Cyclical inactivation of B-type cyclins has been proposed to be required for alternating DNA replication and mitosis. Destruction box-dependent Clb5p degradation is strongly increased in mitotic cells, and constitutive overexpression of Clb5p lacking the destruction box resulted in rapid accumulation of inviable cells, frequently multiply budded, with DNA contents ranging from unreplicated to apparently fully replicated. Loss of viability correlated with retention of nuclear Clb5p at the time of nuclear division. CLB2-Δdb overexpression that was quantitatively comparable to CLB5-Δdb overexpression with respect to Clb protein production and Clb-associated kinase activity resulted in a distinct phenotype: reversible mitotic arrest with uniformly replicated DNA. Simultaneous overexpression of CLB2-Δdb and CLB5-Δdb overexpressors similarly resulted in a uniform arrest with replicated DNA, and this arrest was significantly more reversible than that observed with CLB5-Δdb overexpression alone. These results suggest that Clb2p and not Clb5p can efficiently block mitotic completion. We speculate that CLB5-Δdb overexpression may be lethal, because persistence of high nuclear Clb5p-associated kinase throughout mitosis leads to failure to load origins of replication, thus preventing DNA replication in the succeeding cell cycle.

Testing Cyclin Specificity in the Exit from Mitosis

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Cyclin-dependent kinase activity drives the eukaryotic cell cycle. In Saccharomyces cerevisiae, three G1, or CLN, cyclins and six B-type, or CLB, cyclins bind and activate the cyclin-dependent kinase Cdc28p. CLB function is required for initiation of DNA replication, spindle formation, and initiation of mitosis. With respect to DNA replication and mitosis, the main role of the CLN cyclins is to allow activation of Cdcp-Cdc28p kinase, although the CLN cyclins have additional cell cycle roles (6, 26).

It is likely that all of the CLB cyclins are descendants of a single B-type cyclin-like ancestor, and it has been proposed (28) that a single B-type cyclin regulated both DNA replication and mitosis in a primordial eukaryotic cell. Multiple B-type cyclins derived from gene duplication have diverged in function. Functional divergence could simply reflect different timing of accumulation of functionally interchangeable cyclins; alternatively, specific cyclin coding sequences could have become intrinsically specialized for particular cell cycle roles. Recently, we showed that Clb5p is intrinsically specialized for activation of replication in comparison to Clb2p (8).

Clb2p-Cdc28p kinase drives some essential step(s) in replication, including the binding of Cdc45p and replication protein A (RPA) to the prereplicative complex (PRC) (44, 50). The PRC is formed by Cdc6p-dependent loading of minichromosome maintenance (MCM) proteins onto origins of DNA. The Cdc14p phosphatase (20, 41, 47). Clb5p is not under the control of Cdc14p, and relatively little cell cycle regulation of Clb5p degradation has been observed (39), although its degradation was reported to be destruction box dependent and dependent on components of the APC that are also required for Clb2p ubiquitination and degradation (19). Clb5p and the anaphase inhibitor Pds1p may both be targets of Cdc20p-directed APC degradation, because it has been observed that CLB5 rescues cdc20 pds1 strains which would otherwise arrest in late anaphase, and deleting cdc20 stabilizes Clb5p (40).

Here we report on cell cycle dependence of Clb5p degradation. We also compare the effects of expression of stabilized Clb5p and Clb2p on cell cycle progression, to examine the issue of intrinsic Clb specialization in driving cell cycle events.

MATERIALS AND METHODS

Yeast strains. All strains are isogenic with 15Dau (MATa leu2 ura3 trp1 his2 ade1) (33). Strains were constructed and analyzed by standard genetic methods. DNA transformations were done by the lithium acetate method.

