Holoenzyme proteins required for the physiological assembly and activity of telomerase

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Many proteins have been implicated in the physiological function of telomerase, but specific roles of telomerase-associated proteins other than telomerase reverse transcriptase (TERT) remain ambiguous. To gain a more comprehensive understanding of catalytically active enzyme composition, we performed affinity purification of epitope-tagged, endogenously assembled *Tetrahymena* telomerase. We identified and cloned genes encoding four telomerase proteins in addition to TERT. We demonstrate that both of the two new proteins characterized in detail, p65 and p45, have essential roles in the maintenance of telomere length as part of a ciliate telomerase holoenzyme. The p65 subunit contains an La motif characteristic of a family of direct RNA-binding proteins. We find that p65 in cell extract is associated specifically with telomerase RNA, and that genetic depletion of p65 reduces telomerase RNA accumulation *in vivo*. These findings demonstrate that telomerase holoenzyme proteins other than TERT play critical roles in RNP biogenesis and function.

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Telomerase is a ribonucleoprotein (RNP) reverse transcriptase specialized for addition of telomeric repeats to chromosome 3'-ends. The telomerase reverse transcriptase protein (TERT) contains active-site motifs characteristic of viral reverse transcriptases and other motifs shared only among TERTs (for review, see Kelleher et al. 2002). The telomerase RNA subunit provides a template sequence for telomeric repeat synthesis and also functions in determining enzyme catalytic properties (for review, see Blackburn 2000; Harrington 2003). TERT and telomerase RNA from a few species, if coexpressed in specific cell lysates, can reconstitute a properly templated reverse transcriptase activity (Weinrich et al. 1997). For *Tetrahymena* TERT and telomerase RNA assembled in rabbit reticulocyte lysate, the activity of the minimal recombinant enzyme has notable differences from endogenous enzyme activity in substrate specificity and elongation processivity (Collins and Gandhi 1998; Hardy et al. 2001). These discrepancies could derive from altered protein or RNA folding, differences in the modification state of TERT or telomerase RNA, or differential association of additional RNP factors.

Biochemical fractionation of cell extract suggests that physiologically assembled telomerase RNPs possess a

substantial mass in excess of TERT and telomerase RNA (for review, see Collins 1999). Many proteins that associate with telomerase complexes have been identified using biochemical, genetic, and candidate approaches (for review, see Ford et al. 2002; Harrington 2003). Owing to the scarcity of endogenous telomerase complexes, however, it has been difficult to determine whether individual telomerase-associated proteins interact with a common RNP or with distinct, heterogeneous RNP populations.

One important biological role for telomerase-associated proteins is the assembly of telomerase RNA into stable RNP, a role evidently not fulfilled by TERT itself. In vertebrate cells, the telomerase RNA H/ACA motif is essential for precursor RNA 3'-end processing and mature RNA accumulation (Mitchell et al. 1999a). This motif interacts with dyskerin, NHP2, NOP10, and GAR1 (Mitchell et al. 1999b; Dragon et al. 2000; Pogacic et al. 2000), proteins also associated with a large family of H/ACA small nucleolar (sno) RNPs that catalyze the posttranscriptional, site-specific conversion of uridine to pseudouridine (for review, see Kiss 2002). In *Saccharomyces cerevisiae*, Sm proteins associate with telomerase RNA to enhance RNA accumulation (Seto et al. 1999). To date, none of these proteins implicated in cellular telomerase RNP assembly is unique to telomerase RNP, complicating the analysis of their telomerase biogenesis roles in specific. In the ciliate *Euplotes aediculatus*, telomerase affinity purification by template-comple-

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mentary oligonucleotide hybridization identified two telomerase components, TERT and p43 (Lingner and Cech 1996). *Euplotes* p43 contains an La motif (Aigner et al. 2000), a defining characteristic of the eukaryotic La protein and additional La motif-containing proteins with direct RNA-binding activities (for review, see Wolin and Cedervall 2002). Indeed, recombinant p43 can bind telomerase RNA directly and antibodies against p43 substantially immunodeplete telomerase RNA and activity from *Euplotes* nuclear extracts (Aigner et al. 2000, 2003). A role for p43 in telomerase nuclear retention has been proposed based on changes in the efficiency of telomerase RNA electroelution from macronuclei when p43 is targeted for RNAi (Möllenbeck et al. 2003).

Telomerase-associated proteins also play biologically important roles in coupling telomerase RNP to telomere substrates. In *S. cerevisiae*, mutations in the genes encoding Est1p, Est3p, and Cdc13p can impair telomerase-dependent telomere length maintenance without affecting enzyme activity in cell extract (Lingner et al. 1997). Est1p and Est3p were shown by immunoprecipitation to interact with catalytically active telomerase RNP (Steiner et al. 1996; Hughes et al. 2000). Current models suggest that Est1p either recruits telomerase to a telomeric G-strand 3'-overhang or binds there independently to stimulate the activity of prebound telomerase RNP (Evans and Lundblad 1999; Taggart et al. 2002). Human and *Schizosaccharomyces pombe* EST1 proteins sharing limited conservation with the budding yeast Est1p have recently been identified using a bioinformatics approach. At least a subset of the human EST1 proteins can interact with active telomerase and affect telomere biology (for review, see Lundblad 2003; Vega et al. 2003).

The ciliate *Tetrahymena thermophila* is a favorable model system for the study of telomeres and telomerase. *Tetrahymena* have a silent, diploid germ-line micronucleus containing five pairs of mitotically segregated chromosomes. The transcriptionally active macronucleus, in contrast, has ~20,000 chromosomes and thus ~40,000 telomeres. Fully half of these derive from amplification of a palindromic chromosome encoding ribosomal RNA (rDNA). The macronuclear genome also contains ~45 copies each of ~200 additional chromosomes created during a developmentally programmed process of fragmentation and amplification. The large number of telomeres obliges an abundance of telomerase that can be readily detected in cell extracts (Greider and Blackburn 1985). In addition to this biochemical advantage, *T. thermophila* is highly amenable to genetic and molecular genetic analysis (for review, see Turkewitz et al. 2002). The macronucleus divides amitotically during vegetative growth, producing daughter cells with different allele frequencies at each locus. In the process of phenotypic assortment, a locus disrupted by integration of a selectable marker can be enriched relative to the wild-type locus by culture growth under selective pressure to reach one of two endpoints: a complete gene knockout if the targeted gene is not essential, or a function-limiting gene dose knockdown in the case of essential genes (for review, see Karrer 2000).

