Nucleolar protein PinX1p regulates telomerase by sequestering its protein catalytic subunit in an inactive complex lacking telomerase RNA

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Human TRF1-binding protein PinX1 inhibits telomerase activity. Here we report that overexpression of yeast PinX1p (yPinX1p) results in shortened telomeres and decreased in vitro telomerase activity. yPinX1p coimmunoprecipitated with yeast telomerase protein Est2p even in cells lacking the telomerase RNA TLC1, or the telomerase-associated proteins Est1p and Est3p. Est2p regions required for binding to yPinX1p or TLC1 were similar. Furthermore, we found two distinct Est2p complexes exist, containing either yPinX1p or TLC1. Levels of Est2p–yPinX1p complex increased when TLC1 was deleted and decreased when TLC1 was overexpressed. Hence, we propose that yPinX1p regulates telomerase by sequestering its protein catalytic subunit in an inactive complex lacking telomerase RNA.

[Keywords: Telomerase regulation; telomerase sequestration; alternative telomerase complex; PinX1–telomerase protein complex]

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Telomerase is required for complete replication of chromosomal ends and for genome stability (Blackburn 2000). Without telomerase, loss of telomeric sequence due to incomplete DNA replication and nuclease action leads to telomere shortening and eventual cellular senescence. The core telomerase enzyme is minimally composed of two subunits: the reverse transcriptase protein TERT (Est2p in Saccharomyces cerevisiae) and the template-containing RNA component TER (TLC1 in S. cerevisiae). These two components from various organisms can reconstitute in vitro enzymatic activity (Weinrich et al. 1997; Beattie et al. 1998; Licht and Collins 1999). Although additional proteins are required for telomerase to act in vivo (Blackburn 2000; Blasco 2002), little is known about other factors that may regulate telomerase activity.

Telomerase action on telomeres is regulated during the cell cycle and developmentally. Studies of telomerase in S. cerevisiae demonstrated that the enzyme is most active on shortened telomeres in late S phase or G2/M phase, the time when telomeres are replicated (Diede and Gottschling 1999; Marcand et al. 2000). In the ciliate Euplotus crassus, telomerase complexes with distinct biochemical and structural properties can be purified during different developmental stages (Greene and Shippen 1998). Telomerase activity is constitutively expressed in human germ-line and other dividing cells, but is diminished in many normal somatic cells. The vast majority of primary tumors and tumor-derived cell lines have robust telomerase activity, whereas inhibition of telomerase or interference with its action limits the growth of human cancer cells (Blasco and Hahn 2003). Thus, stimulation of telomerase activity in cancer cells is a key step toward tumorigenesis.

Telomerase activity can be regulated at multiple levels. In yeast, although Est2p protein and TLC1 RNA are present throughout the whole cell cycle, addition of telomeric DNA is restricted to late S phase or G2/M phase. It was proposed that telomerase is only activated during these phases of the cell cycle (Marcand et al. 2000; Taggart et al. 2002, Taggart and Zakian 2003). A similar model was suggested for human telomerase (Loayza and De Lange 2003). TERT expression can be regulated both transcriptionally and posttranscriptionally. In E. crassus, three different TERT variants with different expression profiles have been found, and a switch between different catalytic subunits was proposed to occur during development (Karamysheva et al. 2003). Although hTERT transcription is low in many human tissues, it is com-
monly reactivated in tumors (Blasco and Hahn 2003). In addition, posttranslational phosphorylation of hTERT has been observed during T-cell activation (Liu et al. 2001). Translocation of telomerase between cytoplasm and nucleus, and between nucleolus and nucleoplasm, have also been proposed as activity regulation mechanisms (Seimiya et al. 2000; Liu et al. 2001; Wong et al. 2002).

Sequestration into alternative complexes is a known mechanism for regulating the function of many proteins. For example, NF-κB, a protein important for inflammatory, immune, and antiapoptotic responses in mammals (Ghosh et al. 1998), is sequestered in the cytoplasm by binding to a family of inhibitory proteins, IκBs. NF-κB is released upon ubiquitination of IκBs and subsequently translocates to the nucleus to function as a transcriptional activator (Baldwin 1996). Similarly, separase, the protease that cleaves the chromosomal cohesion complex, is regulated by its binding partner securin, which directly inhibits the proteolytic activity of separase by preventing separase from binding its substrates (Concannon et al. 2001; Hornig et al. 2002; Waizenegger et al. 2002).