Plasmids. All plasmids were derived from CE119, a YCP50-based GAL1::CLB5 URA3 CEN4 ARS1 construct (11, 30). CE119-4, a GAL1::CLB5-Δdb-HA construct was described previously (8). A point mutation, S399P, presumably generated during the PCR-based construction of CE119 (11), was present in CE119-4 and in the GAL1::CLB5-Δdb-H4 construct DB4 were described previously (8). A point mutation, S399P, presumably generated during the PCR-based construction of CE119 (11), was present in CE119-4 and in the GAL1::CLB5-Δdb-H4 construct. The S399P point mutation is relatively innocuous; it slightly reduces Clb5p function, but CLB5-S399P under control of the CLB5 promoter fully rescues the clb5 replication defect (12) and clb3,4,5,6 lethality (37), and GAL1::CLB5-Δdb-S399P is lethal with a phenotype.
similar to that of the S399S version (data not shown). The experiments in Fig. 1, 2, 4, and 5 were performed with S399P versions of GAL1::CLB5. The remaining experiments were performed with constructs in which the S399P mutation was repaired by subcloning the wild-type fragment to make CE119-4R and DB4R. We assume that this does not significantly affect the results. The destruction box-dependent instability of Clb5p documented in Fig. 2 has been found to be similar to those of S399P and S399S versions of Clb5p (compare Fig. 2B and C). The control GAL1-Clb5fsHA construct is a product of the GAL1 promoter at any time before 100 min after release of a block, by incubation in raffinose for 2.5 h to turn off GAL1-CLB5-HA expression, we constructed wild-type strains containing integrated GAL1::CLB5-HA or GAL1::CLB5-db-HA (lacking amino acids 56 to 64) (8). Addition of galactose to cultures of these strains yielded comparable initial accumulation of Clb5p and Clb5Δdbp protein and associated kinase (Fig. 2A). Glucose addition to such galactose-induced cultures could inactivate GAL1-driven transcription showed that the destruction box-containing protein decayed much faster than the destruction box-deleted protein (Fig. 2B). A longer time course revealed that Clb5p lacking its destruction box was still somewhat unstable (Fig. 3A). In addition, at best, minor stabilization of Clb5Δdbp was observed when it was expressed from the endogenous CLB5 promoter (Fig. 3B and C), consistent with a hypothesis that additional regions of Clb5p may contribute to its targeted degradation. Involvement of a Skp1-Cdc53-F box (SCF)-ubiquitinating activity in Clb5p degradation was suggested previously (4). We employed timed glucose addition to shut off GAL1-

**RESULTS**

Clb5p degradation is cell cycle regulated. We constructed a cln1 cln2 cln3 GAL1::CLB5-HA strain, in which CLB5-HA substitutes for the CLN G1 cyclins in driving cell cycle initiation (12, 30). This strain was synchronized by raffinose block in G1, followed by release into the cell cycle with galactose addition to induce expression of the GAL1 promoter. Using timed glucose addition to repress GAL1::CLB5-HA expression, we saw a sharp increase in Clb5p instability approximately coincident with the time of nuclear division (Fig. 1). Consistent with this, shutoff of the GAL1 promoter at any time before 100 min (approximately the time of nuclear division) resulted in failure to bud in the next cell cycle (Fig. 1), consistent with a drop in Clb5p to a nonfunctional level during division (12, 30).
CLB5-HA transcription in cells synchronized by α-factor block-release, in which the cultures were blocked in raffinose plus α-factor medium and released into galactose medium lacking α-factor (YPGal). We did this with cells with one or two copies of GAL1::CLB5-HA or one copy of GAL1::CLB5Δdb-HA. Clb5p was moderately stable in cells completing DNA replication (glucose addition 60 min after release), but by 90 min after release, Clb5p became highly unstable. After division (120 min), Clb5p became highly stable, and stability decreased again later in the second cell cycle. Thus, Clb5p stability is cyclically regulated, with peak instability in dividing cells, consistent with the results in Fig. 1. The destruction box-deleted Clb5p protein was degraded similarly slowly at each time point. We controlled for increased Clb5p levels due to destruction box deletion with the two-copy GAL1::CLB5-HA integrant; the pattern of instability with this strain was similar to that with the one-copy strain (Fig. 2C). Thus, we observed destruction box-dependent and cell-cycle-dependent Clb5p degradation, with peak instability in dividing cells. The GAL1::CLB5Δdb-HA strain did not complete nuclear division during the time course (Fig. 2C), and an accumulation of cells with divided nuclei was observed. A further examination of the effects of GAL1::CLB5Δdb-HA is presented below.

