15-2). We detected a low level of mutant cdc5-1-proA protein in cdc5-1 cells synchronized late in M. We assume that CDC5 message is transcribed at a low level in cells which are cycling from late G1 through S phase and that Cdc5p may accumulate in cells held for prolonged periods in late G1 or S. Alternatively, the process of synchronizing cdc4-1 cells in late G1 and wild-type cells in S with HU may activate the CDC5 promoter.

Cdc5p-associated kinase activity fluctuates during the cell cycle. Although Cdc5p levels peak during G2/M, it was not known when the Cdc5p kinase was active. We measured the Cdc5p-associated kinase activity in cells synchronized in G1 with α-factor and released into fresh medium lacking α-factor. At intervals, samples were lysed, Cdc5-ProA was immunoprecipitated, Cdc5-ProA-associated kinase assays were performed, and the results were measured by immunoblotting (Fig. 8c) and autoradiography (Fig. 8D). Quantitation of the Cdc5p-associated kinase activity is shown in Fig. 8e. Cdc5p-associated kinase activity in these cells peaked in late G2/early M-phase cells 70 and 80 min after release and was not detectable in cells in earlier stages of the cell cycle (Fig. 8d). It is noteworthy that this peak in activity was 20 min after the appearance of Cdc5p (Fig. 8a). This suggests that Cdc5p-associated kinase activity may be cell cycle regulated (compare Fig. 8a and d).

As controls for the Cdc5p-specific nature of the kinase activity in these immunocomplexes, the Cdc5p-associated kinase activity derived from extracts of untagged wild-type cells or ProA-tagged cdc5-1 mutant cells was found to be 7- to 15-fold lower than that obtained from mitotic Cdc5-ProA-tagged cells (Fig. 7d and e). Low levels of Cdc5p-associated kinase activity were observed in ProA-tagged cdc5-1 cells synchronized in telophase by growth at the restrictive temperature (Fig. 7d), cells grown asynchronously, or cells synchronized in metaphase with nocodazole (Fig. 7f). The temperature at which the cells were grown (23 or 37°C) or at which the kinase assay was
performed (23, 30, or 37°C; data not shown) did not alter these
low levels. These results suggest that the cdc5-1 mutant protein
may be unstable when isolated from cells grown under either
permissive or restrictive conditions.

Following up on these results, we examined Cdc5p protein
levels and associated kinase levels in populations of cells syn-
ochronized in specific phases of the cell cycle. As shown in Fig.
7, although Cdc5p was present in cells synchronized in G1
(cdc4-1), early S (α-factor arrest into HU) and S (HU) phases,
the Cdc5p-associated kinase activity at these times was not
above background levels (compare Fig. 7a, c, and d). Quantifi-
tation of the Cdc5p-associated kinase activity in these synchro-
nized samples is shown in Fig. 7e. Cells synchronized in early
(nocodazole [NZ], cdc13-1, and cdc23-1) and late stages of M
(cdc15-2) all had high levels of both Cdc5p protein and asso-
ciated kinase activity. The low level of Cdc5p-associated kinase
activity in cells synchronized in S phase with HU could be due
to the low levels of Cdc5-ProA in the immunoprecipitate (Fig.
7c). However, even when the level of Cdc5-ProA in the immu-
noprecipitate was increased fourfold, the Cdc5p-associated ki-
nase activity was still not significantly above background (Fig.
7c and d, ×4 HU). The results taken from Fig. 6 and 7 suggest
that Cdc5p-associated kinase activity is not active until G2 or
metaphase and may therefore be regulated.

Modification of Cdc5p in cdc13-1 cells requires Mec1p,
Mec2/Rad53p, and Rad9p. During the immunoblot analysis of
Cdc5p levels in cell cycle-arrested cells (Fig. 7), we observed
that the electrophoretic mobility of Cdc5p was modified in
cdc13-1 cells. The migration of Cdc5p in cdc13-1 arrested cells
was notably slower. In contrast, the shifted form of Cdc5p was
not observed in wild-type cells cycling through the cell cycle
(Fig. 8) or in cells grown in either HU or nocodazole, which
arrest in S phase and metaphase, respectively (Fig. 7). The
appearance of the modified form of Cdc5p in cdc13-1 cells was

FIG. 5. Cdc5p is present in cells blocked in S and M phase as detected by immunofluorescence. Wild-type and cdc mutant cells were synchronized at different stages
of the cell cycle using either chemicals or by growth of the cdc mutant at the restrictive temperature of 37°C and processed for immunofluorescence. Four views of each
cell are shown: Nomarski optics (NOM), staining of DNA (DAPI), Cdc5p-ProA staining (Texas red), and Spc42-GFP (direct fluorescence). The cells were synchronized
at the different stages as follows: late G1, with α-factor (YCH301) and cdc4-1 (YCH303); S phase, with HU (YCH301); Metaphase, cdc13-1 (YCH309), nocodazole
(Na) (YCH301), and cdc23-1 (YCH305). Samples of the arrested cells used in this study were also used for FACS, immunoblot, and kinase assay analyses shown in Fig.
7. All images were taken with a 100× objective and printed at the same magnification. Bar = 5 μm.