The possibility that telomerase might be regulated by sequestering TERT in an alternative complex has not been previously explored. Here we report evidence for a novel molecular mechanism for telomerase regulation, in which the yeast telomerase catalytic protein can form an inactive complex that contains nucleolar protein yPinX1p but not the essential RNA component of the telomerase enzyme. Human PinX1p, initially identified as a protein that interacts with the telomeric protein TRF1/Pin2, was previously reported to bind to hTERT and directly inhibit telomerase activity in vitro assays (Zhou and Lu 2001). Depletion of hPinX1p increases tumorigenicity in a mouse host, suggesting that hPinX1p might be a tumor suppressor (Zhou and Lu 2001). However, the molecular mechanisms of PinX1p action were unknown.

The yeast homolog of PinX1p (also called Gno1p, G- patch nucleolar protein 1) is an essential protein involved in rRNA and snRNA maturation (Guglielmi and Werner 2002). yPINX1 deletion cells have poor growth with a doubling time of ∼4.9 h compared with 1.5 h for wild-type cells (Guglielmi and Werner 2002). Guglielmi and Werner (2002) previously concluded, because telomeres in ΔyPINX1 cells were slightly shorter than those in wild-type cells, that yPinX1p is not a telomerase inhibitor, unlike its human counterpart.

Here we show that overexpression of yPinX1p causes telomere shortening and decreased in vitro telomerase enzymatic activity. We provide evidence that the same central region of Est2p is required for binding of yPinX1p or TLC1 RNA. A new in vivo Est2p complex, containing yPinX1p but not TLC1, was detected. The level of this Est2p–yPinX1p complex was increased in a Δtlc1 strain and decreased upon TLC1 overexpression, indicating that yPinX1p competes with TLC1 for binding to Est2p. Taken together, we propose that PinX1p inhibits telomerase activity by sequestering the protein reverse transcriptase subunit into an inactive complex.

**Results**

**Overexpression of yeast PinX1p shortens telomeres and decreases in vitro telomerase activity**

To determine whether yeast PINX1 plays a role in telomere maintenance, we assayed the effects of yPinX1p overexpression on telomere length. We cloned yPINX1 under the GAL1 promoter on a CEN/ARS plasmid. To determine the yPinX1p overexpression level, we tagged both the endogenous yPINX1 gene and the GalPINX1 plasmid with 13 myc tags at the C termini and compared the expression levels by Western blotting using an antimyc antibody. We found that when induced with galactose, yPinX1p was expressed approximately fourfold over its endogenous level (data not shown). Wild-type yeast cells, transformed with the yPINX1 overexpression plasmid or the control vector lacking the yPINX1 ORF, were grown under noninducing (glucose) or inducing (galactose) conditions, and telomere length was examined by Southern blotting. Telomeres in cells overexpressing yPinX1p were ∼70–80 bp shorter than telomeres in controls cells [Fig. 1A, cf. V and P lanes]. This shortening was not caused by a difference in the medium, because telomere length was identical in control cells grown in glucose or galactose medium [Fig. 1A, cf. V lanes + and −]. Telomere shortening was visible by 25 generations of growth on galactose [data not shown], and became more pronounced at the third streak, as shown in Figure 1A. Therefore, these data suggested a role for yPINX1 in telomere maintenance.

Telomere shortening upon yPINX1 overexpression could potentially result from decreased telomerase enzymatic activity per se, or from decreased telomerase action caused by structural and/or conformational changes at the telomeres. In yeast, telomere length is negatively regulated by Rap1p and its binding partners Rif1p and Rif2p. It was proposed that Rap1p binds to telomeric DNA in a sequence-specific manner and recruits Rif1p, Rif2p, and other proteins to form a higher-order complex that limits access of telomerase to telomeres (Marcand et al. 1997). Telomeres become elongated when Rif1 or Rif2 is deleted (Hardy et al. 1992, Wotton and Shore 1997). To determine whether the telomere shortening by yPinX1p overexpression requires Rif1p or Rif2p, we overexpressed yPinX1p in Rif1 or Rif2 deletion strains. If the telomere shortening effect of overexpression of yPinX1p is mediated through the Rif proteins, such shortening should be alleviated by deletion of Rif1p or Rif2p. However, in both Δrif1 and Δrif2 strains, significant telomere shortening was still observed upon yPinX1p overexpression [Fig. 1B]. Hence, telomere shortening caused by overexpression of yPinX1p is not solely dependent on either Rif1p or Rif2p, although some role for either protein cannot be completely excluded.