Even in ciliate systems, telomerase purification has proven to be a difficult task. Telomeres and telomerase are most abundant in spirotrichous ciliates such as *Euplotes*. Affinity purification from *E. aediculatus* extracts using template-complementary oligonucleotide hybridization was successful in the identification of TERT and p43 but was also noted to induce a change in RNP conformation or composition (Lingner and Cech 1996). Human and *Tetrahymena* cell extracts subjected to a similar purification strategy were even more clearly disrupted by this approach, with substantial changes in telomerase RNP mass and composition (Schnapp et al. 1998; data not shown). Conventional chromatography strategies are also disruptive, because of the extent of purification required. The *Tetrahymena* p80/p95 complex identified in this manner by cofractionation with telomerase RNA (Collins et al. 1995) is not required for telomerase catalytic activity and negatively regulates telomere length in vivo (Miller and Collins 2000). We have now developed a more rapid and gentle affinity purification strategy that preserves the integrity of telomerase holoenzyme. Here, we exploit the combined biochemical and genetic advantages of *Tetrahymena* as a telomerase model system to isolate telomerase-associated proteins and characterize their roles in enzyme function.

Results

Transgene expression of epitope-tagged Tetrahymena TERT

Transgenes expressing epitope-tagged TERT were integrated into the *Tetrahymena* genome by replacement of the nonessential β -tubulin 1 (*BTU1*) coding region. Growth of the *Tetrahymena* strain CU522 shows enhanced sensitivity to taxol, because of an amino acid substitution in the β -tubulin protein encoded at *BTU1*. Disruption of *BTU1* does not compromise viability because a second gene also encodes β -tubulin, but in strain CU522, *BTU1* disruption confers increased resistance to taxol (Gaertig et al. 1994). Of the several epitope tags screened for use in large-scale TERT affinity purification, the most effective was a C-terminal tandem affinity purification (TAP) tag (Rigaut et al. 1999). This epitope tag consists of two terminally located protein A domains that bind to IgG, an adjacent TEV protease cleavage site to allow elution from IgG resin, and a calmodulin-binding peptide that binds to calmodulin resin in the presence of calcium and elutes with EGTA (see Supplementary Fig. 1A). The TAP tag has been used in previous studies for the purification of other ribonucleoprotein complexes (Puig et al. 2001; Dragon et al. 2002).

We targeted a transgene encoding TAP-tagged TERT (TERT-TAP) for homologous recombination into the *BTU1* locus, placing TERT expression under control of the endogenous *BTU1* promoter (Gaertig et al. 1999). Correct integration specificity was verified by Southern blot hybridization (data not shown). The resulting strain expressed both TERT-TAP from the *BTU1* locus and

wild-type TERT from the endogenous *TRT* locus, as monitored by immunoblotting with a polyclonal antibody specific for TERT (Fig. 1A). The immunoblot signal from TERT-TAP is substantially amplified by binding of the TERT antibody to the protein A domains of the TAP tag. To determine whether TERT-TAP was produced at a physiologically appropriate level, we incubated total cell

Figure 1. TERT-TAP is expressed and catalytically active. (A) Whole-cell extract from TERT-TAP or a control strain was normalized by total protein concentration and analyzed by immunoblot for TERT. Control cells in panels A, B, and C expressed both wild-type TERT and a tagged TERT differing from TERT-TAP by substitution of the Flag peptide epitope for both protein A domains. Recombinant TERT (rTERT) expressed in *E. coli* was used as an immunoblot standard. A nonspecific cross-reacting polypeptide is shown as a loading control (LC), with equal loading also verified by staining of the immunoblot membrane (not shown). In the TERT-TAP sample, a minor TERT-TAP degradation product (*) migrates with slightly slower mobility than wild-type TERT (see Supplementary Fig. 1 for additional data). (B,C) Extracts normalized by total protein concentration from TERT-TAP cells and control cells were bound to IgG resin. Beads were assayed for telomerase RNA by Northern blot (B) and were assayed independently for telomerase activity (C). The activity assay product ladder is annotated to indicate product lengths from processive synthesis of telomeric repeats. Prior to RNA precipitation for Northern blot analysis, a truncated form of purified recombinant telomerase RNA was added as a loading control (LC). (D) Genomic DNA was digested with HindIII and analyzed by Southern blot from untransformed wild-type cells (WT); wild-type cells transformed with the *TRT* gene disruption cassette, phenotypically assorted and then released from selection (ΔTRT in WT); or TERT-TAP cells similarly transformed with the *TRT* gene disruption cassette, assorted and then released from selection (ΔTRT in TERT-TAP). The probe recognizes the wild-type and disrupted *TRT* loci with equivalent hybridization potential. (E) Macronuclear telomere length was assayed in wild-type cells or TERT-TAP cells disrupted for the wild-type *TRT* gene. Genomic DNA was digested with HindIII, resolved on a nondenaturing agarose gel, and assayed by Southern blot to detect rDNA telomeres. The hybridization probe recognizes the 357-bp subtelomeric region of rDNA chromosomes.

extract with TEV protease to remove the TERT-TAP protein A domains before immunoblot analysis (Supplementary Fig. 1B). Using this method, we determined that epitope-tagged TERT in this and all other TERT transgene strains accumulated to a level less than or equal to endogenous TERT in wild-type cells.

