

# Shared Subunits of *Tetrahymena* Telomerase Holoenzyme and Replication Protein A Have Different Functions in Different Cellular Complexes\*

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In most eukaryotes, telomere maintenance relies on telomeric repeat synthesis by a reverse transcriptase named telomerase. To synthesize telomeric repeats, the catalytic subunit telomerase reverse transcriptase (TERT) uses the RNA subunit (TER) as a template. In the ciliate *Tetrahymena thermophila*, the telomerase holoenzyme consists of TER, TERT, and eight additional proteins, including the telomeric repeat single-stranded DNA-binding protein Teb1 and its heterotrimer partners Teb2 and Teb3. Teb1 is paralogous to the large subunit of the general single-stranded DNA binding heterotrimer replication protein A (RPA). Little is known about the function of Teb2 and Teb3, which are structurally homologous to the RPA middle and small subunits, respectively. Here, epitope-tagging Teb2 and Teb3 expressed at their endogenous gene loci enabled affinity purifications that revealed that, unlike other *Tetrahymena* telomerase holoenzyme subunits, Teb2 and Teb3 are not telomerase-specific. Teb2 and Teb3 assembled into other heterotrimer complexes, which when recombinantly expressed had the general single-stranded DNA binding activity of RPA complexes, unlike the telomere-specific DNA binding of Teb1 or the TEB heterotrimer of Teb1, Teb2, and Teb3. TEB had no more DNA binding affinity than Teb1 alone. In contrast, heterotrimers reconstituted with Teb2 and Teb3 and two other *Tetrahymena* RPA large subunit paralogs had higher DNA binding affinity than their large subunit alone. Teb1 and TEB, but not RPA, increased telomerase processivity. We conclude that in the telomerase holoenzyme, instead of binding DNA, Teb2 and Teb3 are Teb1 assembly factors. These findings demonstrate that *Tetrahymena* telomerase holoenzyme and RPA complexes share subunits and that RPA subunits have distinct functions in different heterotrimer assemblies.

Telomeres, which are the DNA-protein complexes at the ends of eukaryotic chromosomes, are essential for genome sta-

bility and long term cellular proliferation (1, 2). Generally, telomeric DNA is composed of simple sequence repeats arranged as a tract of duplex repeats followed by a single-stranded 3' overhang (3). These telomeric repeats recruit sequence-specific double-stranded and single-stranded DNA-binding proteins to nucleate the assembly of telomere-specific protein complexes, which sequester chromosome termini from DNA damage sensors (3, 4). The accessibility of strand termini is strictly regulated, and as a consequence, the 3' overhang has a fixed length range in any given species. This 3' overhang is critical for telomere end protection, but it must be created anew after genome replication in a manner that obliges a loss of telomeric repeats with each round of cell division (5). Single-celled organisms have a relatively short telomeric 3' overhang and consequently lose a few or tens of base pairs per cell division, whereas human cells have relatively long overhangs on the order of ~100 nucleotides (nt)<sup>3</sup> and correspondingly lose more base pairs of telomeric repeats per cell division (6, 7).

To compensate for incomplete telomere replication by conventional DNA polymerases, most eukaryotes rely on the ribonucleoprotein (RNP) telomerase (8). Each telomeric repeat array is maintained in a dynamic equilibrium of attrition from genome replication and telomerase-mediated *de novo* synthesis. Telomerase acts by reverse transcribing the integral RNA component, TER, with the catalytic telomerase reverse transcriptase protein, TERT (9, 10). By copying a short template sequence within its RNA moiety, telomerase synthesizes the guanosine-rich telomeric DNA strand (G-strand) running 5' to 3' toward a chromosome terminus (*e.g.* repeats of TTGGGG in the ciliate *Tetrahymena* or TTAGGG in vertebrates). TERT and TER assembled in a heterologous cell extract can reconstitute repeat synthesis activity; therefore, an RNP with these two subunits is considered the minimal recombinant RNP (11, 12). For biologically functional telomerase holoenzyme, TER and TERT require a number of other subunits to properly fold TER, assemble TER with TERT, and allow active RNP to elongate telomeres (13, 14). Although telomerase holoenzyme sub-

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<sup>3</sup> The abbreviations used are: nt, nucleotide(s); RPA, replication protein A; RNP, ribonucleoprotein; G-strand, guanosine-rich telomeric DNA strand; OB-fold, oligonucleotide/oligosaccharide-binding fold; C-strand, cytidine-rich telomeric DNA strand; RAP, repeat addition processivity; RRL, rabbit reticulocyte lysate; ZZ, tandem protein A domains; F, triple FLAG peptide; MBP, maltose-binding protein; CTαH, C-terminal α-helix; Ni-NTA, nickel-nitrilotriacetic acid.

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units are evolutionarily divergent in sequence, studies across model organisms have illuminated recurrent functionalities for holoenzyme proteins in RNA stabilization, intracellular RNP trafficking, and RNP recruitment to telomeres (15, 16).

Telomerase binds a chromosome 3' overhang in competition, and also coordination, with other single-stranded DNA (ssDNA)-binding proteins (17, 18). Throughout most of the cell cycle, the telomere 3' overhang is sequestered by DNA binding and telomere remodeling activities of the ssDNA-binding protein Pot1 (18, 19). The Pot1 N-terminal pair of oligonucleotide/oligosaccharide-binding fold (OB-fold) domains interacts sequence-specifically with the telomeric repeat G-strand, whereas the Pot1 C-terminal region interacts with vertebrate TPP1/fission yeast Tpz1/*Tetrahymena* Tpt1 (19, 20). TPP1 and Tpz1 bridge Pot1 with proteins assembled on the double-stranded telomeric DNA repeats (21).

Telomeric repeat ssDNA is also bound, at least transiently, by the general ssDNA-binding RPA heterotrimer of ~70-kDa Rpa1, ~30-kDa Rpa2, and ~15-kDa Rpa3. RPA serves essential roles in DNA replication and repair, recruiting myriad cellular factors to bound ssDNA with specificities that are incompletely understood (22, 23). The RPA heterotrimer has six OB-fold domains, four of which contact DNA: domains A, B, and C in the large subunit Rpa1 and domain D in the middle subunit Rpa2 (Fig. 1A). DNA-binding domains A and B initially engage 8–10 nt of ssDNA, and then subsequent DNA binding by domains C and D extends the footprint to ~30 nt (23, 24). Domains A–D are oriented from 5' to 3' on ssDNA, with interdomain contacts and linker structuring induced by DNA binding (25). Despite high DNA binding affinity from this interdomain cooperation, RPA can diffuse along a bound DNA by a series of individual domain dissociations (23, 24, 26).

Telomeric repeat ssDNA is also bound by the RPA-like CST complex, which is composed of large subunit Cdc13 (in budding yeast), CTC1 (in vertebrate cells), or p75 (in *Tetrahymena*; see below) as well as the middle and small subunits Stn1 and Ten1, respectively. CST has evolutionarily variable ssDNA binding properties and variable biological roles linked to a high degree of large subunit divergence (18, 27). Vertebrate CST contributes to DNA replication at sites throughout the genome and, with distinct structural requirements, to telomere-specific processes, such as the post-replication cytidine-rich strand (C-strand) fill-in by polymerase  $\alpha$ -primase (28–30). Vertebrate CST also has been proposed to inhibit telomerase access to chromosome ends, although this role is not uniformly evident across different studies (30, 31). Budding yeast CST function is telomere-specific; it stimulates C-strand fill-in and contributes to chromosome end-capping, and, when disassembled in S-phase, its Cdc13 subunit recruits telomerase holoenzyme (18). A single OB-fold domain within budding yeast Cdc13 is necessary and sufficient for sequence-specific recognition of G-strand ssDNA (32), whereas detectable binding of vertebrate or *Tetrahymena* CST to DNA requires all three subunits (28, 31, 33). *Tetrahymena* CST assembles as a stable subcomplex of the telomerase holoenzyme, where it is proposed to couple G-strand synthesis to C-strand fill-in (33, 34).