We overexpressed telomerase protein subunit Est2p in addition to yPinX1p. Expression of Est2p under its endogenous promoter from a high-copy 2µ plasmid caused slight lengthening of telomeres [Fig. 1C, left part, cf. V and P + E lanes when PinX1p was not overexpressed].
Interestingly, telomere length was restored to wild-type upon overexpression of both Est2p and yPinX1p together, consistent with the possibility that yPinX1p influences telomerase enzymatic activity directly [Fig. 1C, cf. P and P + E lanes under yPinX1p overexpression conditions]. This possibility was confirmed by comparison of in vitro telomerase activity in extracts of cells overexpressing yPinX1p to those containing only the vector control. Telomerase was partially purified by DEAE chromatography as previously described (Cohn and Blackburn 1995). To make quantitative comparisons, the TLC1 peak fraction was used, and each assay contained equal amounts of TLC1 RNA. Cells overexpressing yPinX1p had a decreased in vitro activity (33% of the control, after normalization and correction for background signal; see Fig. 1D). Taken together, these results show that overexpression of yPinX1p decreases telomerase enzyme activity and telomere length in yeast cells.

yPinX1p associates with Est2p in vivo

Because yPinX1p is involved in rRNA and snoRNA processing, we first tested whether overexpression of yPinX1p alters the levels of two RNAs involved in telomerase activity: TLC1 RNA and EST2 mRNA. Induction of yPinX1p overexpression did not change either the TLC1 RNA or the EST2 mRNA levels [Supplemental Fig. S1A]. Furthermore, Western blotting analysis of Est2p tagged with 13 tandem myc epitopes [Est2p-myc] expressed from its endogenous locus showed no change in the total Est2p protein level upon yPinX1p overexpression [Supplemental Fig. S1B]. However, despite the same levels of total Est2p-myc in whole cell extracts, we found that the amount of Est2p-myc immunoprecipitable with anti-myc antibody was significantly decreased in the cells overexpressing yPinX1p, compared with the vector control cells [Supplemental Fig. S1C]. These results indicated that even though yPinX1p overexpression does not alter the total levels of Est2p, the myc epitopes on Est2p became less accessible to antibody binding.

Human PinX1p has been shown to interact with hTERT both in vitro and in vivo (Zhou and Lu 2001). We therefore examined whether yPinX1p is associated with Est2p in vivo by coimmunoprecipitation. yPinX1p was tagged with the TAP tag (Puig et al. 2001), and Est2p was tagged with the 13-myc tag in strain yEHB4114, with both proteins expressed from their endogenous chromosomal loci. Strain yEHB4008, containing myc-tagged Est2p, but untagged yPinX1p, was used as the control. Both strains grew as healthily as a wild-type strain and had normal telomere length (data not shown), indicating that the tagged proteins were fully functional. Est2p-myc copurified with yPinX1p-TAP on IgG beads, which bind to the ZZ domain in the TAP-tag (Puig et al. 2001), but was not copurified in the control strain that contained the untagged yPinX1p [Fig. 2A], suggesting that the two proteins associate in vivo. To further confirm the interaction between Est2p and yPinX1p, we performed the same experiment in strains with Est2p-TAP-tagged and yPinX1p-myc-tagged. As with the previous tagged strains, the Est2p-TAP/yPinX1p-myc [yEHB4109] and yPinX1p-myc [yEHB4107] strains had normal growth and telomere length [data not shown]. Consistent with the result above, yPinX1p-myc also copurified with Est2p-TAP on IgG beads [Fig. 2B]. Taken together, these results suggest that yPinX1p binds, directly or indirectly, to Est2p in vivo.