Clb5p, but not Clb5Δdbp, is degraded in the nucleus at mitosis. We replaced the HA tag with a Myc tag to allow immunofluorescent detection of Clb5p (Fig. 4). (In our hands, the HA tag is not suitable for immunofluorescent detection.) In unbudded cells, small budded cells, and most large budded cells with an undivided nucleus, Clb5p is concentrated in the nucleus. In large budded cells with an undivided DNA mass near or spanning the bud neck and in large budded cells with two DNA signals, Clb5p is distributed diffusely throughout the nucleus.
This pattern indicates that Clb5p accumulates in the nucleus before budding and during DNA replication, but during mitosis, Clb5p nuclear abundance is strikingly reduced.

We compared the localization patterns of Myc-tagged protein throughout the cell cycle in GAL1::CLB5-MYC, GAL1::CLB5△db-MYC, and control strains blocked in raffinose plus α-factor medium and released into galactose medium lacking α-factor. Strains were either released into yeast extract-peptone-dextrose (YPD) (B) or released into YPD and rearrested with α-factor (+αF) 60 min later (C). Synchrony was gauged by counting the percentage of unbudded cells. CLB5△db expression was monitored by immunoblots (anti-HA) and associated histone H1 kinase blots (H1-P).

We compared the localization patterns of Myc-tagged protein throughout the cell cycle in GAL1::CLB5-MYC, GAL1::CLB5△db-MYC, and control strains blocked in raffinose plus α-factor medium and released into galactose medium lacking α-factor. The synchrony between samples was similar during DNA replication and entry into nuclear division, but the GAL1::CLB5△db-MYC culture did not complete nuclear division during the time course (Fig. 5). The accumulation of binucleate cells in this α-factor synchrony protocol is reduced to very low levels in the nucleus of most cells when long spindles became abundant, and then became concentrated in the nucleus of most cells again after cytokinesis (Fig. 5), consistent with the results with asynchronous culture (Fig. 4). Clb5-△dbp remained concentrated in the nucleus throughout the time course, including in cells with DAPI staining, similar to that observed in cells with long spindles (Fig. 5 and data not shown). Our localization results using overexpressed Clb5p agree with those recently reported for endogenously expressed Clb5p (40). Our results also suggest that the observed nuclear persistence of Clb5-△dbp around the time of division is destruction box dependent. Removal of Clb5p from the nucleus is likely to be due to degradation resulting from Cdc20p-dependent targeting of the APC to Clb5p (40).

Removal of Clb5p regulation. CLB5 is controlled in three ways: it is transcriptionally induced early in the cell cycle (12), its associated kinase activity is inhibited by Sic1p (36), and it is degraded in a destruction box-dependent manner (14, 19 [see above]). To address the biological significance of proteolytic control, we deleted the CLB5△db allele (8). GAL1::CLB5 expression results in the accumulation of cells with 2C DNA content (see Fig. 8). GAL1::CLB5 expression becomes lethal in a σ1c1 background (data not shown). Deletion of the SWE1 inhibitory kinase (5)
GAL1-CLB5MYC

currence (Anti-Myc) from 1255-5C (wild type) strains carrying integrated copies of staining (DAPI), and Myc-tagged protein detected by indirect immunofluorescence (DIC), nuclei visualized by DAPI (data not shown).

had little additional effect on these phenotypes (data not shown).

In contrast, CLB5-Dlpb expressed from its own promoter was not lethal (S), even in the absence of SIC1 (data not shown). Therefore, loss of transcriptional control (of periodicity, levels or both), in addition to loss of either proteolytic or Sic1p control, was required to elevate Clb5p activity sufficiently to achieve lethality. Clb5p-Dlpb expressed from the CLB5 promoter was moderately if at all stabilized in cell cycle time courses, in contrast to the strong stabilization observed with GAL1::CLB5-Dlpb (Fig. 2 and 3). It may be that overexpression of Clb5p-Dlpb is required to saturate some means of Clb5p degradation that is independent of the identified destruction box. These observations on expression of CLB5-Dlpb pose a paradox with respect to the results of Shirayama et al. (40). If Cdc20-dependent degradation of Clb5p is destruction box-dependent, and if failure of Cdc20-dependent degradation of Clb5p (expressed from its own promoter) blocks mitotic exit, then CLB5-Dlpb expression should similarly result in significant persistence of Clb5p and a block to mitotic exit. There may be additional unidentified destruction boxes in Clb5p (S. Holloway, unpublished data), or there may be Cdc20-dependent but destruction box-independent means of Clb5p degradation. We are exploring these possibilities.