Remarkably, the *Tetrahymena* telomerase holoenzyme contains another RPA-like heterotrimer in addition to CST: the

TEB heterotrimer composed of Teb1, Teb2, and Teb3 (34)(Fig. 1B). The holoenzyme RNP catalytic core (TERT, TER, and the RNA-binding protein p65) interacts with the central hub protein p50 (with an OB-fold domain structurally and functionally related to TPP1), which in turn binds independently to the CST and TEB heterotrimers (34–36). The RPA-like TEB complex includes large subunit Teb1, middle subunit Teb2, and small subunit Teb3 (34, 37). Teb1 has an N-terminal OB-fold domain that does not contribute to DNA binding, two central DNA-binding OB-fold domains (Teb1A and Teb1B), and a C-terminal OB-fold domain (Teb1C) that improves DNA binding by Teb1AB (37, 38). Teb1A and Teb1B each bind sequence-specifically to the G-strand of *Tetrahymena* telomeric repeats (38, 39). Teb1C interacts with p50, probably threading ssDNA between the telomerase active site and Teb1AB (34, 35). In cells, high affinity ssDNA binding by Teb1 is a major determinant of telomerase association to telomeres. Teb1C mutations that disrupt p50 interaction do not reduce Teb1 binding to telomeric DNA, but Teb1 interaction with p50 is required for robust telomere association of the rest of the telomerase holoenzyme (40).

In comparison with Teb1, little is known about the function(s) of Teb2 or Teb3. A structural model from cryo-electron microscopy (34) indicates that the Teb2 OB-fold domain (the N-terminal half of the protein) and the Teb3 OB-fold domain (the full-length protein) interact with Teb1C to form the heterotrimer core, which is stabilized by RPA-like bundling of the  $\alpha$ -helices immediately following the OB-fold domains (Fig. 1A). Teb1 alone or TEB heterotrimer dramatically increases telomerase repeat addition processivity (RAP), but for proteins expressed in rabbit reticulocyte lysate (RRL), TEB mediates this stimulation more effectively than Teb1 (34, 36, 37). We suggested previously that Teb2 and Teb3 could be *Tetrahymena* RPA subunits as well as subunits of telomerase holoenzyme (34). Phylogenetic analysis grouped Teb2 with Rpa2 and Teb3 with Rpa3 proteins of other species (34). Furthermore, Teb2 and Teb3 have mRNA expression levels higher than Teb1 and the other telomerase-specific holoenzyme subunits, approaching the mRNA level of the previously characterized RPA large subunit Rpa1 (34, 38). Together, these observations raise the hypothesis that Teb2 and Teb3 could be shared subunits of telomerase holoenzyme and RPA.

To test this hypothesis and to better understand the function(s) of Teb2 and Teb3 in cells, we investigated their endogenous interaction partners by purification of cellular complexes. These studies and complementary ssDNA-binding assays performed with purified recombinant heterotrimers establish that Teb2 and Teb3 are subunits of telomerase holoenzyme and *Tetrahymena* RPA complexes. Curiously, the same proteins make different functional contributions to different heterotrimers. Understanding the complexity of RPA and RPA-like complexes in *Tetrahymena* provides a precedent for studies of alternative RPA subunits in other organisms, including humans (41–43).

## Results

*Teb2 and Teb3 Are Not Telomerase-specific Proteins*—To further our functional understanding of Teb2 and Teb3, we

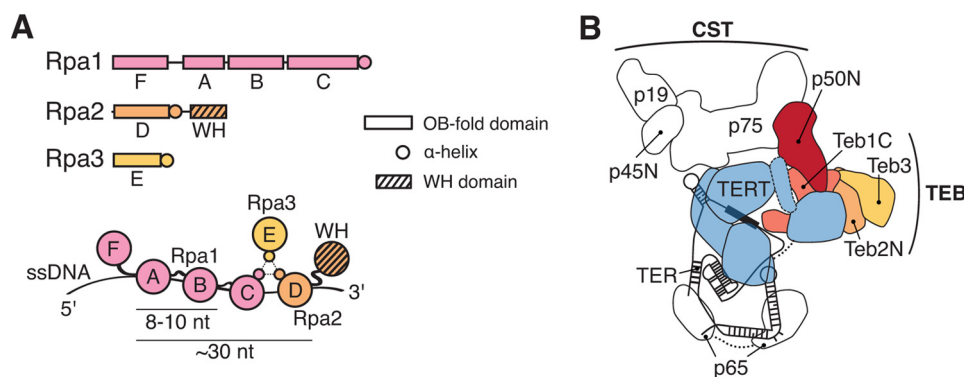


FIGURE 1. RPA domains and TEB subunit organization in telomerase holoenzyme. *A*, schematic diagram of RPA subunit domains (*top*) and their DNA contact (*bottom*). *B*, simplified diagram of *Tetrahymena* telomerase holoenzyme architecture (34). Subunits are colored as follows: TERT (blue), p50 N-terminal domain (red), Teb1C (pink), Teb2 N-terminal domain (gold), and Teb3 (yellow). The remaining holoenzyme proteins and TER are labeled without color; two p65 RNA-binding domains are connected by a dashed line, and only the N-terminal OB-fold domain of p45 (p45N) is included in the model.

sought to characterize endogenously assembled complexes containing these proteins. In addition to the telomerase holoenzyme (Fig. 1*B*), Teb2 and Teb3 could be subunits of RPA or RPA-like complexes likely to be more abundant than telomerase. We performed unbiased affinity purification of Teb2 and Teb3 using N-terminally tagged versions of the proteins expressed from their endogenous gene loci. We targeted each endogenous gene locus to insert an N-terminal tag of tandem Protein A domains (ZZ) and a triple FLAG peptide (F), optimized for affinity purifications from *Tetrahymena* cell extracts (37). The integrated construct also contained a blasticidin resistance cassette (Fig. 2*A*), allowing for a standard protocol of assortment to maximal recombinant chromosome copy number in replacement of the endogenous locus chromosome (44). Cells released from selection were used for genomic DNA analysis to discriminate whether the recombinant chromosome had entirely substituted for the wild-type chromosome. Southern blotting hybridization confirmed full macronuclear replacement of endogenous chromosomes with ZZ-F-Teb2 or ZZ-F-Teb3 chromosomes (Fig. 2*B*). The silent micronuclear gene locus gives a very faint wild-type locus signal that is discriminated as micronuclear because it does not rebound in copy number with release of cells from selection in blasticidin (44).