yPinX1p has previously been shown to be involved in rRNA and snoRNA processing (Guglielmi and Werner 2002). Because telomeres in ΔypinX1 cells were slightly shorter than those in wild-type cells, it was concluded that yPinX1p is not a telomerase inhibitor, unlike its human counterpart [Guglielmi and Werner 2002]. However, such slight shortening might have been explained by a pleiotropic effect of the defective processing of rRNA and snoRNA in the deletion strain. We were able to uncouple the effects of yPinX1p on telomere length from its role in snoRNA and rRNA processing. A point mutant, yPinX1p-W38A, does not support cell growth when the wild-type yPINX1 gene is replaced by it, pre-
sumably because of its defects in rRNA and snoRNA processing [data not shown]. In agreement with this, mutation of the same amino acid residue to serine, W385, was reported to be defective in rRNA and snoRNA processing [Guglielmi and Werner 2002]. Notably, however, when overexpressed, yPinX1p-W38A was still capable of causing telomere shortening to the same degree as the wild-type yPinX1p. Furthermore, yPinX1p-W38A still copurified with Est2p [Supplemental Fig. S2]. Therefore, the functions of yPinX1p in rRNA/snoRNA processing and telomere maintenance are separable.

**Est2p associates with yPinX1p in the absence of TLC1, Est1p, or Est3p**

In addition to Est2p and TLC1, yeast telomerase holoenzyme contains at least two protein components that are essential for its function in vivo, Est1p and Est3p. Est1p binds to TLC1 and telomeric DNA and is proposed to recruit/activate telomerase at the telomeres [Evans and Lundblad 1999, 2002, Seto et al. 2002, Taggart et al. 2002]. Est3p is another component of telomerase holoenzyme whose biochemical function remains unknown [Hughes et al. 2000]. We found that the association between Est2p and yPinX1p is not dependent on TLC1, Est1p, or Est3p. This was shown by coimmunoprecipitation experiments in strains singly deleted for each one of these three genes [see Table 1 for strains used in this study]. EST2 was cloned into a 2µ plasmid together with its endogenous promoter and ADH terminator, and was tagged at its C terminus with 13 myc epitopes. This EST2 plasmid (pRS424Est2myc), together with a plasmid expressing TAP-tagged yPinX1p from the GAL1 promoter [pRS316GalPinX1TAP, see Table 2], was transformed into yeast strain YEH4084, which is deleted of its endogenous EST2 and maintains its telomeres by the alternative Rad52-dependent recombination pathway [Lundblad and Blackburn 1993]. We immunopurified yPinX1p-TAP in two steps with IgG beads and calmodulin beads using the published protocols for TAP-tag purification [Puig et al. 2001] and then performed Western blot analysis to detect Est2p-myc. As with the endogenously expressed tagged Est2p and yPinX1p, wild-type Est2p-myc overexpressed from the 2µ plasmid was readily detectable in the purified fractions [Fig. 3A]. Furthermore, wild-type Est2p-myc still efficiently copurified with yPinX1p-TAP in TLC1, EST1, or EST3 deletion strains [Fig. 3B]. Hence, Est2p-myc associates with yPinX1p independently of TLC1 RNA, Est1p, or Est3p.

The same region in Est2p is required for yPinX1p and for TLC1 binding

The Est2p protein contains six recognizable conserved domains [Fig. 4A] found in other TERTs [Xia et al. 2000]. The Q domain in yeast Est2p is required for its binding to Est3p, another component of the yeast telomerase holoenzyme [Hughes et al. 2000, Friedman

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**Figure 2.** Coimmunoprecipitation of yPinX1p and Est2p. (A) Coimmunoprecipitation of Est2p-myc with yPinX1p-TAP using IgG beads. (B) Coimmunoprecipitation of yPinX1p-myc with Est2p-myc using IgG beads. For Western blotting (WB), 2% of the extract used for coimmunoprecipitation experiments was loaded in whole-cell extract lanes.

**Table 1. Strains used in this study**

| Strain       | Genotype                                      |
|--------------|-----------------------------------------------|
| VY4705       | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 |
| yEHB0234     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 rif1Δ:TRP1 |
| yEHB0235     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 rif2Δ:KAN |
| yEHB4008     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 est2::EST2-13myc-KanMX6 |
| yEHB4084     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 est2::HIS3 |
| yEHB4100     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 est2::HIS3 tlc1Δ:KAN |
| yEHB4104     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 est2::HIS3 tlc1Δ::TRP1 |
| yEHB4107     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 pinX1::PINX1-13myc-KanMX6 |
| yEHB4109     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 pinX1::PINX1-13myc-KanMX6 |
| yEHB4114     | MATa ade2Δ:hisG his3Δ200 leu2Δ0 lys2Δ0 met15Δ0 trp1Δ63 ura3Δ0 est2::EST2-13myc-KanMX6 |
| yEHB4120     | MATa ade2Δ:hisG his3Δ200 met15Δ0 trp1Δ63 ura3Δ0 est1Δ::HIS3 |
| yEHB4121     | MATa ade2Δ:hisG his3Δ200 met15Δ0 trp1Δ63 ura3Δ0 est3Δ::HIS3 |