Since GAL1::CLB5-Dlpb and GAL1::CLB5 yield nearly comparable levels of protein through most of the cell cycle, it seemed likely that the lethality of GAL1::CLB5-Dlpb may require specific persistence of Clb5p through mitosis, when Clb5p but not Clb5-Dlpbp is degraded. Since Sic1p, protein accumulation is induced during late mitosis, this may explain the lethality of GAL1::CLB5 in sic1 strains: a low level of residual Clb5p escaping degradation may require Sic1p inhibition to avoid lethality due to active Clb5p complexes persisting through mitosis. Overexpressed Clb5p-Dlpb may saturate the available Sic1p.

Confirming that high levels of Clb5p activity are required to induce lethality, the introduction of a double mutation (K253A, E282A) that partially interferes with Cdc28p kinase activation into the GAL1::CLB5-Dlpb strain relieved lethality (S). Deletion of SIC1 made expression of GAL1::CLB5-KA,EA-Dlpb lethal (data not shown), again confirming that destruction box-dependent degradation and Sic1p inhibition can coregulate Clb5p activity, probably specifically in mitosis. GAL1::CLB5-KA,EA was not lethal even in the simultaneous absence of sic1 and swel (data not shown). The KA,EA mutation does not significantly affect degradation rates of Clb5p with or without its destruction box (data not shown).

Characterization of GAL1::CLB5-Dlpb lethality. GAL1::CLB5-Dlpb cells grown in raffinose medium arrest after several hours of galactose induction. Arrest is associated with lethality in that the plating efficiency of these cells on glucose medium drops approximately 1,000-fold by 4 to 6 h of galactose incubation (see Fig. 8C and 9C). After approximately 2 h of expression, the GAL1::CLB5-Dlpb cells appear to delay as binucleate large budded cells with a 2C DNA content, resembling a GAL1::CLB2-Dlpb-induced arrest (data not shown), but cells escape this mitotic block over the succeeding 2 h. By 4 to 6 h, the cells are heterogeneous with respect to DNA content: approximately half of the cells have a 2C (replicated) DNA content, and almost half have only 1C (Fig. 8B and 9B). There is usually some accumulation of cells that appear to have intermediate DNA contents. The cells are predominately large budded, and frequently they rebud after 4 h of galactose induction (Table 1). Tubulin staining by indirect immunofluorescence indicates that most of the cells are arresting with postmitotic spindles (Fig. 6). The DNA signals (DAPI) are heterogeneous in strength, compared to those of wild-type controls, possibly the result of an unequal distribution into the mother and daughter buds (Fig. 6). Combined with the FACs analysis showing a significant population of cells with approximately 1C DNA content, these results suggest that some cells have undergone abortive mitosis despite failure of DNA replication (S).

We interpret this phenotype as indicating that Clb5p-Dlpbp overexpression delays completion of mitosis, but many cells nevertheless ultimately divide in the presence of Clb5p-Dlpbp. These cells do not efficiently replicate DNA after division, accounting for the accumulation of cells with 1C DNA content. The GAL1::CLB5-Dlpb rebudding phenotype (Table 1) may be due to the initiation of a G1 cell cycle program without properly completing the later stages of mitosis and cytokinesis. The rebudding phenotype was significantly reduced when the cells were processed for indirect immunofluorescence. This most likely results from digestion of the cell wall during sample preparation and could indicate that cytokinesis (but not cell...
separation) is largely complete in many of the apparently rebudded cells.