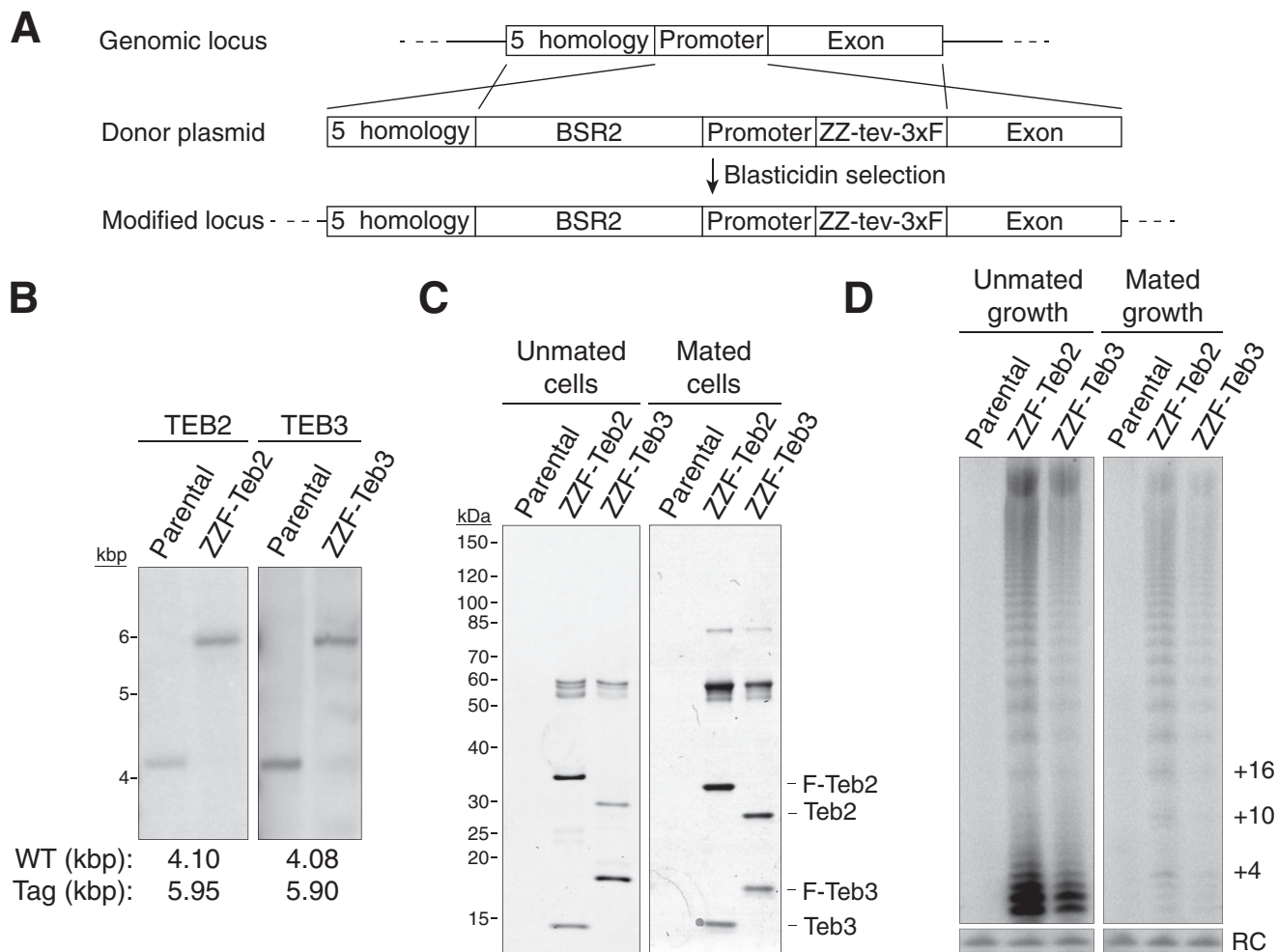
We performed tagged protein affinity purifications from extracts of cells in synchronized, asexual (vegetative) growth or mated cells in the process of sexual reproduction, two conditions that demand high rates of new DNA synthesis (45). Tandem affinity purification of ZZ-F-Teb2 or ZZ-F-Teb3 from whole-cell lysates co-purified several polypeptides not recovered in parallel mock purifications from cell extracts without tagged protein (Fig. 2*C*). Proteins with the SDS-polyacrylamide gel mobilities of F-Teb2 and Teb3 or F-Teb3 and Teb2 were readily detectable (note that the ZZ portion of the tag was removed during purification). Aside from Teb2 and Teb3, the proteins in the purifications did not co-migrate with telomerase holoenzyme proteins (37), such as the ~130-kDa TERT. Although it contributes a minority of the total associated proteins, telomerase holoenzyme was co-purified with F-Teb2 and F-Teb3, as judged by specific enrichment of telomerase activity assayed using direct telomeric primer extension with radiolabeled dGTP and dTTP (Fig. 2*D*). Telomerase enriched by ZZ-F-Teb2 or ZZ-F-Teb3 had more low RAP activity with the purifi-

cations from cells in vegetative growth *versus* the purifications from mated cells, which, based on our experience, is likely to reflect more proteolysis in extracts of growing cells.

To identify unknown proteins that interact with Teb2 and Teb3, we submitted the entire pool of proteins associated with F-Teb2 or F-Teb3 for mass spectrometry (MS), using the purifications from mated cells. Proteins detected by MS in the mock purification from the parental strain were subtracted from the list of proteins specifically associated with Teb2 or Teb3, which we then rank-ordered by number of unique sequence peptides (Table 1). Both purifications co-enriched *Tetrahymena* Rpa1 as the top-ranked associated protein. We had previously characterized Rpa1 as a general ssDNA-binding subunit genetically essential for *Tetrahymena* growth (38). In addition to Rpa1, the Teb2 purification co-enriched Teb3, and the Teb3 purification co-enriched Teb2. The representation of Rpa1, Teb2, and Teb3 clustered them together as a top-ranking group (Table 1). Also well represented were likely RPA-interacting factors involved in DNA repair (22, 23), including DNA mismatch repair proteins (*Tetrahymena* gene names *MSH2* and *MSH6*) and other DNA repair factors (Ku80 and Ku70). The telomerase holoenzyme proteins p75, p65, p50, and p45 were detected at low representation in the Teb2 purification (Table 1), which had a higher yield of total protein than the Teb3 purification in the samples used for MS. Due to the scarce amount of telomerase compared with DNA replication and repair factors, it is not surprising that only a subset of telomerase holoenzyme subunits was detected in only the higher yield affinity purification.

Interestingly, Teb3 co-purified two RPA-like proteins not associated with Teb2: the RPA large subunit paralog Rlp1 (RPA-like protein 1), which we previously characterized in parallel with Rpa1 in cells and as recombinant protein (38), and a putative RPA middle subunit paralog THERM\_00459400, here designated Rlp2 (RPA-like protein 2). Rlp1 has a sequence and predicted domain structure similar to Rpa1 and Teb1, except that Rlp1 lacks a regulatory N-terminal OB-fold domain (Fig. 3*A*). Unlike Rpa1 and Teb1, Rlp1 was not genetically essential for cell growth (37, 38). Rlp2 is a hypothetical protein predicted from genome sequence (46). BLAST of Rlp2 against all sequences in GenBank<sup>TM</sup> identified *Tetrahymena* Teb2 and the protist *Phytophthora infestans* Rpa2 in the top 10 scores, the remainder of which were hypothetical proteins, including a

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**FIGURE 2. *Teb2* and *Teb3* are not telomerase-specific.** *A*, genomic locus targeting strategy to generate ZZF-Teb2 and ZZF-Teb3 cell lines. *B*, genomic DNA Southern blots showing macronuclear replacement of an endogenous locus with the recombinant locus. *C*, affinity purifications from extracts of cells in synchronized vegetative growth (*left*) or mated cells (*right*) profiled by colloidal Coomassie staining after SDS-PAGE. *D*, telomerase activity assays for the purifications shown in *C*. RC, recovery control for telomerase product precipitation. Numbers of nt added to the primer to complete the first three telomeric repeats are indicated at the *right*.

marine ciliate protein annotated only as “nucleic acid-binding OB-fold.” Several domain prediction methods confirm that Rlp2 could harbor an OB-fold, which will require future direct structural analysis to confirm. Because both *Teb2* and *Teb3* co-purified Rpa1 but only *Teb3* co-purified Rlp1 and Rlp2, we suggest that *Tetrahymena* cells assemble at least two RPA or RPA-like heterotrimers beyond the TEB and CST complexes of telomerase holoenzyme: an abundant RPA complex containing Rpa1, *Teb2*, and *Teb3* and complex(es) containing Rlp1, Rlp2, and *Teb3* (see “Discussion”). Cells could also assemble a complex of Rlp1, *Teb2*, and *Teb3* that we did not detect by MS. An Rlp1, *Teb2*, and *Teb3* heterotrimer could be low in abundance or not assembled at the specific state of sexual reproduction that cells were harvested in for large scale purifications. Because the goal of this work is to characterize *Teb2* and *Teb3*, we compared heterotrimers containing these two subunits and each alternative large subunit. For convenience, we will refer to a recombinant complex containing Rlp1, *Teb2*, and *Teb3* as RTT.

*Teb2* and *Teb3* Contribute to the DNA Binding Affinity of RPA but Not TEB—We next investigated the DNA binding properties of the RPA large subunit paralogs *Teb1*, Rpa1, and

Rlp1 with or without co-assembled *Teb2*-*Teb3*. We expressed and purified N-terminally six-histidine ( $\text{His}_6$ )-tagged *Teb1*, Rpa1, and Rlp1 from *Escherichia coli* either alone, as done previously for *Teb1* and Rpa1 (38), or co-expressed with maltose-binding protein (MBP)-tagged *Teb2* and untagged *Teb3* (Fig. 3A). MBP-*Teb2* and *Teb3* were also co-expressed in the absence of an RPA large subunit paralog.  $\text{His}_6$ -tagged large subunit proteins were purified using Ni-NTA resin. *Teb2*-*Teb3* complex was purified using amylose resin. Heterotrimer complexes were purified using Ni-NTA resin and then amylose resin in series. Overall, these recombinant protein purifications (Fig. 3B) indicate that each large subunit protein can assemble with *Teb2*-*Teb3*, at least in the absence of competing cellular factors. Furthermore, *Teb1*, Rpa1, and Rlp1 each formed complexes with roughly stoichiometric amounts of MBP-*Teb2* and *Teb3* (Fig. 3B).