FIG. 6. Cdc5p is present in cells synchronized in telophase as detected by
immunofluorescence. CDC5-ProA cdc15-1 cells (YCH307) were synchronized in
telophase by growth at the restrictive temperature and processed for immuno-
fluorescence; CDC5 and SPC42 are not tagged in cdc15-2* cells (YCH238). Four
views of each cell are shown: Nomarski optics (NOM), staining of DNA (DAPI),
Cdc5p-ProA staining (Texas red), and Spc42-GFP (direct fluorescence). All
images were taken with a 100× objective and printed at the same magnification.
Bar = 5 μm. ND, not done.
FIG. 7. Cdc5p is detected in late G1; S- and M-phase synchronized cells, but Cdc5p-associated kinase activity is restricted to cells arrested in M Phase. Wild-type and cdc mutant cells were arrested in specific stages of the cell cycle using either chemicals or by growth at the restrictive temperature of 37°C, respectively. The cells were synchronized at the various stages as follows: late G1, with α-factor (YCH301) and cdc4-1 (YCH303); G1/S, cells were initially blocked in G1 with α-factor (YCH199) and then released into fresh medium containing HU; S phase, with HU (YCH301); metaphase (Meta), with cdc13-1 (YCH309), nocodazole (NZ) (YCH301), and cdc23-1 (YCH305); and telophase (Tel.), with cdc15-2 (YCH307) and cdc5-1 (YCH214). Extracts from asynchronous (Async) cells were derived from cultures of Cdc5-ProA (YCH199) and strain W303a expressing an untagged Cdc5p. Samples for FACS analysis and extract preparation were taken when >95% of the cells in the culture were appropriately arrested, as detected by light microscopy. Cdc5-ProA (a) and actin (b) in the crude extract were detected by immunoblotting with anti-IgG and antiactin antibodies, respectively. (c) Cdc5-ProA was routinely immunoprecipitated from 400 μg of extract with IgG-Sepharose beads and detected as above by immunoblotting. (d) Kinase activity in these immunocomplexes was measured as described in Materials and Methods with casein as the substrate, except in the case of the 4× HU lane, where 1.6 mg of extract was used. (e) A bar graph of the [32P]-casein levels is shown. Levels were quantitated with a Molecular Dynamics PhosphorImager. (f) Cdc5-ProA kinase activity was determined for cdc5-1-proA (YCH214) cultures grown either asynchronously (Async) (23°C), in the presence of nocodazole (NZ) (3 h, 23°C) or in the presence of nocodazole (3 h, 23°C) followed by a 1-h shift to 37°C (NZ→37°C). Cdc5-ProA kinase activity was also determined for a wild-type strain expressing Cdc5-ProA (YCH199) arrested in the presence of nocodazole and loaded on the same gel. (g) FACS analysis of DNA content.
further monitored in a time course following a shift of the cdc13-1 culture to the restrictive temperature (Fig. 9a). The shift was apparent after only 1 h of growth of the cdc13-1 strain at the restrictive temperature. The cdc13-1 mutant cells are known to activate the DNA damage checkpoint when they are grown under restrictive conditions (7). Therefore, we next asked whether the MEC1, MEC2/RAD53, and RAD9 genes were necessary for the modification. These three gene products are required for the DNA damage checkpoint pathway (7). We monitored the shift of Cdc5p in cdc13-1 checkpoint defective strains by immunoblotting (cdc13-1 rad53-21, cdc13-1 rad9::URA3, and cdc13-1 mec1-1) following 3 h of growth at the restrictive temperature (Fig. 9b). In all three strains, the shifted form of Cdc5p was not observed. In addition, the shifted form was not observed after extended growth (7 h) of the double mutant strains at the restrictive temperature (Fig. 9c). Thus, accumulation of the shifted form of Cdc5p is dependent upon Mec1p, Mec2/Rad53p, and Rad9p function.