*These strains are survivors.
et al. 2003). The CP, QFP, and T domains of telomerases are involved in binding to TER in various organisms [Friedman and Cech 1999; Bryan et al. 2000; Armbruster et al. 2001; Lai et al. 2001; Moriarty et al. 2002b]. The RT domain shares sequence homology with other reverse transcriptases and contains amino acid residues that directly participate in catalysis [Nakamura et al. 1997]. The function of the very N-terminal region is unclear. Deletional and mutational analyses have shown that all six regions are necessary for normal telomerase function [Kelleher et al. 2002].

To determine which region in Est2p is required for its interaction with yPinX1p in vivo, we tested six EST2 mutants, each deleting one single conserved domain as indicated in Figure 4A (see also Table 2). Each EST2 domain mutant was cloned into a 2µ plasmid with the endogenous EST2 promoter and the ADH terminator, and tagged with 13-myc epitopes at the C terminus. Replacement of the wild-type EST2 gene by each mutant EST2 plasmid caused telomere shortening and senescence at the same rate as an EST2 deletion [data not shown]. We examined whether each EST2 domain deletion mutant protein was stably expressed by Western blot analysis using an anti-myc antibody. Whereas Est2pΔN-myc was expressed at a lower level than the wild type, the other five mutants were at levels similar to wild type (Fig. 4C).

To determine whether these mutant EST2 proteins could copurify with yPinX1p-TAP, each EST2 domain deletion mutant plasmid was transformed into yeast strain yEB4084, together with plasmid pRSGALPinX1-TAP. We immunopurified yPinX1p-TAP and performed Western blot analysis to detect Est2p-myc. Three of the six mutants—ΔCP, ΔQFP, and ΔT—which delete the central region of Est2p, failed to copurify with yPinX1p-TAP [Fig. 4B], whereas the other three mutants—ΔN, ΔGQ, and ΔRT—still copurified with yPinX1p-TAP [Fig. 4B]. Thus, the central region of Est2p encompassing the CP, QFP, and T domains is required for yPinX1p association. Interestingly, immunoprecipitated ΔCP, ΔQFP, and ΔT mutant proteins also failed to pull down TLC1 RNA, in contrast to the other three mutants [Fig. 4C]. This result is consistent with previous studies of human and Tetrahymena TERT proteins, where deletion or mutation of the CP, QFP, and T domains each reduced or abolished TER binding [Armbruster et al. 2001; Lai et al. 2001; Moriarty et al. 2002b]. Thus, the regions in Est2p necessary for either TLC1 or yPinX1p binding overlap.

Two distinct Est2 complexes contain either TLC1 or yPinX1

The above results showed that association of Est2p with yPinX1p or TLC1 required the same Est2p region. To determine whether these three molecules are in the same or different complexes, we fractionated telomerase using the previously reported DEAE chromatography method [Prescott and Blackburn 1997a,b] from a yeast strain expressing wild-type Est2p-myc from a 2µ plasmid and yPinX1p-myc from its endogenous chromosomal locus. Cells were lysed, and the S100 extract was loaded onto a DEAE column. As shown by Northern blot analysis, the vast majority of TLC1 bound to the column, with no TLC1 detected in the flowthrough (Fig. 5A). However,
Western blot analysis showed that a significant portion of Est2p was in this flowthrough (Fig. 5A), suggesting that a fraction of Est2p is TLC1-free. TLC1 was eluted, and its concentration peaked in fraction 5 (Fig. 5A, bottom panel). Therefore, active telomerase complex containing Est2p and TLC1 RNA does not contain detectable levels of yPinX1p.