To investigate the functional contribution of *Teb2*-*Teb3* to each heterotrimer, we tested the recombinant protein preparations above for binding to a panel of four ssDNA oligonucleotides differing in length (18 or 30 nt) and in telomeric versus non-telomeric sequence (Fig. 3C). In electrophoretic mobility shift assays, the *Teb2*-*Teb3* complex had undetectable DNA

TABLE 1

## Proteins co-purified with Teb2 or Teb3 as detected by MS

The column labeled "ZZF" indicates the tagged protein used for affinity purification. Listing order of MS-identified proteins associated with ZZF-Teb2 or ZZF-Teb3 is by decreasing number of unique-sequence peptides (sequence count) detected for the top 15 associated proteins, followed by any telomerase holoenzyme subunits. Teb2 recovered 70 total proteins and 35 hits with more than one peptide read. Two additional proteins fall between p75 and the top 15 associated proteins, whereas 28 additional proteins fall between p45 and the top 15 associated proteins. Teb3 recovered 40 total proteins and 24 with more than one peptide read. Boldface type indicates an RPA subunit paralog or telomerase holoenzyme protein.

ZZF	Locus	Gene name	Description	Sequence count	Protein coverage
					%
Teb2	THERM_00106890	<i>RFA1</i>	Replication protein A large subunit	73	53.8
	THERM_001113129	<i>TEB2</i>	<b>Teb2</b>	32	39.8
	THERM_00439320	<i>TEB3</i>	<b>Teb3</b>	19	47.6
	THERM_00295920	<i>MSH2</i>	MutS domain III family protein	17	20.2
	THERM_00194810	<i>MSH6</i>	MutS domain III family protein	17	14.4
	THERM_00502600	<i>PARP6</i>	WGR domain-containing protein	11	6.4
	THERM_00046920	<i>RVB2</i>	DNA helicase RBV2 homolog	8	21.3
	THERM_00865240	None	DNA ligase I	8	15.6
	THERM_00125640	<i>SSA3</i>	HSP70a paralog	8	14.1
	THERM_00558440	<i>SSA5</i>	HSP70a paralog	8	11.7
	THERM_00216140	None	DNA topoisomerase family protein	8	9.6
	THERM_00633360	<i>HTB1</i>	Histone H2B	7	29.5
	THERM_00492460	<i>TKU80</i>	Ku80 ortholog	7	12.2
	THERM_00105110	<i>HSP70</i>	Putative 70-kDa heat shock protein	7	11.8
	THERM_00043780	<i>POLN1</i>	DNA polymerase I family protein	7	8.0
	THERM_00059040	<i>TAP75</i>	<b>p75</b>	3	8.2
	THERM_00083360	<i>TAP45</i>	<b>p45</b>	1	4.3
	THERM_000318539	<i>TAP65</i>	<b>p65</b>	1	2.6
	THERM_01049190	<i>TAP50</i>	<b>p50</b>	1	2.4
	Teb3	THERM_00106890	<i>RFA1</i>	Replication Protein A large subunit	70
THERM_001113129		<i>TEB2</i>	<b>Teb2</b>	42	57.6
THERM_00439320		<i>TEB3</i>	<b>Teb3</b>	28	92.4
THERM_00194810		<i>MSH6</i>	MutS domain III family protein	21	18.8
THERM_00216140		None	DNA topoisomerase family protein	19	21.7
THERM_00672190		<i>THD13</i>	Histone deacetylase 13	17	11.8
THERM_00316500		<i>HTA2</i>	Histone H2A	7	24.8
THERM_00295920		<i>MSH2</i>	MutS domain III family protein	7	13.7
THERM_00726370		<i>RLP1</i>	RPA-like protein 1	5	9.7
THERM_00502600		<i>PARP6</i>	WGR domain-containing protein	4	2.1
THERM_00691710		None	Toprim domain-containing protein	4	3.3
THERM_00561799		<i>TKU70</i>	Ku70 ortholog	4	5.1
THERM_00070820		None	U-box domain containing protein	4	8.9
THERM_00459400		<i>RLP2</i>	RPA-like protein 2	4	15.2
THERM_00328620		None	Hypothetical protein	4	17.1

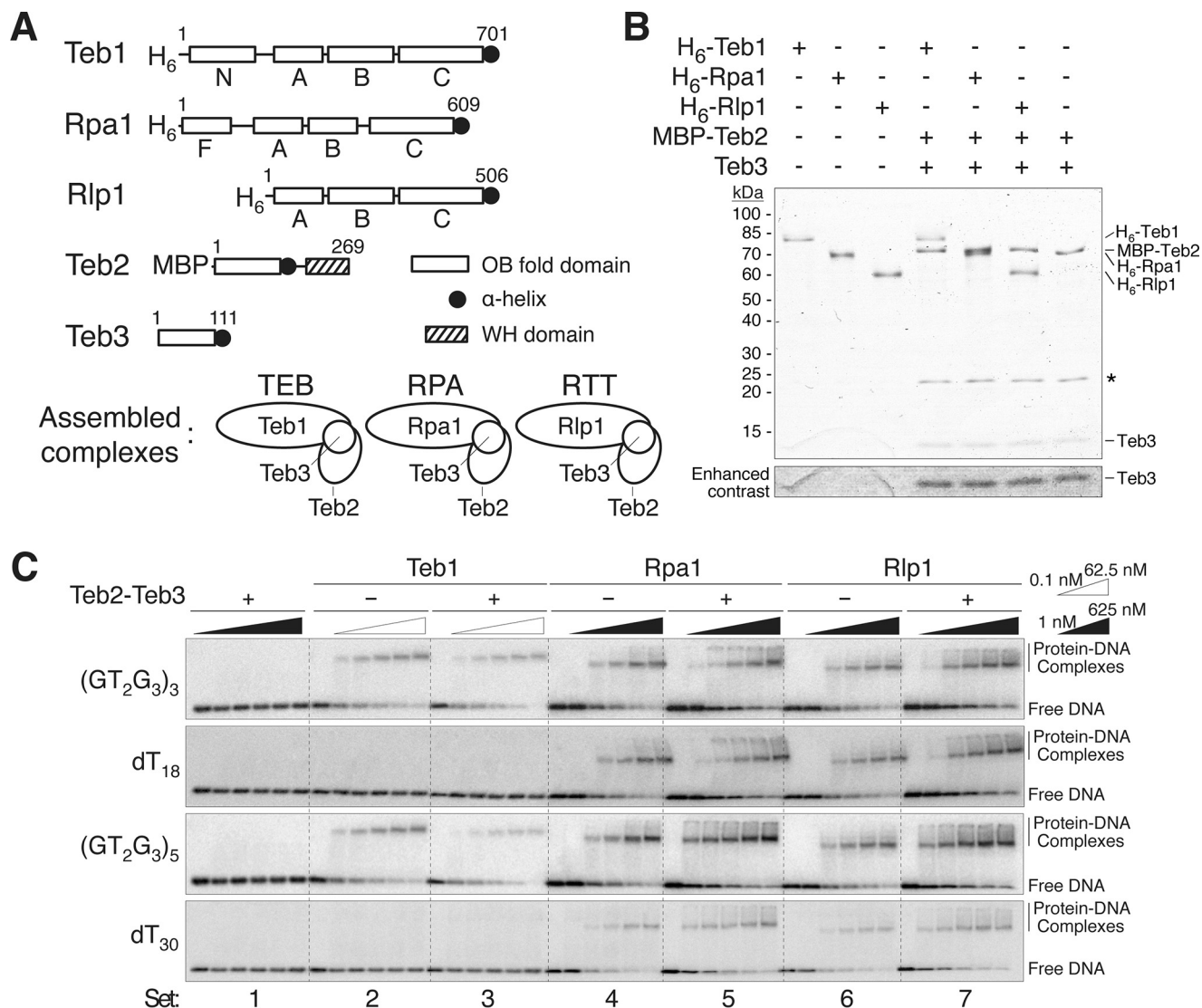
binding affinity for any ssDNA (Fig. 3C, set 1). This finding parallels results observed for human Rpa2 and Rpa3 (47). Teb1 alone bound 18- and 30-nt telomeric ssDNAs with similar affinity and did not bind either length of polythymidine (Fig. 3C (set 2) and Table 2), as expected from previous observations (38). Also consistent with previous assays (38), Rpa1 alone bound all four ssDNAs with approximately equal affinity (Fig. 3C (set 4) and Table 2), as did Rlp1 (Fig. 3C (set 6) and Table 2). The TEB complex had DNA binding properties indistinguishable from Teb1 alone, even on the longer ssDNAs (Fig. 3C (set 3) and Table 2), indicating very little or no contribution of Teb2-Teb3 to DNA contact by TEB. In contrast, RPA and RTT each bound 30-nt ssDNA with higher affinity than the corresponding large subunit alone (5–7-fold for RPA, 3–4-fold for RTT; Fig. 3C (sets 4–7) and Table 2). Binding of RPA and RTT to the shorter 18-nt ssDNAs was not improved relative to the corresponding large subunit alone (Fig. 3C (sets 4–7) and Table 2). These results strongly support the conclusion that Teb2 and Teb3 are canonical RPA subunits when in complex with Rpa1; they increase DNA binding affinity and in parallel extend the length of ssDNA that contributes to protein interaction (Fig. 1A) (23, 24).