Cdc5p-associated kinase activity is phosphorylation dependent. The phosphorylation state of a given kinase has often been found to play a key role in determining the activity of the kinase (27). To determine whether the Cdc5p-associated kinase activity was dependent on phosphorylation, Cdc5-ProA was immunoprecipitated from lysates of nocodazole-arrested
cells and incubated with calf intestinal phosphatase (CIP) prior to the kinase assay. The kinase activity of the CIP-treated sample was dramatically lower than that of the untreated sample (Fig. 10a, right panel). In contrast, only a slight reduction was observed if the CIP was boiled for 10 min prior to use. The lack of full activity in the presence of boiled CIP may simply be due to incomplete inactivation by boiling. As a further control, the phosphatase reaction was repeated in the presence of β-glycerophosphate, a phosphatase inhibitor. Under these conditions, no reduction in Cdc5p-associated kinase activity was observed (Fig. 10b). It is interesting to note that Cdc5p may possess autophosphorylation activity as Cdc5p appeared to be phosphorylated in the untreated lane of Fig. 10a (right panel). In contrast, only a slight reduction was observed if the CIP was boiled for 10 min prior to use. The kinase activity of the CIP-treated sample was dramatically lower than that of the untreated sample (Fig. 10a, right panel). Further in vitro studies will be required to determine the nature of the phosphorylation that may be regulating Cdc5p-associated kinase activity.

DISCUSSION

Based on immunoblot and indirect immunofluorescence analyses, we report here that the levels of the conserved Polo kinase Cdc5p are strictly regulated during the cell cycle and steeply decline as cells exit mitosis. Cdc5p accumulates in the nuclei of G2 and early M-phase cells and disappears from cells steeply decline as cells exit mitosis. Cdc5p accumulates in the nuclei of G2 and early M-phase cells and disappears from cells as they complete anaphase. We show that the levels of Cdc5p are sensitive to mutations in components of the APC (Fig. 2) and its disappearance from cells as they complete anaphase (Fig. 1) suggest that Cdc5p is targeted for degradation by the APC in the same manner as the mitotic cyclins are. While this research was being done, similar results were reported by other labs (3, 32), including one report that shows that Cdc5p is indeed ubiquitinated in an APC-dependent manner (32). Cdc5p is a member of the Polo family of kinases. Other members of the family including Pklp1 from mammals exhibit the same cell cycle-regulated pattern of message and protein levels as Cdc5p, and therefore, their levels late in M may also be regulated by the APC (10, 25).

Recently, it has been shown that Cdc5p plays a positive role in regulating cyclin-specific APC activity (3, 32). The APC-associated cyclin ubiquitin ligase activity was reduced in cdc5-1 cells even at the permissive temperature and increased in wild-type cells which overexpress CDC5 (3). Our finding that cdc5-1 cells have low Cdc5p-associated kinase activity, at any temperature, suggests that Cdc5p-associated kinase activity is required to activate the APC. By analogy, studies on Polo kinases from mammals suggest that Polo kinases might play a direct role in activating the APC. The mammalian homologue of Cdc5p, Pklp1, interacts with and phosphorylates three components of the APC and this phosphorylation activates the APC to ubiquitinate cyclin B in vitro (22). Further support for a role of Polo kinases in this regulatory process comes from dominant negative and immunodepletion experiments with Xenopus, which indicate that Pkl1 (Polo-like in Xenopus) is required for M-phase exit and destruction of mitotic cyclins (5).

It is interesting to note that a regulator of the APC, Cdc5p, is itself a target of the APC. Removal of Polo kinases late in mitosis may allow G1 cells to target a different or G2 class of targets for APC-mediated degradation. Alternatively, the removal of Cdc5p late in mitosis may serve to prepare cells for the eventual inactivation of the APC late in G1 (19). However, it is not clear from our study or any of the published Cdc5p analyses how the cell coordinates the APC-mediated degradation of Cdc5p with that of Cdc5p or how it maintains its APC activity toward Cdc5p in the absence of Cdc5p during...
Clearly, other factors, including perhaps Hct1p, may play roles in the activation and maintenance of APC function during G1 (30, 42). Further studies are required to understand the complex links between Cdc5p and the APC and their ramifications for progression through and completion of mitosis.

**Regulation of Cdc5p kinase activity.** In addition to the regulation of Cdc5p levels by the APC, the timing of Cdc5p kinase activity is also posttranslationally regulated. In studies of synchronized populations of cycling cells, we determined that the peak in Cdc5p-associated kinase activity significantly lagged behind that of Cdc5p levels. Cdc5p was detected in HU- and nocodazole-arrested cells, which have activated the DNA replication and spindle assembly checkpoints, respectively. However, Cdc5p-associated kinase activity was detected only in the population of nocodazole-arrested cells. We also showed that the DNA damage checkpoint, which is activated in these HU-treated cells (7), regulates Cdc5p kinase activity. In this model, the activation of Cdc5p-associated kinase activity may be dependent upon the prior completion of DNA replication. Alternatively, as suggested above, Cdc5p-associated kinase activity may simply be dependent upon an M-phase-specific activity, such as Clb/Cdk, for its activation.