To determine whether TERT-TAP was assembled into a catalytically functional telomerase RNP, we assayed for cofractionation of telomerase RNA and activity with TERT-TAP after purification on IgG resin. Both telomerase RNA and activity were recovered on IgG resin from extracts of strains expressing TERT-TAP but not from control strains expressing TERT C-terminally fused to a TAP tag variant containing a Flag epitope in place of the protein A domains (Fig. 1B,C). This result indicates that TERT-TAP associates with telomerase RNA in enzymatically functional telomerase RNP complexes. Notably, upon purification of the epitope-tagged TERT, we failed to detect any association with coexpressed wild-type TERT even when the wild-type TERT was much more abundant (Supplementary Fig. 2). This result suggests that *Tetrahymena* TERT is monomeric in the active telomerase RNP purified from cell extract.

To determine whether TERT-TAP telomerase RNPs were functional *in vivo* as well as *in vitro*, we disrupted the endogenous *TRT* gene to eliminate the expression of wild-type TERT protein. We used phenotypic assortment to test whether TERT-TAP could substitute genetically for wild-type TERT during vegetative growth (see Supplementary Fig. 3). Because *TRT* is an essential gene in wild-type cells, it is impossible to completely replace the endogenous *TRT* locus with a transgene conferring paromomycin resistance (Miller and Collins 2000). Instead, under maximum selective pressure, a combination of wild-type and disrupted loci is observed. This genotype reverted to predominantly wild-type *TRT* in the absence of selection (Fig. 1D, lane 1). In contrast, in strains containing the TERT-TAP transgene, the endogenous *TRT* locus could be fully replaced with a transgene encoding paromomycin resistance. No back-assortment to wild-type *TRT* was detected even after multiple generations of growth in the absence of selective pressure (Fig. 1D, lane 2). Immunoblot analysis confirmed that *TRT* knockout cells with the TERT-TAP transgene do not express wild-type TERT protein (Supplementary Fig. 1B). Moreover, these cells expressed TERT-TAP protein at the normal physiological level of the endogenous TERT protein (Supplementary Fig. 1B). These experiments demonstrate that TERT-TAP fulfills the essential functions of wild-type TERT *in vivo*. Cells expressing TERT-TAP instead of wild-type TERT had no growth defect and no apparent change in telomerase activity assayed in cell extract (data not shown). Cells expressing only TERT-TAP maintained macronuclear telomere lengths within the variability between wild-type strains (Fig. 1E; data not shown), confirming that the tagged protein functions in telomerase-dependent telomere length maintenance. All subsequent experiments shown here were performed using the TERT-TAP transgene strain lacking wild-type TERT.

Affinity purification of epitope-tagged TERT reveals additional telomerase proteins

Extracts from TERT-TAP cells were used for telomerase affinity purification with only minor changes from the standard TAP protocol (Puig et al. 2001). Telomerase activity (Fig. 2A), telomerase RNA (data not shown), and TERT protein (Fig. 2B) were recovered and enriched with each specific binding and elution step. Importantly, telomerase activity in the purified samples retained the high repeat addition processivity of the authentic cellular holoenzyme. By SDS-PAGE, four major polypeptides in addition to TERT were evident by silver stain (Fig. 2C) or Coomassie stain (data not shown) in several variations of purification conditions and with numerous extract preparations. These polypeptides were not recovered in

purifications performed in parallel from extracts of control strains (data not shown). Notably, p80 and p95 were not detected in the purified material either by silver stain (Fig. 2C) or immunoblot (data not shown; see Discussion).

Peptide sequences were obtained from proteolytic digests of each protein and used to design primers for degenerate PCR. PCR products were obtained for which translation in silico corroborated additional peptide sequence not used in primer design. Subsequently, RT-PCR and RACE were used to obtain full-length cDNA sequences, and PCR was used to obtain the sequences of the macronuclear genomic loci (see Supplemental Material). Each full-length gene is predicted to encode a protein with molecular weight closely matching the apparent molecular weight by SDS-PAGE. For chronological reasons, we began our analysis of telomerase holoenzyme proteins with p65 and p45.

We designated the p45 gene *TAP45*, for TERT- or telomerase-associated protein of 45 kDa. The protein encoded by the full-length cDNA exhibits no apparent homology to other known proteins when examined using BLAST searches of the GenBank database. We termed the p65 gene *TAP65*. BLAST searches of GenBank with p65 revealed limited homology to other proteins within an N-terminal region representing a divergent La motif (Fig. 2D). This motif, strikingly also found in *Euplotes* p43, was originally discovered in the eukaryotic La protein and has subsequently been found in additional direct RNA-binding proteins (for review, see Wolin and Cedervall 2002). In its most characterized role, La binds to the 3' polyuridylyl tract of primary transcripts of RNA polymerase III and to other small RNA precursors, protecting them from exonucleolytic degradation and promoting subsequent assembly to mature RNP. Both the La motif and a putative RNA recognition motif (RRM) C-terminal to the La motif were detected in a p65 query of the Prosite database (<http://us.expasy.org/prosite>). Sequence homology between *Tetrahymena* p65 and *Euplotes* p43 was contained almost exclusively within the La motif, not the adjacent putative RRM. The full-length forms of the two proteins share ~20% amino acid identity and ~40% amino acid similarity overall.

Figure 2. Four major proteins copurify with TERT-TAP. (A) Telomerase activity was assayed in the TEV protease and EGTA elutions of the affinity purification. The activity assay product ladder is annotated to indicate product lengths from processive synthesis of telomeric repeats. (B) TERT recovered in an aliquot of the EGTA elution was examined by immunoblot in comparison with 5 ng of recombinant TERT standard (rTERT). (C) Aliquots of the TEV protease and EGTA elutions were resolved by SDS-PAGE, and proteins were visualized by silver stain. The molecular weights of proteins in the Marker lane are indicated at left, and the most abundant polypeptides in the EGTA elution are indicated at right. (D) The La motif in p65 was aligned with La motifs in other proteins. Identical residues are boxed and shaded dark. Similar residues are lightly shaded. The numbering at left in the lower alignment panel indicates the number of amino acids not shown between conserved sequences at the end of the top panel and the beginning of the bottom panel.