**Teb2-Teb3 Function in Telomerase Holoenzyme Is Dependent on Teb1**—When expressed in RRL, Teb2 and Teb3 without Teb1 did not stimulate telomerase activity or RAP (34). The very low amount of protein produced in RRL and its uncertain

folding presented caveats to the interpretation of this finding. Bacterially expressed proteins generated in this study offered new opportunity to investigate whether Teb2 and Teb3 directly contribute to RAP stimulation. We also sought to investigate whether *Tetrahymena* RPA, like TEB, stimulates telomerase RAP *in vitro*. RRL-assembled telomerase RNP catalytic core was combined with p50 and CST (p75, p45, p19) containing C-terminally F-tagged p45, which has biological function (37). The telomerase subunit complex was purified from RRL using anti-FLAG antibody resin and mixed with purified Teb2-Teb3, Teb1, TEB, Rpa1, RPA, Rlp1, or RTT added at final concentrations of 40 or 200 nM. These reconstitutions were then assayed for telomerase product synthesis by direct primer extension. Human RPA (48) or *Tetrahymena* RPA (data not shown) added at high concentration to an activity assay sequestered primer from telomerase. Here we sought to test the functional interplay of elongating telomerase and the ssDNA binding factors, so we used a 200 nM concentration of an 18-nt telomeric repeat primer with 40 or 200 nM added protein or protein complex.

The addition of Teb1 or TEB converted the RNP catalytic core with p50 to high RAP activity (Fig. 4, lanes 1 and 3–6). The addition of the Teb2-Teb3 complex alone had no influence (Fig. 4, lane 2). The addition of Rpa1 did not affect activity or RAP, consistent with previous studies (38), and neither did the addition of RPA (Fig. 4, lanes 7–10). For Rlp1 or RTT, protein(s)

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**FIGURE 3. Comparison of DNA binding by Teb1, TEB, Rpa1, RPA, Rlp1, and RTT.** *A*, domain structures for Teb1, Rpa1, Rlp1, Teb2, and Teb3 and complexes formed by co-assembly that were assayed for DNA binding affinity. *B*, colloidal Coomassie staining after SDS-PAGE of proteins and complexes used for DNA binding assays. \*, a background protein in the Teb2-Teb3 purification. Enhanced contrast for the bottom of the gel is shown below the main gel panel. *C*, DNA binding by Teb2-Teb3, Teb1, TEB, Rpa1, RPA, Rlp1, and RTT. A fixed concentration of the indicated end-labeled ssDNA oligonucleotide (~10  $\mu$ M) was incubated with protein or protein complex added at steps of final concentration over the indicated range of 0.1–62.5 or 1–625 nM.

added at a concentration equimolar with the 200 nM DNA primer resulted in modest inhibition of overall activity but no change in RAP (Fig. 4, lanes 11–14). Thus, only Teb1 or TEB stimulates RAP, not other ssDNA binding proteins or protein complexes containing Teb2 and Teb3. These activity assays, along with the gel mobility shift assays, suggest that the role of Teb2-Teb3 in telomerase holoenzyme is indirect through Teb1.

**TEB Heterotrimer Formation Enhances Teb1 Assembly in Telomerase Holoenzyme**—We turned to the hypothesis that Teb2 and Teb3 favor a conformation of Teb1 optimal for its holoenzyme assembly and high RAP stimulation. If Teb2-Teb3 function is indirect through Teb1, it would be dependent on TEB heterotrimer assembly. In RPA, the C-terminal  $\alpha$ -helix (CT $\alpha$ H) of Rpa1C, Rpa2 OB-fold domain, and Rpa3 together form the trimerization interface (49). Similarly, the OB-fold domains of Teb1C, Teb2, and Teb3 are each followed by an  $\alpha$ -helix that forms their trimerization interface (Fig. 5, *A* and *B*),

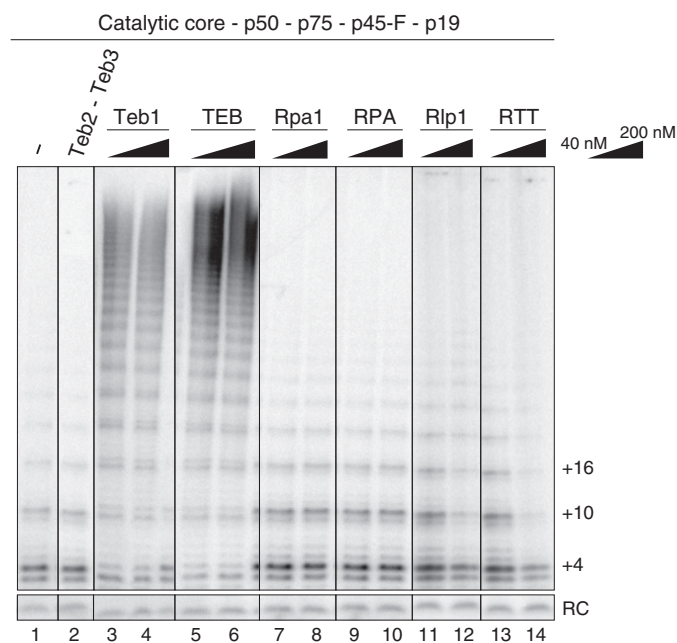
which is on the far side of Teb1C from the contact surface with TERT (34). Reconstitution of high RAP telomerase activity in assays with bacterially expressed Teb1 was robust without Teb2 and Teb3 and showed no requirement for the Teb1C CT $\alpha$ H (38). However, deletion of this  $\alpha$ -helix precluded assembly of Teb1 with other telomerase holoenzyme subunits *in vivo* (40). Because RRL expression and assembly of Teb1 sensitized high RAP activity reconstitution for stimulation by Teb2 and Teb3 (34), we used this system to test the significance of the TEB heterotrimer interface.

We generated expression constructs that removed the CT $\alpha$ H of Teb1, Teb2 OB-fold domain, or Teb3 (Fig. 5*B*). Recombinant telomerase complexes were assembled containing the RNP catalytic core, p50, and N-terminally F-tagged Teb1, Teb2, or Teb3 in the presence or absence of the other TEB subunits. Complexes containing an F-tagged protein were enriched by binding to anti-FLAG-agarose and then assayed for co-purified telom-

**TABLE 2****Dissociation constants for oligonucleotide binding to recombinant proteins and complexes**

The column labeled "Protein" indicates the recombinant protein or protein complex used for assays of binding to the column header DNA oligonucleotides. Numbers for Teb1, TEB, Rpa1, RPA, Rlp1, and RTT are given in nM as calculated from three experimental replicates of the gel mobility shift assays. S.E. was calculated for each mean to give an estimate of the variation among the replicates. All values had  $p < 0.06$  for goodness of fit for the one-site binding model used to calculate the dissociation constant. —, binding affinity too low to quantify using the gel mobility shift assay conditions. Rows labeled "-Fold change" indicate the relative increase in binding comparing the preceding large subunit protein alone to the heterotrimer.