Comparison between the GAL1::CLB5-Δdb and GAL1::CLB2-Δdb phenotypes. GAL1::CLB5-Δdb was reported to cause uniform arrest in late mitosis with replicated DNA (43), unlike the GAL1::CLB5-Δdb phenotype we observed; these cells also did not rebud, unlike the GAL1::CLB5-Δdb cells (Table 1). These differences could be due to intrinsic differences in the ability of Clb2p and Clb5p to block exit from mitosis; alternatively, levels of expression could differ between the two GAL1::CLB-Δdb constructs. Therefore, we replaced the CLB5 coding sequence with the CLB2 coding sequence in the GAL1::CLB5-Δdb construct. We found that these constructs yielded comparable levels of Clb protein and associated kinase activity (Fig. 7). As reported previously (43), GAL1::CLB2-Δdb expression results in late mitotic arrest, with few or no cells with unreplicated DNA detected (Fig. 8) and almost all cells displaying elongated spindles with the replicated DNA separated into the mother cell and the bud (Fig. 6). This phenotype was similar with or without the HA tag on GAL1::CLB2-Δdb (data not shown). In addition, the rapid induction of inviability by GAL1::CLB5-Δdb was not observed with GAL1::CLB2-Δdb (Fig. 8). The multiple-budding phenotype of GAL1::CLB5-Δdb was also not observed with GAL1::CLB2-Δdb (Table 1). Thus, even at comparable levels of expression,

### Table 1. Budding and rebudding percentages resulting from GAL1-induced CLB2 and CLB5 constructs

<table>
<thead>
<tr>
<th>Genotype</th>
<th>% of cells</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>Expt 1</td>
</tr>
<tr>
<td></td>
<td>Budded</td>
</tr>
<tr>
<td>Set I</td>
<td></td>
</tr>
<tr>
<td>Wild type</td>
<td>67</td>
</tr>
<tr>
<td>GAL1::CLB2</td>
<td>59</td>
</tr>
<tr>
<td>GAL1::CLB2-Δdb</td>
<td>88</td>
</tr>
<tr>
<td>GAL1::CLB5</td>
<td>67</td>
</tr>
<tr>
<td>GAL1::CLB5-Δdb</td>
<td>56</td>
</tr>
<tr>
<td>Set II</td>
<td></td>
</tr>
<tr>
<td>Wild type + vector</td>
<td>54</td>
</tr>
<tr>
<td>GAL1::CLB5-Δdb + vector</td>
<td>68</td>
</tr>
<tr>
<td>GAL1::CLB5-Δdb + pGAL1::CLB2</td>
<td>64</td>
</tr>
<tr>
<td>GAL1::CLB5-Δdb + pGAL1::CLB2-Δdb</td>
<td>54</td>
</tr>
</tbody>
</table>

*Percentages of budded and rebudded cells were determined by counting 200 cells for each sample in two separate experiments. The samples were processed by the methods described for Fig. 7 (set I) and 8 (set II) following 6 h of galactose induction. Budded cells represent only singly budded cells, and rebudded cells represent all variations of multiply budded cells (typically either three or four cell bodies).
the phenotypes due to overexpression of stabilized Clb2p and Clb5p are different.

We also carried out experiments in which strains were synchronized in G1 by using α-factor and then released into galactose medium to induce GAL1::CLB2Δdb or GAL1::CLB5Δdb expression. For reasons that we do not understand, in this protocol, GAL1::CLB5Δdb expression resulted in a long preanaphase delay with apparently fully replicated DNA (data not shown); such a delay was not detected in the experiments (Fig. 8) in which galactose was added to asynchronous cultures. This difference makes it hard to directly compare results between the two protocols. Despite this, a significant population of the GAL1::CLB5Δdb cells eventually escaped this block and divided their nuclei, and, overall, about 30% of the cells then rebudded (data not shown). In contrast, the GAL1::CLB2Δdb cells arrested stably late in mitosis with divided nuclei and without rebudding, as was seen in the experiments where galactose was added to asynchronous cultures (Fig. 6 and 8).

These results suggest that a new cell cycle is being initiated despite the presence of overexpressed Clb5p, but that this is blocked by overexpressed Clb2p, consistent with the results when galactose was added to asynchronous cultures (Fig. 8).