Endogenous p65 and p45 are components of telomerase holoenzyme

Because no extensive purification protocol, including our affinity purification, can recover all of the tagged protein, telomerase RNA, or telomerase activity in the starting material, it was important to determine using whole-cell extract whether p65 and p45 associate with a large percentage of the total telomerase RNP pool. We thus performed immunodepletion studies with polyclonal antibodies raised and affinity-purified against full-length p65 and p45 overexpressed using synthetic genes in *Escherichia coli* (R. Prathapam, K.L. Witkin, and K. Collins, unpubl.). These antibodies recognized their cognate proteins specifically (Supplementary Fig. 4). We subjected wild-type *Tetrahymena* whole-cell extract to

serial depletions in successive, 1-h incubations with bead-immobilized antibody. Antibody specific for p65 or p45 recruited telomerase activity and telomerase RNA to beads (Fig. 3A, lanes 1–4), whereas preimmune serum (Fig. 3A, lanes 5–6) or affinity-purified antibody raised against an unrelated human protein (data not shown) did not. Depletion with p65 or p45 antibody reduced telomerase activity in an aliquot of the supernatant to a barely

detectable (after round 1) or undetectable (after round 2) level (Fig. 3A, lanes 7–10), whereas preimmune serum (Fig. 3A, lanes 11–12) or another control antibody (data not shown) did not. Northern blot hybridization revealed that telomerase RNA remaining in the second-round immunodepletion supernatant was reduced to <10% of the amount of telomerase RNA in the control antibody supernatant (Fig. 3B). In contrast, other small RNAs including SRP RNA, U3 snoRNA, U2 small nuclear (sn) RNA, and U6 snRNA remained in the supernatant after immunodepletion with either antibody. Notably, none of the antibodies depleted a nonspecific radiolabeling activity evident in the whole-cell extracts (Fig. 3A, products marked by an asterisk). Using either p65 or p45 antibody, the majority of telomerase activity and RNA could be removed from supernatants and recovered on antibody beads. This finding demonstrates that both proteins are stable telomerase RNP components, and furthermore that the predominant pool of telomerase RNP contains both p65 and p45 bound together to each molecule of telomerase RNA.

p65 associates specifically with telomerase RNA

To further confirm the association of p65 and p45 with active telomerase RNP, we created *Tetrahymena* strains expressing an epitope-tagged version of each protein. Proteins fused at their C termini to a TAP tag were expressed from transgenes integrated into the CU522 macronuclear genome at the *BTU1* locus as described for TERT-TAP above. We confirmed the intended integration events by Southern blot (data not shown). In each case, the tagged protein accumulated at a much lower level than TERT-TAP and likely for this reason was unable to substitute genetically for the wild-type protein. Although p65-TAP and p45-TAP expression levels were too low to permit detection of associated proteins after affinity purification from cell extract, purification could recover telomerase activity substantially enriched over any background recovered from wild-type control extract (Fig. 4A). The relative recovery of telomerase activity is not a meaningful comparison in this experiment, because individual TAP-tagged proteins are expected to bind IgG resin at different efficiencies and are present at different levels in the starting extract as well.

The presence of an La motif in p65 raises the intriguing possibility that it could be the *Tetrahymena* La protein, playing a general role in binding RNA polymerase III transcripts including telomerase RNA. We used two sequential steps of affinity purification of p65-TAP to investigate whether p65 is preferentially associated with telomerase RNA, as expected of a telomerase-specific protein, or whether like La it associates with the large variety of RNA species. We compared the spectrum of RNAs that copurified with TERT-TAP and p65-TAP from whole-cell extracts, also monitoring any background RNAs recovered in a parallel mock purification from extract of a control strain. RNA was precipitated from the purified samples, radiolabeled by addition of a single 3'-nucleotide, then resolved by denaturing

Figure 3. Endogenous p65 and p45 are associated with telomerase activity and RNA. Whole-cell extract was subject to immunodepletion with antibodies against p65 (α p65) and p45 (α p45) or with preimmune serum (Pre). (A) A small aliquot of extract samples following first (1) or second (2) round immunodepletion (Supernatants) and a 30-fold higher relative amount of each bead-bound sample (Beads) were assayed for telomerase activity. Asterisks indicate telomerase-independent background from radiolabeling of nucleic acids in the crude extract. Telomerase RNA associated with an aliquot of each antibody bead sample was detected by Northern blot. (B) The extent of supernatant RNA depletion was quantitated following the second round of immunodepletion by comparing p65 (α p65), p45 (α p45), and preimmune (Pre) antibody depletions to control for any RNA degradation over the incubation in crude extract. A single blot was probed for telomerase RNA, recombinant truncated telomerase RNA added prior to precipitation of RNA samples as a loading control (LC) and endogenous control RNAs (at right). Hybridization signal intensity was quantitated by PhosphorImager analysis to determine the specificity of immunodepletion (at left) using antibodies against p65 (light gray) and p45 (dark gray). Recovery of control RNAs normalized to the preimmune immunodepletion control ranged from 80% to 120% in the single experiment shown, reflecting small irreproducible differences in sample recovery.

Figure 4. Epitope-tagged p45 and p65 copurify active telomerase RNP. (A) Whole-cell extracts from TERT-TAP, p65-TAP, p45-TAP, and wild-type strains normalized by total protein concentration were incubated with IgG resin. Specifically bound complexes were eluted from the resin with TEV protease and assayed for telomerase activity. (B) RNAs recovered after the complete TAP purification protocol from TERT-TAP or p65-TAP cell extracts were radiolabeled by a single 3'-nucleotide addition. Because p65-TAP was expressed at a lower level than TERT-TAP, 90% (lane 1) and 10% (lane 2) of the TERT-TAP purification were loaded in separate lanes for better comparison to 100% of the p65-TAP purification (lane 3). RNAs were resolved by denaturing gel electrophoresis. The arrow indicates the predominant RNA species copurified with TERT-TAP and p65-TAP that also comigrated with a purified telomerase RNA standard (data not shown). The asterisk indicates a background RNA present in the control purification from wild-type cells (lane 4). The Marker lane contains end-labeled DNA ladder.

gel electrophoresis (Fig. 4B). The TERT-TAP-associated RNA profile (Fig. 4B, lanes 1–2 loaded at two different concentrations) matched the p65-TAP associated RNA profile (Fig. 4B, lane 3) and was composed of a single predominant species not present in the wild-type extract control purification (Fig. 4B, lane 4). The RNA associated with TERT-TAP and p65-TAP migrated consistent with an expected telomerase RNA size of 159 nt and comigrated with a recombinant telomerase RNA standard (data not shown). Some RNA degradation occurred during the 3'-radiolabeling protocol, resulting in a variable ladder of smaller RNA products in all samples including the purified RNA standard. The similarity of the TERT- and p65-associated RNA profiles suggests that both proteins are preferentially associated with telomerase RNA.