Protein	(GT <sub>2</sub> G <sub>3</sub> ) <sub>3</sub>	dT <sub>18</sub>	(GT <sub>2</sub> G <sub>3</sub> ) <sub>5</sub>	dT <sub>30</sub>
Teb2-Teb3	—	—	—	—
Teb1 (nM)	3.4 ± 1.3	—	2.5 ± 1.0	—
TEB (nM)	2.9 ± 0.8	—	2.2 ± 1.7	—
-Fold change	1.2	—	1.1	—
Rpa1 (nM)	36 ± 11	41 ± 20	36 ± 15	39 ± 16
RPA (nM)	24 ± 9	30 ± 12	7.4 ± 3.0	5.3 ± 0.6
-Fold change	1.5	1.4	4.8	7.3
Rlp1 (nM)	68 ± 22	75 ± 9	73 ± 28	81 ± 14
RTT (nM)	45 ± 16	56 ± 25	27 ± 13	21 ± 7
-Fold change	1.5	1.3	2.7	3.8



**FIGURE 4. Comparison of telomerase activity stimulation by Teb2-Teb3, Teb1, TEB, Rpa1, RPA, Rlp1, or RTT.** The same amount of RRL-reconstituted telomerase complex (composed of the RNP catalytic core, p50, p75, p45-F, and p19) was assayed in the presence or absence of the indicated recombinant proteins at a concentration of 40 or 200 nM. Teb2-Teb3 was added at 200 nM.

erase catalytic activity (Fig. 5C). Each F-tagged TEB subunit co-purified a similar amount of high RAP activity in the presence of all three TEB subunits (Fig. 5C, lanes 2, 5, and 7), indicating that the tagged proteins assembled into telomerase holoenzyme. F-Teb1 recovered a low amount of high RAP activity in the absence of Teb2 and Teb3 (Fig. 5C, lanes 1 and 2). F-Teb1 $\Delta$ CT $\alpha$ H co-purified the same level of activity co-purified by Teb1 without Teb2 and Teb3 even if Teb2 and Teb3 were present (Fig. 5C, lanes 3 and 4). In comparison, purification through F-Teb2 $\Delta$ CT $\alpha$ H or F-Teb3 $\Delta$ CT $\alpha$ H failed to recover any telomerase activity, even in the presence of other TEB subunits (Fig. 5C, lanes 6 and 8). These findings establish

that assembly of Teb2-Teb3 into telomerase holoenzyme requires the formation of the heterotrimer  $\alpha$ -helix bundle. Deletion of any one of the three  $\alpha$ -helices precluded Teb2-Teb3 association with or stimulation of high RAP activity.

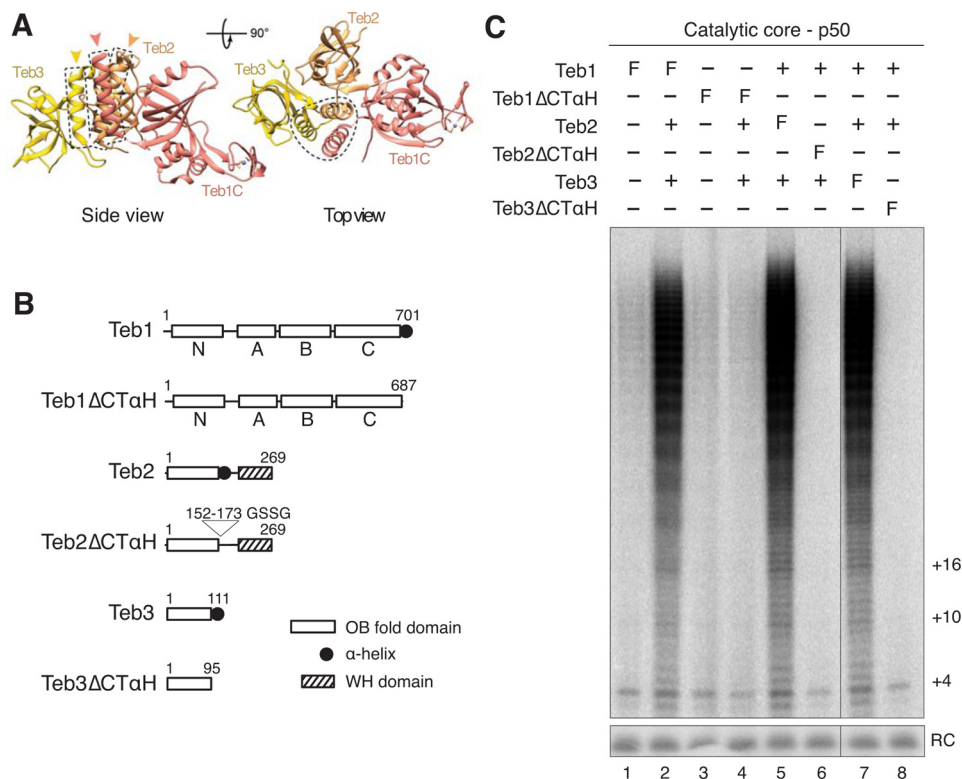
Previous assays were not designed to detect a subtle influence of Teb2-Teb3 on RAP or the rate of repeat synthesis. Therefore, we profiled repeat synthesis over a time course of 40 min for telomerase enzymes reconstituted with Teb1 or TEB. We purified Teb1- or TEB-containing telomerase complexes from reconstitutions with the p50-bound RNP catalytic core using F-Teb1. In comparison, we purified the p50-bound RNP catalytic core reconstituted without TEB subunits using p50-F. Activity assays confirmed that both Teb1 and TEB dramatically increase the rate of tandem repeat synthesis and the amount of high RAP product relative to the p50-bound RNP catalytic core alone (Fig. 6). There was no major difference in the profile of high RAP repeat synthesis with Teb1 alone *versus* TEB. However, F-Teb1 co-purified much less activity in the absence than in the presence of Teb2-Teb3 (Fig. 6, lanes 6–10 *versus* lanes 11–15 from the same gel, but the *top panel* of lanes 6–10 is shown with amplified signal intensity relative to lanes 1–5 and 11–15). With equal inputs of F-Teb1 and p50-bound RNP catalytic core, the simplest explanation for the results is that Teb2-Teb3 increased the amount of Teb1 associated with active telomerase. Background binding of RRL-expressed telomerase subunits to purification resin makes direct quantification of this influence unreliable (data not shown). In addition, some activity of p50-bound RNP catalytic core alone was detectable in assays of enzyme purified by F-Teb1 without Teb2-Teb3, but not enzyme with TEB (Fig. 6, compare low RAP with high RAP activity in lanes 6–10 and 11–15). We suggest that some p50-bound RNP catalytic core dissociated from Teb1, but not TEB, during the activity assay reaction.

Altogether, results of this work implicate Teb2 and Teb3 as assembly factors for Teb1 incorporation into telomerase holoenzyme. The importance of this Teb2-Teb3 activity *in vivo* is supported by the inability of Teb1 $\Delta$ CT $\alpha$ H to interact with active telomerase in cells (40).

## Discussion

In this work, we characterized the proteins Teb2 and Teb3 previously isolated as subunits of the *Tetrahymena* telomerase holoenzyme (34). Although the other *Tetrahymena* telomerase holoenzyme subunits are telomerase-specific (37), we show here that Teb2 and Teb3 are not (Fig. 7A). Teb2 and Teb3 have all of the properties expected for RPA middle and small subunits, respectively. Phylogenetic alignments clustered Teb2 with Rpa2 and Teb3 with Rpa3, and their mRNA abundance indicated an expression level higher than other telomerase proteins (34). More definitively, in this study, we demonstrate that in cells, Teb2 and Teb3 are bound to *Tetrahymena* Rpa1 as the majority fraction of their cellular pool. In addition to this physical interaction evidence, DNA binding assays indicate that Teb2 and Teb3 increase the DNA binding affinity of heterotrimeric Rpa1-Teb2-Teb3 relative to Rpa1 alone. Specifically, Teb2 and Teb3 improved Rpa1 interaction with 30-nt ssDNAs but not 18-nt ssDNAs, regardless of ssDNA sequence. In summary, our purification and reconstitution approaches strongly

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**FIGURE 5. Roles of the TEB subunit OB-fold domain C-terminal  $\alpha$ -helices.** *A*, model of Teb1C-Teb2N-Teb3 based on fitting of the human RPA heterotrimer (RPA70C-RPA32N-RPA14) (Protein Data Bank entry 1L10) into a cryo-electron microscopy map, followed by replacement of RPA70C with Teb1C, except for the RPA70C C-terminal  $\alpha$ -helix (34). Human Rpa2 OB-fold domain and Rpa3 fit the density of Teb2 OB-fold domain and Teb3, respectively. The three-helix bundle is highlighted with a *dashed line*. *B*, diagram of proteins without or with OB-fold CT $\alpha$ H removal. *Numbers*, amino acids in the full-length proteins; GSSG, a glycine/serine linker added to replace the CT $\alpha$ H of the Teb2 OB-fold domain. *C*, telomerase activity reconstituted using the proteins diagrammed in *B* after purification using anti-FLAG antibody resin.

support a complex of Rpa1, Teb2, and Teb3 as the general ssDNA-binding *Tetrahymena* RPA (Fig. 7*A*, *middle*).