Clb5p is inefficient at blocking mitotic exit. Clb2p is probably the major B-type cyclin active in mitosis (26, 27, 43). Previous studies have shown that Clb2p-associated kinase activity must be eliminated for completion of division and for proper loading of DNA replication origins during G1 (10, 15, 43). Clb2p can be stabilized by Clb2p-Cdc28p or Clnp-Cdc28p (1). If Clb5p-Cdc28p also stabilizes Clb2p, then some aspects of the phenotype due to overexpression of CLB5Δdb could be indi-
rect, due to stabilized Clb2p-associated kinase activity. To test this, we constructed GAL1-CLB5 clb2::LEU2 strains, with or without the destruction box and with or without the KA,EA mutation in CLB5. clb2 deletion enhanced inviability due to CLB5, since GAL1::CLB5 and GAL1::CLB5-KA,EA-D were lethal in a clb2 background. clb2 deletion also enhanced the speed of induction of irreversible lethality due to GAL1::CLB5-Ddb, although this effect was somewhat variable (data not shown). In almost all cases, GAL1::CLB5-Ddb induction in clb2 strains resulted in a significantly greater accumulation of cells with 1C DNA content than was observed in CLB2 strains (data not shown).

These results suggested that Clb2p might be restraining mitotic exit in the CLB5-Ddb overexpressers. We constructed a strain expressing both GAL1::CLB5-Ddb and GAL1::CLB2-Ddb. The strain arrested as large budded cells with a 2C DNA content (Fig. 9) where the replicated DNA was separated into the mother and daughter cell bodies (data not shown), typical of a GAL1::CLB2-Ddb-induced arrest. This is consistent with the idea that the accumulation of 1C DNA content in cells with GAL1::CLB5-Ddb alone requires mitosis, which may be blocked by GAL1::CLB2-Ddb expression. Similarly, GAL1::CLB2-Ddb expression significantly reduces the number of cells displaying the characteristic GAL1::CLB5-Ddb rebudding phenotype (Table 1), consistent with the inhibitory effects of Clb2p on cell polarization and bud emergence reported previously (2, 23). Although the reversibility of the arrest in the GAL1::CLB5-Ddb/GAL1::CLB2-Ddb strain was reduced from that observed in GAL1::CLB2-Ddb cells, viability was increased 10-fold from strains expressing GAL1::CLB5-Ddb alone (Fig. 9). These results support the idea that Clb2p can restrain mitotic exit in GAL1::CLB5-Ddb expressers and that the severity of GAL1::CLB5-Ddb-induced lethality is correlated with an accumulation of cells containing a 1C DNA content. Accumulation of 1C cells may not be the sole cause of irreversible arrest, because GAL1::CLB5-Ddb-expressing cells arrest with a heterogeneous population of DNA content. This is difficult to interpret fully, though, because these cells may ultimately divide upon plating for the viability assay.

**DISCUSSION**

**Clb5p degradation.** We find that Clb5p degradation is cell cycle regulated, and Clb5p is most unstable in dividing cells. These differences in stability correlate with loss of detectable nuclear accumulation of Clb5p in dividing cells (although significant cytoplasmic signal remains). We speculate that the specific loss of Clb5p from the nucleus may be due to nucleus-localized degradation, but further work is required to eliminate other possibilities (for example, that Clb5p is exported from the nucleus during mitosis and is independently rendered less stable at this time).

Clb5p-Dbp expressed from the CLB5 promoter has at best minor phenotypes, even in the absence of sic1, and destruction box deletion from endogenously expressed Clb5p also results in, at most, minor stabilization of the protein, monitored with HA-tagged Clb5p in synchronous culture (Fig. 3 and data not shown). Destruction box-independent degradation of Clb5p...
may be responsible, or there may be other unidentified destruction boxes in Clb5p. In either case, the overexpression of Clb5p from the GAL1 promoter may swamp out these alternative means of degrading Clb5p.