Telomerase p65 and p45 are encoded by essential genes

The genes encoding TERT and telomerase RNA are essential for *Tetrahymena* viability (Miller and Collins 2000). To determine whether the genes encoding p65 and p45 are similarly essential, we performed gene disruption of *TAP65* and *TAP45* loci in a manner identical to the *TRT* gene disruption described above. In strains taken to maximal phenotypic assortment and then released from selective pressure for an extended interval of growth, analysis of *TAP65* (Fig. 5A) and *TAP45* (Fig. 5B) loci revealed that both wild-type and disrupted genes remained present. The persistence of the wild-type loci through phenotypic assortment indicates that both p65 and p45 are encoded by essential genes (see Supplementary Fig. 3).

Short telomeres in p65 and p45 knockdown strains

We used the maximally assorted gene knockdown strains to examine the impact of limiting p65 or p45 function on the physiological role of telomerase in telomere length maintenance. Half of all macronuclear telomeres have precisely the same subtelomeric sequence, derived from the palindromic rDNA chromosome ends. To examine telomere length in p65 and p45 knockdown strains, purified genomic DNA was digested with *HindIII*, resolved by denaturing gel electrophoresis, and probed for the terminal fragment of the macronuclear rDNA chromosome using an oligonucleotide complementary to the subtelomeric region on the telomeric re-

Figure 5. The genes encoding p45 and p65 are essential. Cells were transformed with a construct designed to disrupt the endogenous gene encoding either p65 (*TAP65* locus) or p45 (*TAP45* locus). (A) Genomic DNA from wild-type cells and from cells transformed with the *TAP65* disruption construct (65KD), maximally assorted and released from selective pressure was digested with *BbsI* and *SacI* and then analyzed by Southern blot. The probe recognizes wild-type and disrupted *TAP65* loci with equivalent hybridization potential. (B) Genomic DNA from wild-type cells and from cells transformed with the *TAP45* disruption construct (45KD), maximally assorted and released from selective pressure was digested with *SpeI* and *SacI* and then analyzed by Southern blot. The probe recognizes wild-type and disrupted *TAP45* loci with equivalent hybridization potential.

peat AC-rich strand. Previous studies using numerous experimental controls have demonstrated that telomere length maintenance is not affected by the presence of selective drug alone (Miller and Collins 2000; data not shown). All cultures were maintained at a selective drug concentration that permitted a growth rate comparable to that of wild-type cells grown in the absence of drug selection.

We found that telomeres were dramatically shorter in p65 and p45 knockdown strains (Fig. 6A, lanes 2–3) than in wild-type cells (Fig. 6A, lane 1). Each hybridizing DNA fragment contains a precise 357 nt of subtelomeric sequence and a variable length of telomeric repeat array. We estimate that under these assortment conditions, the telomeric repeat tract in p65 and p45 knockdown strains was predominantly ~150–350 bp. In contrast, the telomeric repeat tract in wild-type cells was predominantly ~350–550 bp. Genomic DNA digested with the frequently cutting restriction enzyme *MseI* and probed using a telomeric repeat oligonucleotide confirmed the substantial telomere loss on rDNA and other chromosome ends in p65 and p45 knockdown strains (data not shown).

In wild-type strains, macronuclear telomere length increases by 3–10 bp per generation when cultures are kept in continuous log-phase growth at 30°C (Larson et al. 1987). We examined whether this telomere length regulation was affected in p65 and p45 knockdown strains. Cultures maintained under selection (day 0) were released into drug-free media and maintained in log-phase growth for 10 d (~80–100 doublings total). Telomere length was assayed for rDNA as described above, except that nondenaturing gel electrophoresis was used to re-

solve longer telomeres (Fig. 6B). In wild-type cells, telomere length increased from day 0 to day 5 as expected (Fig. 6B, WT lanes). In contrast, telomere elongation was marginal in p65 and p45 knockdown strains. By 10 d of growth in the absence of selection, telomere length in the knockdown strains had increased detectably but still remained less than telomere length in wild-type cells at day 0 (data not shown). Analysis of rDNA and other telomeres using genomic DNA digested with *MseI* and probed with a telomeric repeat oligonucleotide confirmed these altered telomere length dynamics (data not shown).

A telomerase RNA accumulation defect in p65 knockdown strains

One mechanism that could account for the defect in telomere maintenance in the p65 and p45 knockdown strains is a failure in accumulation of the telomerase RNP. Total RNA prepared from cultures at days 0 and 5 of the time course described above was analyzed for telomerase RNA by Northern blot hybridization. Telomerase RNA accumulation was unaffected by continuous log-phase growth in two independent wild-type cell cultures (Fig. 7A, lanes 1–4). No change in telomerase RNA level was detected in p45 knockdown cultures under selective pressure (Fig. 7A, lane 5) or after 5 d of continuous culture in the absence of selection (Fig. 7A, lane 6). In contrast, telomerase RNA accumulation in the p65 knockdown strain was inhibited dramatically under selective pressure (Fig. 7A, lane 7) and only partially recovered in 5 d of continuous culture after release from selection (Fig. 7A, lane 8).