Next, we used an independent method to test whether TLC1 RNA is present in the yPinX1p–Est2p complex. TAP-tagged yPinX1p was purified from yeast using IgG beads as described above. As shown above, Est2p-myc was found to copurify with yPinX1p (Fig. 5B, bottom left panel). However, no TLC1 was detected in the yPinX1p-TAP pull-down (Fig. 5B, top left panel). As a positive control, the same extract was immunoprecipitated with anti-myc antibody to pull down Est2p-myc, and TLC1 RNA was readily detected (Fig. 5B, right panels). Taken together, these results show that there are at least two distinct Est2p complexes: the enzymatically active telomerase containing Est2p and TLC1 (possibly with other proteins), which does not contain yPinX1p, and an inactive complex containing Est2p–yPinX1p, but not TLC1 RNA.

Evidence that active telomerase is a dimer with two active sites and two interacting RNAs has been reported previously (Prescott and Blackburn 1997a, b; Beattie et al. 2001; Wenz et al. 2001; Ly et al. 2003). In human telomerase, TERT itself is known to interact via TERT–TERT protein–protein binding in vitro (Arai et al. 2002). Using coimmunoprecipitation experiments with cells expressing HA-tagged Est2p from one allele and myc-tagged Est2p from the other allele, we found that Est2p–Est2p in vivo association could still occur even in strains in which the other essential telomerase holoenzyme components Est1p, Est3p, or the telomerase RNA gene (TLC1) were deleted (Supplemental Fig. S3). This property of independence from other telomerase holoenzyme components was shared by the Est2p–yPinX1p association described above. Therefore, we delineated the domains of Est2p that are required for this Est2p–Est2p interaction in vivo, by coexpressing the wild-type Est2p (tagged with 13×myc) and each of the Est2p single-domain deletion mutants (tagged with 3×HA), both from the endogenous promoter on 2µ plasmids. Deletion of the T or RT domain, but not of the CP or QFP domains, significantly disrupted the in vivo physical association between Est2p molecules (Supplemental Fig. S4). Hence the region(s) of Est2p required for its physical Est2p–Est2p interaction in vivo overlaps with, but is not identical to, the common region required for the mutually exclusive PinX1p or TLC1 binding. These results suggest the possibility that yPinX1p binding may also interfere with telomerase dimerization.
**TLC1 levels influence yPinX1–Est2p binding efficiency**

The above results showed that Est2p exists in alternative complexes, containing either yPinX1p or TLC1 RNA. The overlap of the Est2p regions required for binding of PinX1p or TLC1 further suggested that PinX1p and TLC1 may compete for Est2p. Therefore, we tested whether TLC1 RNA levels influence the level of the yPinX1–Est2p complex. We determined the effects of TLC1 deletion or overexpression on the extent of coimmunoprecipitation of Est2p-myc with yPinX1p-TAP. TAP-tagged yPinX1p was expressed from a GAL1 promoter in strains containing either wild-type TLC1, TLC1 deletion (Δtlc1), or TLC1 overexpressed from a GAL1 promoter. Compared with the wild-type TLC1 strain, the total level of Est2p-myc protein slightly increased in the TLC1 overexpression strain, and reproducibly decreased dramatically in the Δtlc1 strain [Fig. 6A]. Strikingly, despite this lower level of total Est2p-myc, in the Δtlc1 strain more Est2p-myc was reproducibly coimmunoprecipitated by yPinX1p-TAP than in the control TLC1 wild-type strain [Fig. 6A]. In contrast, in the TLC1 over-

**Discussion**

The results of this study support a novel mechanism for telomerase activity regulation. Our results show that Est2p forms two distinct complexes: the active telomerase enzyme containing TLC1, and an alternative complex containing yPinX1p but no TLC1. Furthermore, we have provided evidence that yPinX1p competes with TLC1 for Est2p binding in vivo. Thus, we propose that yPinX1p sequesters Est2p to prevent assembly of Est2p and TLC1 RNA into the enzymatically active telomerase RNP [Fig. 6B].

**PinX1p: a dual-function protein**

yPinX1p is an essential nucleolar protein required for the processing of rRNA and some snoRNAs [Guglielmi and Werner 2002]. The poor growth in yPINX1 deletion cells is very likely caused by an inability to process rRNA and snoRNAs. The rRNA and snoRNA processing functions of PinX1p are conserved as the human PINX1 gene complemented the slow growth and rRNA maturation defects of the yeast deletion strain [Guglielmi and Werner 2002].