Cyclin specificity. The six CLB B-type cyclins have distinct roles in vivo, although they overlap significantly in function. Distinct in vivo roles could be due to time of expression during the cell cycle or to intrinsic specialization of the different CLB coding sequences. Previously, we showed that Clb5p was much more potent at inducing DNA replication than Clb2p, even when time of expression and protein accumulation were made comparable, by placing both under control of the CLB5 promoter (8). Here, we examine the cell cycle-inhibitory activity of Clb5p compared to that of Clb2p. To make this comparison, we eliminated differential transcriptional control by using the GAL1 promoter and eliminated differential proteolytic control by removing the proteins’ destruction boxes. We find that while Clb2p is able to block exit from mitosis, as reported previously, Clb5p is relatively weak at this activity, even when strongly overexpressed. Even the limited apparent capacity of Clb5p to inhibit mitotic exit is likely to be dependent on endogenous Clb2p. Clb5p could recruit endogenous Clb2p for this role by stabilizing it, perhaps by phosphorylation of Cdh1p (1, 49). In addition, the potency of existing Clb2p may be increased due to Clb5p-dependent phosphorylation of Sic1p (46). The idea that Clb5p restrains mitotic exit by phosphorylation of Cdh1p and Sic1p was also suggested by Shirayama et al. (40).

Accumulation of cells with unreplicated DNA in GAL1::CLB5Δdb cells may be due to failure to license replication origins due to high Clb5p-associated kinase (9), combined with the permissiveness of high Clb5p-associated kinase for completion of mitosis. The consequence may be that cells divide without being able to initiate DNA replication after division. If cells under these conditions pass the “point of no return” (31) when endogenous Clb-associated kinases block reloading of replication origins, then even after shutoff of GAL1::CLB5-Δdb, irreversible lethality is predicted. It is also possible that some of these cells undergo anaphase without DNA replication, leading to unequal segregation of the haploid DNA content (31), a clearly lethal event.

The molecular basis for differences in cyclin specificity is unknown. The results reported here and previously (8, 34) do not suggest that differential accumulation of the protein or associated kinase is responsible, although subtle differences in timing or levels are hard to rule out unambiguously. A candidate substrate-targeting domain (the hydrophobic patch [8, 34]) could function differently between Clb5p and Clb2p, potentially leading to differential substrate targeting. This region contributes to but is not essential for lethality of GAL1::CLB5-Δdb (8). The analogous region in Clb2p may be required for efficient blocking of mitotic exit in GAL1::CLB5-Δdb, GAL1::CLB2-Δdb overexpressors in the assay shown in Fig. 9 (data not shown). Failure to detect a strong requirement for this region for some effects of Clb-Δdb overexpression may be due to masking the role for the region due to high expression levels. The region is required for efficient lethality of Clb5p in the absence of cdc20 and pds1 and is also required for efficient Clb2p mitotic function at lower levels of expression (7).

S and M cyclins and the organization of the yeast cell cycle. CLB5 and CLB6 have been called S-phase cyclins and CLB1 and CLB2 have been called M-phase cyclins, due to their time of expression and evident function as deduced from null phenotypes (12, 15, 26, 27, 37, 43). In fission yeast, cdc13 appears to be the predominant M-phase cyclin, and cig1 (with help from cig2) may be the major S-phase cyclin. Despite this differentiation of function among B-type cyclins, models have been proposed for both budding yeast and fission yeast in which a single generic B-type cyclin-dependent kinase activity could be sufficient to control both the S and M phases in proper alternation (13, 26, 42). These models suggest that functional differences in the roles of different B-type cyclins (as deduced from null phenotypes) are due solely to differential accumulation due to transcriptional or proteolytic controls. At least for budding yeast, this appears to be an oversimplification. Previously we showed that Clb5p is intrinsically specialized for induction of DNA replication, functioning much better
at this activity than Clb2p (8). The results reported here suggest that Clb2p has a specific function of restraining mitotic exit, which is weak or absent in Clb5p. This difference is not due to defects in Clb5p protein accumulation or kinase activation, and thus the difference is likely to be intrinsic to the protein. Thus, at least for these two activities, one early in the cell cycle and one at the end, the yeast cell cycle is driven by a single B-type cyclin capable of driving events in DNA replication and in mitosis. In metazoans, it is likely that cyclins A and E are specialized for induction of S phase and that B-type cyclins are specialized for induction of mitosis, and this situation may be closer to the budding yeast system when cyclins are expressed at different levels.

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