The preferential interaction of p65 with telomerase RNA in extract (Fig. 4B) would predict a specific telomerase RNA accumulation defect in the p65 knockdown strain. To test this hypothesis, we reprobbed the same Northern blot membrane to detect additional small RNAs. Steady-state levels of 5S rRNA, SRP RNA, and U2 snRNA (Fig. 7A) and also U6 snRNA and U3 snoRNA (data not shown) were not affected by growth conditions in any strain. These RNAs share properties with telomerase RNA including transcription by RNA polymerase III (5S, SRP, U6) or a nuclear site of function (U2, U6, U3). We also investigated whether TERT levels decreased in p65 knockdown strains in parallel with the decrease in telomerase RNA. We compared the levels of telomerase RNA and TERT in whole-cell extracts of the wild-type and knockdown strains. Telomerase RNA detected by Northern blot hybridization (Fig. 7B, top) and wild-type TERT detected by immunoblot (Fig. 7B, middle) were both reduced in extracts from two independently generated p65 knockdown strains (Fig. 7B, lanes 3–4), in comparison to extracts from wild-type or p45 knockdown strains (Fig. 7B, lanes 1–2). These findings provide genetic evidence that p65 functions *in vivo* to promote telomerase RNA accumulation, playing a crucial and apparently specific, physiological role in telomerase holoenzyme RNP assembly.

Figure 6. p65 and p45 are required for telomere maintenance in the macronucleus. (A) Genomic DNA from wild-type, p45 knockdown, and p65 knockdown cells was digested with *HindIII* before Southern blot analysis to detect rDNA telomeres. DNA was resolved on a denaturing acrylamide gel. Markers were detected by hybridization to the same membrane to calculate telomere length as precisely as possible. (B) Wild-type, p45 knockdown, and p65 knockdown strains were grown under maximum drug selection for 24 h followed by 5 d of continuous log-phase growth in the absence of selection. Cells were harvested each day from 0 to 5 d after release from selection for telomere length analysis as in A, except that DNA was electrophoresed on a nondenaturing agarose gel to resolve telomere lengths >1 kb. Markers were detected by ethidium bromide staining of a gel slice removed before the remainder of the gel was transferred to membrane and should be considered as an approximation.

Figure 7. Telomerase RNA and TERT levels decrease in p65 knockdown strains. (A) Total RNA was prepared from equal numbers of wild-type cells in log-phase growth under no drug selection (WT-1) or from wild-type and knockdown strain cells in log-phase growth under the maximum sublethal drug selection (WT-2, 45KD, 65KD). These same cell cultures were also grown for an additional 5 d of continuous log-phase growth in the absence of selection. RNA levels were analyzed by Northern blot for telomerase, 5S, SRP, and U2 RNAs. (B) Whole-cell extracts from wild-type, p45 knockdown, and two independent p65 knockdown strains normalized by total protein concentration were used for Northern blot analysis of telomerase RNA (top) or immunoblot analysis of TERT (middle). A nonspecific cross-reacting polypeptide is shown as a loading control (LC) for the immunoblot (bottom), with equal loading also verified by staining of the immunoblot membrane (data not shown). (C) This model depicts the proposed role for p65 in establishing the level of telomerase RNA accumulation, with the subsequent incorporation of TERT, p45, p75, and p20 to form a biologically active telomerase RNP holoenzyme.

Discussion

Telomerase holoenzyme purification

To accomplish a rapid and gentle affinity purification of telomerase RNP, we expressed TAP-tagged *Tetrahymena* TERT in substitution of the wild-type protein. Telomerase RNPs that incorporate the tagged TERT were shown to have full catalytic activity in vitro and in vivo. Four major proteins copurified with TERT and telomerase RNA, with several less abundantly recovered proteins also coenriched in the purified material (Fig. 2C). Our studies do not address the composition of any hypothetical telomerase RNP lacking TERT or the potential association of developmentally stage-specific holoenzyme proteins.

A conserved role for p65

In yeast and mammalian systems, telomerase proteins other than TERT confer telomerase RNA stability and

promote RNP accumulation in vivo. In vertebrate telomerase RNAs, an H/ACA motif serves as the binding site for cooperative assembly of dyskerin, NHP2, NOP10, and GAR1 (Mitchell et al. 1999b; Dragon et al. 2000; Pogacic et al. 2000); the yeast telomerase RNA precursor instead assembles with Sm proteins (Seto et al. 1999). Our findings here reveal that a role for holoenzyme proteins in determining steady-state telomerase RNA accumulation is universal, even though the mechanism of their interaction with telomerase RNA is not. Ciliate telomerase RNAs (~150 nt) are much smaller than yeast (>1 kb) or vertebrate (~450 nt) telomerase RNAs and are transcripts of RNA polymerase III rather than RNA polymerase II (for review, see Collins 1999). Nonetheless, telomerase RNA accumulation in *Tetrahymena* requires the holoenzyme protein p65. By extension, we anticipate that telomerase RNA accumulation in *Euplotes* will require p43.

Several lines of evidence suggest that p65 functions directly to increase the cellular stability of telomerase RNA (Fig. 7C). In *Tetrahymena*, overexpression of transgene-encoded telomerase RNA or TERT does not increase the absolute level of subunit accumulation; instead, it competitively reduces endogenous subunit expression to maintain the same overall level of accumulation (Yu et al. 1990; D. Cunningham and K. Collins, unpubl.). This finding suggests that a third factor establishes the steady-state level of telomerase RNP. We propose that p65 is this factor. The presence of an La motif and putative RRM indicate that p65 should bind telomerase RNA directly. The dramatic reduction in telomerase RNA accumulation in p65 gene knockdown strains supports this conclusion. Although TERT accumulation is also reduced in these strains, we believe that TERT is destabilized as a secondary effect of telomerase RNA depletion. This would be consistent with the observation in *S. cerevisiae* that accumulation of the yeast TERT Est2p is reduced in the absence of telomerase RNA (Taggart et al. 2002). Results from other *Tetrahymena* telomerase RNP purification strategies reveal that telomerase RNP can be isolated lacking TERT but that TERT is not stable independent of telomerase RNA (K.L. Witkin and K. Collins, unpubl.). Together, these findings suggest that telomerase RNA is destabilized in the absence of p65, and that TERT is degraded in the absence of stable telomerase RNA (Fig. 7C).