Telomerase “appropriation” of Teb2 and Teb3 from RPA raises the question of what the subunits of a general ssDNA-binding factor contribute to telomeric repeat synthesis. The Teb2 OB-fold domain is displaced from the RPA-like configuration on DNA by Teb1C interaction with the telomerase RNP catalytic core (Fig. 7*A*, *top*). Also, DNA binding by TEB-assembled Teb2 would be unnecessary given the high affinity of Teb1 alone for telomeric ssDNA. We suggest that Teb2 and Teb3 influence Teb1 function indirectly by favoring a Teb1 conformation productive for holoenzyme assembly. Teb2-Teb3 function depends on the Teb1C CT $\alpha$ H, because Teb1 $\Delta$ CT $\alpha$ H stimulation of high RAP activity lost its enhancement by Teb2-Teb3. Teb1 $\Delta$ CT $\alpha$ H expressed in *Tetrahymena* does not co-purify telomerase activity from cell extracts (40), yet bacterially expressed Teb1 $\Delta$ CT $\alpha$ H can reconstitute high RAP telomerase activity *in vitro* (38). Combined, these findings support the conclusions that the telomerase holoenzyme role of Teb2-Teb3 is indirect through Teb1, is critical under biological conditions, and is better recapitulated by TEB expressed and assembled with active RNP in RRL than by bacterially expressed and purified TEB. Our findings also reveal that the same proteins make different functional contributions in an RPA *versus* TEB heterotrimer.

The original annotation of the open reading frame for *Tetrahymena* Rpa1 suggested an  $\sim$ 70-kDa protein, which did not

match the mobility of a major ZZP-Teb2- or ZZP-Teb3-associated protein. We initially suspected that a post-translation modification of endogenous Rpa1 altered its SDS-PAGE migration. However, upon further investigation, GenBank<sup>TM</sup>-deposited *Tetrahymena* mRNA expressed sequence tags support the possibility of an extended N terminus. The longest Rpa1 mRNA would encode an  $\sim$ 80-kDa protein and from an alternative start site an  $\sim$ 60-kDa protein (Fig. 7*B*), which are the sizes of the predominant polypeptides co-purified with Teb2 and Teb3 (Fig. 2*C*). The  $\sim$ 60-kDa Rpa1 protein could also result from proteolysis in the linker between the regulatory N-terminal OB-fold domain and the DNA-binding OB-fold domains. MS analysis here validated expression of the longest open reading frame, because sequenced peptides mapped within its unique N-terminal region (Fig. 7*B*). Because C-terminal tagging of *Tetrahymena* Rpa1 inactivated its biological function (38), a different approach will be required to test whether a start codon other than that for the  $\sim$ 80-kDa protein is also used for Rpa1 translation in cells. Our biochemical characterizations used Rpa1 expressed from a synthetic gene encoding the originally annotated sequence (38), which is truncated for the N-terminal region of the N-terminal OB-fold domain. For *in vitro* assays the  $\sim$ 70-kDa protein should be representative, because the Rpa1 N-terminal OB-fold does not influence DNA binding affinity or assembly of Rpa1 into heterotrimer (23, 24).

Beyond the general ssDNA-binding *Tetrahymena* RPA composed of Rpa1, Teb2, and Teb3, our findings suggest that cells



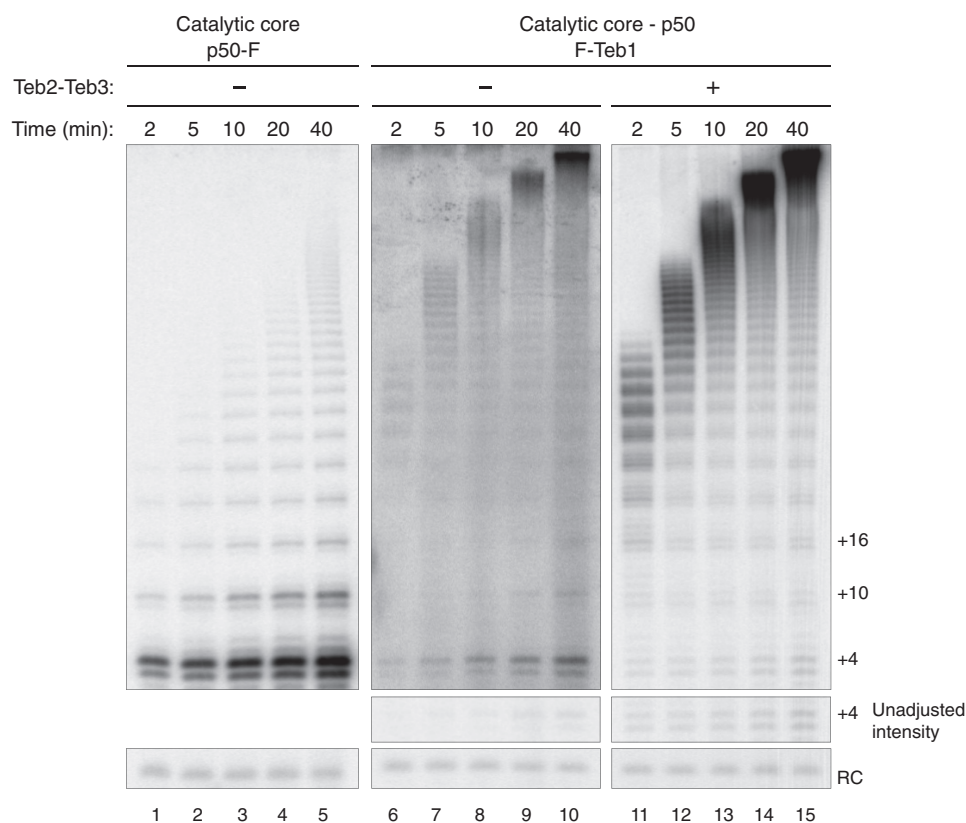


FIGURE 6. **Comparison of telomerase assembly and activity with Teb1 or TEB.** A time course of product synthesis was monitored for enzyme without Teb1 (lanes 1–5), with Teb1 alone (lanes 6–10), or with TEB (lanes 11–15). All lanes are from the same gel, but the *top panel* of the *middle set* of lanes is shown at amplified signal intensity relative to the *flanking panels*.

undergoing sexual reproduction assemble an additional RPA-related complex composed of Rlp1, Rlp2, and Teb3. Also, it remains possible that Rlp1 assembles with Teb2 and Teb3 in cells under conditions not studied in this work. Based on the sequence-nonspecific DNA binding specificity of Rlp1, any heterotrimer with Rlp1 is likely to have general ssDNA binding activity. Rlp1 lacks an N-terminal regulatory OB-fold domain, as do some predicted Rpa1 paralogs in protozoan parasites (50). Rpa1, Teb1, Teb2, and Teb3 mRNAs are detectable in cells in vegetative growth, starvation, and sexual reproduction (Fig. 7C, *top* and *middle*; Teb2 abundance could be analyzed only by expressed sequence tags, not shown). In contrast, Rlp1 and Rlp2 mRNAs are expressed at extremely low, if any, level in vegetative growth (Fig. 7C, *bottom*). Consistent with this expression pattern, *RLP1* knock-out was not deleterious for vegetative growth (38). The expression specificity of Rlp1 and Rlp2 suggests that they and Teb3 could function together specifically in mated cells. The complexity of RPA and RPA-like complexes in *Tetrahymena* provides new insights and opportunities to understand the function of alternative RPA subunits, which, based on genome sequencing, appear widespread across organisms ranging from apicomplexan parasites to plants to mammals (41–43).