Because telomeres in ΔypinX1 cells were slightly shorter than in wild-type cells, a previous report concluded that yPinX1p is not a telomerase inhibitor, unlike its human counterpart [Guglielmi and Werner 2002]. However, such slight shortening might have been explained by a pleiotropic effect of the defective processing of rRNA and snoRNA in the deletion strain. In agreement with this, two yPinX1p mutants (W38S and Δ1–24) that caused slight shortening of telomeres are both defective for rRNA processing; whereas another mutant (Δ150–271) that retained rRNA/snoRNA functions, had wild-type telomere length [Guglielmi and Werner 2002]. Furthermore, as described above [see Results and Supplemental Fig. 2S], the functions of yPinX1p in rRNA/snoRNA processing and telomere maintenance are separable. We have shown here that yeast PinX1p binds to CP, QFP, and T regions of telomerase catalytic protein Est2p. Because these regions are conserved among all telomerase catalytic subunits of all species, a similar mechanism may exist for human PinX1p–TERT binding.

In support of a conserved role for PinX1-mediated telomerase regulation, we overexpressed human PinX1p in yeast and observed decreased telomere length (data not shown). Thus, we propose that yeast PinX1p, like its human counterpart, is also a dual-function protein involved in both rRNA/snoRNA processing and telomerase regulation.

**Role of the nucleolus in telomerase activity**

This proposed sequestration of telomerase into alternative complexes may, at least in part, underlie the trans-

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**Figure 6.** TLC1 levels influence yPinX1–Est2p binding efficiency. (A) Deletion of TLC1 results in more Est2p associated with yPinX1p, and overexpression of TLC1 results in less Est2p associated with yPinX1p. (O/E) TLC1 overexpressed from the GAL1 promoter. PGK serves as the loading control for Western blotting. The amount of TLC1 RNA is shown on the bottom with actin mRNA as the loading control. **(B)** A model for yPinX1p-mediated inhibition of telomerase activity. See text for description.
that this increases Est2p levels in the enzymatically active state, which is typically found in telomeres under conditions of environmental stress (Gasch et al. 2000, 2001). We suggest that increased levels of Est2p are a mechanism to regulate telomerase activity? PinX1p, like overexpressed Est2p, is predominantly a nucleolar protein, but may also be involved in a different level of regulation of in vivo telomerase activity (Seimiya et al. 2002).

The nucleolus has been strongly implicated in the biogenesis of telomerase and regulation of its activity. Mammalian telomerase RNA shares domains with small nuclear RNAs that bind to several known nucleolar proteins (Mitchell et al. 1999; Pogacic et al. 2000; Fu and Collins 2003). Recently, movement of hTERT between subnuclear compartments has been reported. hTERT in primary cells was shown to culture to localize in the nucleolus, but is apparently released into the nucleoplasm during the late S/G2 stages of the cell cycle, coincident with the replication of telomeres (Wong et al. 2002). In contrast, in transformed cells, hTERT has a diffuse nucleoplasmic distribution (Wonget al. 2002). Interestingly, the nucleolar localization of hTERT has been shown to interact with hTERT and prevent its nuclear export. This suggested that 14–3–3 proteins are required for the nuclear accumulation of hTERT, and may thus be involved in another level of regulation of in vivo telomerase activity (Zhou and Lu 2001; Lin and Blackburn 2003). Recently, movement of overexpressed Est2p correlated with changes in vitro telomerase activity, consistent with the interpretation that enzymatically active telomerase is in the nucleoplasm outside the nucleolus (Teixeira et al. 2002).

Under what physiological situations might PinX1p act to regulate telomerase activity? PinX1p, like overexpressed Est2p, is predominantly a nucleolar protein, but is also found in the nucleoplasm (Zhou and Lu 2001; Teixeira et al. 2002). Interestingly, the nucleolar localization domain of hTERT maps to amino acids 326–620 (Etheridge et al. 2002). The homologous region in Est2p includes the yPinX1p-interacting region we have reported here. We therefore propose that PinX1p is involved in controlling the movement of TERT between nucleolus and nucleoplasm. In support of this model, human telomerase is found in the nucleoplasm at the time during the cell cycle when telomeres are replicated, and also upon transformation, which promotes cell cycling. In addition, telomerase enzymatic activity detectable in cell extracts is very rapidly up-regulated in human leukemia HL-60 cells treated with clinical doses of the DNA-damaging drug etoposide (Morarity et al. 2002a). In S. cerevisiae, the level of yPINX1 mRNA decreases significantly upon DNA damage or general environmental stress (Gasch et al. 2000, 2001). We suggest that this increases Est2p levels in the enzymatically active complex under these conditions.