Although the La motif is a hallmark feature of the eukaryotic La protein, it is shared by a larger family of proteins distinct from La (for review, see Wolin and Cedervall 2002). Pairwise comparisons of the sequences of *Tetrahymena* p65, *Euplotes* p43, and authentic La proteins did not reveal substantial homology beyond the La motif. La itself can associate with the human telomerase RNA transcribed by RNA polymerase II (Ford et al. 2001). However, our experimental evidence suggests that p65 is an La-motif, telomerase-specific protein instead of ciliate La. First, whereas telomerase RNA level decreased in p65 knockdown strains, the accumulation of other RNA polymerase III transcripts was unaffected. Second, the purification of p65-TAP readily recovered

telomerase RNA but not other RNA polymerase III transcripts. We note that BLAST searches of the *Tetrahymena* macronuclear genome (<http://www.tigr.org/tdb/e2k1/ttg>) can yield at least one region other than the *TAP65* locus with potential to encode an La motif. Whether this region produces an mRNA transcript and corresponding protein, and if so whether the protein represents ciliate La, remain to be determined.

Other telomerase holoenzyme protein roles

In addition to directing telomerase RNA folding and RNP stability, telomerase holoenzyme proteins must also coordinate RNP interactions with intranuclear storage sites (Fang and Cech 1995; Wong et al. 2002) and with telomeres (for review, see Vega et al. 2003). Genetic criteria demonstrate that p45 is required for telomere length maintenance but not for cellular accumulation of telomerase RNA or TERT, suggesting that p45 acts at a step after stable RNP biogenesis (Fig. 7C). Telomerase holoenzyme proteins including p45 could be required for catalytic activation. Because knockdown strains retain some wild-type telomerase enzyme even at maximal phenotypic assortment, we cannot reliably assay the activity of an RNP lacking p45 by using p45 knockdown cell extract. Further analysis will be required to determine the precise role of p45 in telomerase function and to investigate roles for p75 and p20 in telomerase-dependent telomere length maintenance.

Does the set of five proteins and telomerase RNA recovered by TERT-TAP affinity purification represent a complete telomerase holoenzyme? Telomerase activity in *Tetrahymena* extract fractionates at ~270 kDa by glycerol gradient sedimentation or ~500 kDa by gel filtration (Collins and Greider 1993; Wang and Blackburn 1997). The RNP recovered through TERT-TAP affinity purification would be consistent with these mass estimates. A strikingly similar fractionation has been reported for active telomerase in extracts of *E. aediculatus*, comparing both glycerol gradient sedimentation and gel filtration (Lingner and Cech 1996; Aigner et al. 2003). This suggests that the telomerase-associated proteins identified here could also be components of a *Euplotes* telomerase holoenzyme, with yet additional proteins joining to form larger holoenzyme complexes detected in mated *Euplotes crassus* undergoing developmentally programmed chromosome fragmentation (Greene and Shippen 1998).

Our TERT-TAP affinity purification did not recover p80 and p95, two proteins characterized by co-fractionation with telomerase RNA through multiple steps of conventional chromatography (Collins et al. 1995). While purified recombinant p80 and p95 bind to telomerase RNA and telomeric repeat substrate DNA in vitro (Gandhi and Collins 1998), they are not important for catalytic activity in vitro or in vivo and at the physiological level are negative regulators of telomere length (Miller and Collins 2000). It has been suggested that p80 and p95 are not associated with active telomerase be-

cause the two proteins more effectively coimmunoprecipitate each other than they coimmunoprecipitate TERT, telomerase RNA or telomerase activity from cell extracts (Mason et al. 2001). Such results are difficult to interpret conclusively given that the p80/p95 complex is in substantial molar excess of telomerase RNP (K.L. Witkin and K. Collins, unpubl.). However, in addition, immunopurification of TERT does not obligatorily copurify p80/p95 (Mason et al. 2001). We find that under some conditions—for example, RNP purification using 2'-O-methyl RNA oligonucleotides complementary to the template region—p80 and p95 are recovered with telomerase RNA to a much greater extent than is TERT (K.L. Witkin and K. Collins, unpubl.). The nucleic acid binding properties of p80/p95 may promote an interaction with misfolded telomerase RNP in cell lysate under purification conditions that are less gentle than used here for purification of TAP-tagged RNP. Disruptive conditions may allow the formation of an RNP complex that does not reflect a physiologically functional state of telomerase RNP. We note that even under optimized conditions, whether using crude *Tetrahymena* extract or extract enriched for telomerase, antibodies against p80 or p95 immunoprecipitated a much smaller fraction of the total enzyme pool than did antibodies against p45 or p65 shown here. This comparison indicates that p80 and p95 are not stable holoenzyme components in the same manner as are p45 and p65. The litmus test of immunodepletion remains to be performed for p20 and p75, if appropriate antibody reagents can be produced. Finally, we note that neither p65 nor p45 has appreciable sequence homology to full-length Est1p or the motifs conserved among EST1 proteins (for review, see Lundblad 2003). The ongoing efforts in *Tetrahymena* genome sequencing, assembly, and annotation will be helpful in determining whether this ciliate encodes a potential Est1p ortholog.

Materials and methods

Tetrahymena growth and transformation

Cells were grown in 2% proteose peptone, 0.2% yeast extract, 10 μ M FeCl₃ with 250 μ g/mL ampicillin and streptomycin and 1.25 μ g/mL Fungizone (amphotericin B). When indicated, cells were starved in 10 mM Tris-HCl (pH 7.5) for 12–24 h. All transformations were performed on starved cells by particle bombardment (Cassidy-Hanley et al. 1997). For transformations into the *BTU1* locus of CU522, cells were allowed to recover for 2–5 h at 30°C then were selected in 20–30 μ M paclitaxel (Sigma). Cells were passaged to progressively higher paclitaxel concentrations for 2–3 d until cells were growing in 50 μ M paclitaxel, after which they were passaged daily in 50 μ M paclitaxel for 5 d. Transgenes encoding p45-TAP and p65-TAP were transformed into CU522 strains partially assorted for p45 and p65 gene disruptions. For *TERT* gene knockout, cells were transformed and assorted as described previously (Miller and Collins 2000). Cells were subjected to maximal selective pressure by stepwise passaging to the maximum tolerated paromomycin concentration, continued passaging at this drug concentration for 16 d (~100 doublings), then passaging for 8 d (~50 doublings) in the absence

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of selection. For *TAP45* and *TAP65* gene knockdowns, cells were grown for ~150 doublings in the presence of paromomycin and 100 doublings after release from drug selection.