**Role for Cdc5p in adaptation to the DNA damage response.** We observed that the electrophoretic mobility of Cdc5p is modified under conditions that induce the DNA damage checkpoint. In cdc13 mutant cells grown at the restrictive temperature, the mobility of Cdc5p is slower than that of Cdc5p derived from wild-type cells (Fig. 9a). In cdc13 arrested cells, Cdc5p-associated kinase activity is high and its modification is dependent upon the factors required to transduce the DNA damage signal including Mec1p, Mec2/Rad53p, and Rad9p (7). It has recently been reported that although the kinase activity associated with the mutant cdc5-ad protein in cdc13-arrested cells is high, its stimulation of APC activity under these conditions is dramatically lower than that of wild-type Cdc5p (3). The failure of cdc5-ad cells to adapt may merely reflect its inability to specifically target and to activate the APC. It will be interesting to determine whether the Cdc5-ad mutant protein is modified under conditions, which activate the DNA damage checkpoint, and whether this modification is required for the adaptation response. In summary, Cdc5p, a regulator of adaptation to the DNA damage response, may also be regulated by the DNA damage response checkpoint.
Additional roles for Cdc5p. The results of our previous study suggests that Cdc5p may have a role in DNA replication (12). We suggested in our previously published study that Cdc5p could play a role in activating the degradation of the Clb-Cdk kinase activity that is implicated in the exit from mitosis (12). Proteolysis of the Clb cyclins is required for the timely transition during anaphase/telophase, between the post- and prereplication complexes present at origins (28). The low level of cyclin-specific APC activity in cdc5-1 mutant cells late in mitosis may affect the efficient formation of prereplication complexes at origins at that time (3). Such a model could help explain the cdc5-1 mutant plasmid loss defect that can be suppressed by the addition to the plasmid of multiple replication origins (12).

Additional support for a Cdc5p role in regulating replication initiation is the interaction between Cdc5p and the origin interacting factor Dbf4p (12). Alternatively, the Cdc5p-Dbf4p interaction may reflect a role for Dbf4p during the late stages of mitosis. In support of this hypothesis, we have determined that Dbf4p is absent from cells during early G1, begins to accumulate in G1, persists through anaphase, and is degraded as cells finish mitosis (our unpublished data). Interestingly, we have found that like Cdc5p, the levels of Dbf4p are sensitive to mutations in a key APC component, CDC23 (our unpublished data). These results suggest that the APC-mediated removal of Dbf4p from cells late in mitosis may be important for the timing of the transition at origin complexes.

Additional factors required to complete mitosis. In addition to Cdc5p, there is a group of factors which are required to complete mitosis. These factors include the kinases Cdc15p (14) and Dbf2p (39), a Ras-like GTPase (Tem1p) (31), a nucleotide exchange factor (Lte1p), and a phosphatase (Cdc14p) (43). A number of genetic interactions between these genes have been reported, and they suggest a possible concerted role during the completion of mitosis (17, 21, 38, 39). cdc15-2 cells grown at the restrictive temperature arrest uniformly in telophase with high levels of Clb/Cdk and Cdc5p-associated kinase activity (this report) but low levels of APC activity (14, 17). These results show that Cdc15p is not required to activate Cdc5p-associated kinase activity and that this activity alone is not sufficient to activate the APC. To move the field forward, the role of these other factors in regulating not only APC and subsequent Cdc5p/Cdk function at the end of mitosis but also that of Cdc5p must be further investigated.

ACKNOWLEDGMENTS

We thank L. Hartwell, K. Nasmyth, J. Cooper, D. Lew, O. Cohen-Fix, D. Koshland, and M. Rout for strains and plasmids; S. Dowdy and lab for excellent help with FACs; S. Wente for reading the manuscript; H. Piwnica-Worms for helpful discussions; J. A. Cooper and T. Karlab for excellent help with FACS; S. Wente for reading the manuscript; Fix, D. Koshland, and M. Rout for strains and plasmids; S. Dowdy and V. A. Nigg. 1995. Cell cycle regulation of the activity and subcellular localization of PLK1, a human protein kinase implicated in mitotic spindle function. J. Cell Biol. 129:1617–1628.

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This work was supported by grants RPG-97-162-01-CCG from ACS and GM578801 from NIH to C.J.H.

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