In summary, the results reported here, together with the above findings, suggest that release of Est2p from its sequestering PinX1p, to allow binding of telomerase RNA, may constitute an efficient and responsive mechanism to rapidly regulate telomerase.

Materials and methods

Southern blot analysis of telomeres

Cells were streaked on plates selective for the plasmids as specified. Genomic DNA was prepared and digested with XhoI and run on 0.8% agarose gels. DNA was transferred from gels to Hybond N+ membranes (Amersham) and probed with a γ-32P end-labeled wild-type telomeric repeat oligonucleotide as described previously (Prescott and Blackburn 1997b).

Immunoprecipitation and Western blot analysis

Cells were grown in selective media to an OD600 of 0.5–0.6, lysed with glass beads in IPP150 buffer (25 mM HEPES, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA, 0.1% NP-40, 10% glycerol, 1 mM DTT, and protease inhibitors). The lysates were centrifuged at 13,000 rpm for 15 min and 40,000 rpm for 30 min to obtain the extracts. Typically, 2–3 mg of total protein was immunoprecipitated with anti-myc antibody 9E10 (Covance) and protein A beads (Sigma) in a 400-μL total volume for 4 h, washed with IPP150, and resuspended in SDS sample buffer. Immunoprecipitation of TAP-tagged proteins was performed as reported using 100–150 μg of total protein (Puig et al. 2001). Briefly, extracts were incubated with IgG beads for 2 h at 4°C, and the beads were washed with IPP150 buffer and resuspended in TEV cleavage buffer (25 mM HEPES, 150 mM NaCl, 0.5 mM EDTA, 1 mM EGTA, 0.1% NP-40, 10% glycerol, 1 mM DTT). TEV protease (Invitrogen) was added, and the sample was rotated overnight at 4°C. Eluate was collected from the TEV cleaved beads. Three volumes of calmodulin binding buffer (10 mM β-mercaptoethanol, 25 mM HEPES, 150 mM NaCl, 1 mM Mg-acetate, 1 mM imidazole, 2 mM CaCl2, 0.1% NP-40, 10% glycerol), and calmodulin beads (Stratagene) were added, and the eluate was rotated for 1 h at 4°C. The calmodulin beads were washed with CBP binding buffer and eluted with calmodulin elution buffer (10 mM β-mercaptoethanol, 25 mM HEPES, 150 mM NaCl, 1 mM Mg-acetate, 1 mM imidazole, 2 mM EGTA, 0.1% NP-40, 10% glycerol). Typically, 0.2% of total extract, 2% of IgG beads after binding, 2% of IgG beads eluate, and 20% of calmodulin elute were loaded for Western blotting, except in Figures 2 and 5, where 2% and 3% of the extract used for co-immunoprecipitation experiments were loaded in whole-cell extract lanes for Western blotting.

For Western blot analysis, peroxidase–anti-peroxidase (Sigma) antibody was used to detect TAP tag, and anti-myc clone 9E10 was used to detect myc tag. The same membrane was probed for PGK using anti-PGK antibody (Molecular Probes) as the loading control.

Northern blot analysis

RNA was extracted from total protein lysates, or from immunoprecipitated samples with phenol/chloroform and treated with glyoxyl (Sigma). RNA was run on 1.3% agarose gels, transferred to Hybond-NX membrane (Amersham) and hybridized with probes made by random primers (RediPrime II, Amersham). Signals were quantified with PhosphorImager (Storm 860, Molecular Dynamics). The same membrane was probed for actin mRNA as the loading control.
DEAE fractionation and in vitro telomerase activity assay

DEAE fractionation was performed as described [Prescott and Blackburn 1997b]. RNA was prepared from total extract, flowthrough, the last wash, and each fraction for Northern blot analysis. Western blot analysis was performed with the same samples.

In vitro telomerase activity assay was performed as described [Prescott and Blackburn 1997b]. Wild-type yeast cells (BY4705) were transformed with either the vector control plasmid or the yPinX1p overexpression plasmid. Cells were grown in galactose medium to induce yPinX1p overexpression. S100 extraction and DEAE fractionation were carried out as described. To normalize, peak DEAE fractions from the vector control and yPinX1p plasmids containing the same amount of TLC1 were used for quantitative comparison.

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