Transformation constructs

The C-terminal TAP tag was fused to the synthetic TERT gene (Collins and Gandhi 1998) by subcloning of PCR-amplified tag sequence from the vector pRSAB1234 (a gift from Erin O'Shea, University of California, San Francisco). The entire TERT-TAP coding region was cloned into pBICH3 (a gift from Jacek Gaertig, University of Georgia, Athens) with *NsiI* and *BamHI*, inserting the sequence between the *BTU1* promoter and 3'-flanking genomic sequence. Expression constructs for p65-TAP and p45-TAP were constructed as for TERT-TAP except that the wild-type genomic loci were used as the coding regions. The *TRT* gene disruption construct, as described (Miller and Collins 2000), consisted of a neomycin-resistance expression cassette designed to recombine into the *TRT* locus, replacing a segment of the gene. The *TAP45* gene disruption construct contains the same neomycin-resistance expression cassette cloned between an ~520-bp and an ~580-bp segment of the *TAP45* gene, designed to replace the encoded protein from V150 to E215. Similarly, the *TAP65* disruption construct contains the neomycin-resistance cassette cloned between an ~600-bp and an ~735-bp segment of the *TAP65* gene, replacing the region encoding G181 to Q302.

Protein, RNA, DNA, and activity assays

To make whole-cell extracts, starved cells were lysed with 0.2% NP-40 in T2MG buffer (20 mM Tris-HCl at pH 8.0, 1 mM MgCl₂, 10% glycerol) in the presence of protease inhibitors and β-mercaptoethanol, and then centrifuged at 130,000 × g for 1 h. Affinity-purified rabbit polyclonal antibody raised against a peptide from the *Tetrahymena* TERT motif AB-loop region was used for immunoblot detection (Collins and Gandhi 1998). Activity assays were performed by direct primer extension in reaction buffer with 50 mM Tris-acetate (pH 8.0), 10 mM spermidine, 5 mM β-mercaptoethanol, 2 mM MgCl₂, 0.4 mM dTTP, 0.3 μM [α-³²P]dGTP (800 Ci/mmol, NEN), and 0.4 μM DNA primer (G₄T₂)₃. Samples from affinity purification were washed or diluted into T2MG to make up half of the activity assay reaction volume, such that elution buffer made minimal contribution to the final reaction conditions. Assays were incubated for 1 h at 30°C.

Total RNA was prepared by standard methods (Ausubel et al. 1996). For RNA 3'-end labeling, RNA isolated from the affinity purification EGTA elution was incubated with 0.4 μM cordycepin (5000 Ci/mmol; PerkinElmer Lifesciences) and 120 U of poly(A) polymerase (USB) for 20 min at 30°C, extracted with phenol/chloroform and then resolved by electrophoresis on a 9% polyacrylamide denaturing gel. Small RNA Northern blots were probed with 5'-end-labeled DNA oligonucleotides complementary to the intended RNA target. Genomic DNA isolation was performed as described previously (Ahmed et al. 1998). Southern blots to assess integration events were analyzed with random hexamer-primed probes corresponding to the genomic region immediately 5' of the integrated transgene. Telomeric Southern blot hybridization was performed as described (Miller and Collins 2000), except that DNA was separated on a 4% acrylamide or 0.7% agarose gel.

Affinity purification

For large-scale purification, 800 mL of extract was used corresponding to ~200 L of starved cells at 2–4 × 10⁵ cells/mL. Ex-

tracts were adjusted to 0.1 M NaCl and incubated in batch with IgG agarose (Sigma) for 1–2 h at room temperature. Resin was washed three times in wash buffer (WB: 20 mM Tris-HCl at pH 8.0, 2 mM MgCl₂, 0.2 mM EGTA, 0.1 M NaCl, 0.1% NP-40, 10% glycerol, 1 mM DTT, 0.1 mM PMSF) and twice in TEV protease elution buffer (10 mM Tris-HCl at pH 8.0, 0.1 M NaCl, 0.1% NP-40, 0.5 mM EDTA, 1 mM DTT). Elution was accomplished by incubation in elution buffer for 1–1.5 h mixing at room temperature with recombinant TEV protease. The elution was adjusted to 3 mM CaCl₂ and mixed with 3 volumes of calmodulin binding buffer (CBB: 10 mM Tris-HCl at pH 8.0, 0.1 M NaCl, 1 mM magnesium acetate, 1 mM imidazole, 2 mM CaCl₂, 0.1% NP-40, 10 mM β-mercaptoethanol), then incubated with calmodulin affinity resin (Stratagene) for 1 h at 4°C. Resin was washed three times in CBB then eluted with three sequential additions of calmodulin elution buffer (10 mM Tris-HCl at pH 8.0, 0.1 M NaCl, 1 mM magnesium acetate, 1 mM imidazole, 0.1% NP-40, 10 mM EGTA, 10 mM β-mercaptoethanol). For large-scale purifications, samples were precipitated with trichloroacetic acid. Proteins were resolved by SDS-PAGE, transferred to nitrocellulose, stained with Ponceau S, excised, and digested with Lys-C protease. Peptides were separated by HPLC and sequenced by Edman degradation.

For immunodepletion, affinity-purified rabbit polyclonal antibodies raised against full-length p45 and p65 and control rabbit polyclonal antibodies were prebound to Protein A Sepharose (Amersham Biosciences) and then incubated with whole-cell extract adjusted to 50 mM NaCl for 1 h at 4°C for each round of immunodepletion. Antibody resin was subsequently washed in WB as above except for the presence of 50 mM NaCl.

GenBank accession numbers

The accession numbers for the sequences reported here are AY280525 (*TAP45*) and AY280524 (*TAP65*).

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Holoenzyme proteins required for the physiological assembly and activity of telomerase

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