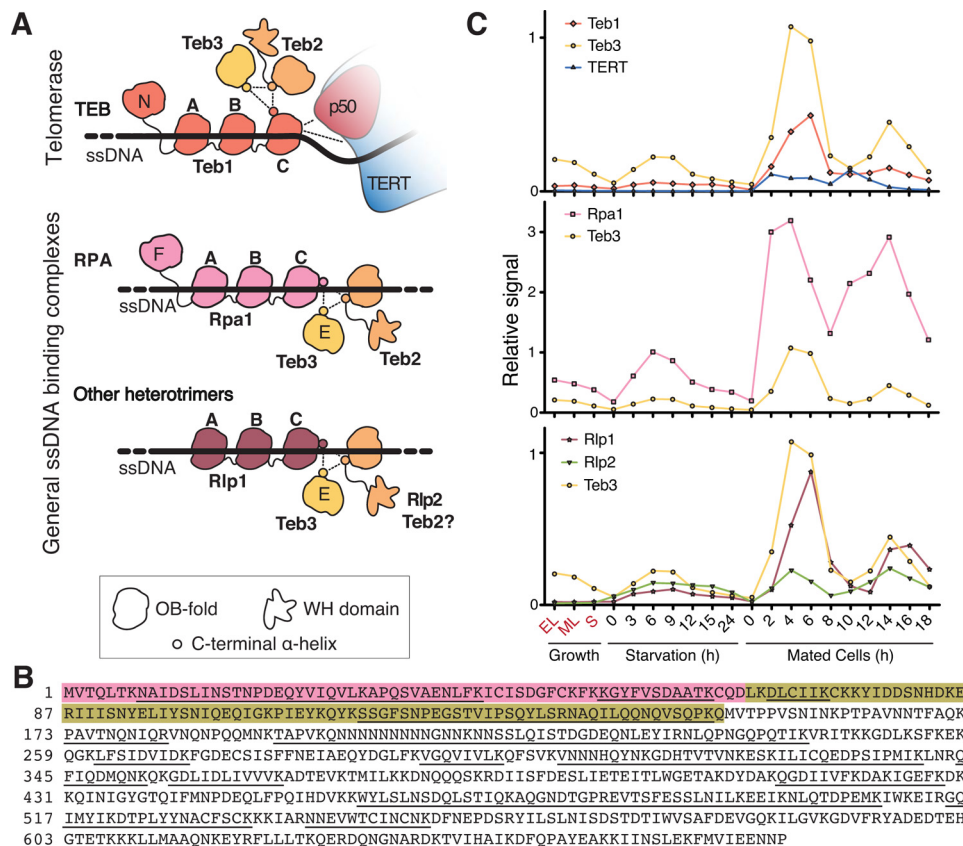
## Experimental Procedures

**Tetrahymena Strain Construction and Growth**—*Tetrahymena* strains expressing tagged Teb2 or tagged Teb3 instead of the endogenous untagged protein were generated by cassette

integration at the respective genomic loci using the BSR2 cassette (44). N-terminal tag fusion was chosen due to predicted protein domain structures and the loss of function imposed by fusion of the same tag to the C terminus of *Tetrahymena* Rpa1 (38). Cells were grown in modified Neff medium (0.25% proteose peptone, 0.25% yeast extract, 0.2% dextrose, 30  $\mu$ M FeCl<sub>3</sub>) to mid-log phase ( $3 \times 10^5$  cells/ml). For mating, cells were starved in 10 mM Tris (pH 8.0) for 16 h and mixed in a 1:1 ratio with complementary mating type SB1969 at  $2 \times 10^5$  cells/ml. To maximally synchronize mating, cells were shaken at 180 rpm for 30 min followed by a 30-min rest period three times. Conjugating cells were harvested 5 h after the final shake period. For cells synchronized in vegetative growth, starved cell cultures described above were re-fed with modified Neff medium at  $3 \times 10^5$  cells/ml for 4 h.

**Affinity Purification and Mass Spectrometry**—Cell lysis (completed at 4 °C) and subsequent steps (completed at room temperature) used T2EG50 (20 mM Tris, pH 8.0, 2 mM EGTA, 10% glycerol, 50 mM NaCl) supplemented with 0.2% Igepal CA-630 and 2 mM dithiothreitol (DTT). After binding to IgG-agarose (Sigma) and washing in T2EG50 with 0.03% Igepal CA-630 and 2 mM DTT, bound complexes were incubated with 30  $\mu$ g/ml tobacco etch virus protease for 0.5–1 h. Eluted samples were bound in batch to 10  $\mu$ l of EZView Red anti-FLAG M2 resin (Sigma) per 50-ml initial extract volume in low retention tubes for 1 h. Washed resin was eluted into T2MG (20 mM Tris, pH 8.0, 2 mM MgCl<sub>2</sub>, 10% glycerol) with 1 mM DTT and 150 ng/ $\mu$ l

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**FIGURE 7. RPA and RPA-like complexes in *Tetrahymena*.** *A*, illustration of the inferred DNA contacts within TEB, RPA, and Rlp1-containing heterotrimer(s). Teb2–Teb3 interaction with Teb1 stabilizes the interaction of Teb1C with p50 and the RNP catalytic core. Teb2–Teb3 interaction with Rpa1 or Rlp1 instead extends the DNA-binding surface of the heterotrimer compared with the large subunit alone. *B*, *Tetrahymena* Rpa1 coding region. *Pink*, extra sequence in ~80-kDa versus ~70-kDa predicted proteins; *yellow*, extra sequence in ~70-kDa versus ~60-kDa predicted proteins. The original ~70-kDa predicted protein had three additional N-terminal amino acids not shown. Peptide sequences identified by MS are *underlined*. We note that the protein open reading frame is validated by peptide sequence to the C terminus that we annotated, instead of the C terminus from an alternative reading frame predicted in the most recent genome database update. *C*, microarray mRNA expression profiles for Teb3 (*yellow*) compared with other telomerase subunits (*top*), other RPA subunits (*middle*), and RPA-like protein subunits (*bottom*). There are no microarray expression data for Teb2. Graphs were generated using information from the *Tetrahymena* Gene Expression Database (52, 53). *Vertical axes*, relative microarray signal intensity. *Horizontal axis*, a progression of life cycle states: in vegetative growth at low density (EL), log phase (ML), or stationary phase (S); in starvation medium for the indicated time in hours following transfer from growth media; mating time points in hours following mixing of two mating types.

of 3× FLAG peptide for 60 min. For mass spectrometry, samples were washed with T2EG50 supplemented with 2 mM DTT but no Igepal CA-630 before elution and dialyzed into the same buffer to remove the FLAG peptide.

MS peptide digests used sequencing grade trypsin (Promega). The IP2 program suite (Integrated Proteomics) was used for peptide and protein identification. The proLuCid search engine was used with the *Tetrahymena thermophila* predicted proteome database (ciliate.org) downloaded on June 27, 2014. The set of three samples was purified and processed for MS in parallel, with MS data shown from one biological replicate (of multiple replicates that had a similar SDS-PAGE protein profile).

**Recombinant Protein Expression and Purification**—Recombinant Teb1, Rpa1, Rlp1, Teb2, and Teb3 (34, 37, 38) were expressed in *E. coli* BL21 (DE3) using synthetic open reading frames, which is necessary due to expanded *Tetrahymena* codon usage. Each RPA large subunit paralog was expressed from pET28a with an N-terminal His<sub>6</sub> tag, as described previously (38), or in combination with pCDFDuet vector expressing Teb2 with an N-terminal MBP tag and untagged Teb3. Cells were lysed by sonication for 3 min at 4 °C in T2MG50 (20 mM Tris, pH 8.0, 2 mM MgCl<sub>2</sub>, 10% glycerol, 50 mM NaCl) supple-

mented with 0.1% Igepal CA-630, 1 mM DTT, and 20 mM imidazole. His<sub>6</sub>-tagged proteins were isolated by purification on Ni-NTA-agarose (Qiagen) and eluted into T2MG with 2 mM DTT and 500 mM imidazole. Complexes that also contained an MBP-tagged subunit were isolated by subsequent purification on amylose resin (New England Biolabs) and eluted into T2MG with 1 mM DTT and 10 mM maltose. The complex of MBP–Teb2 and Teb3 was purified using amylose resin only.

**Telomerase Activity and DNA Binding Assays**—Activity assays used a standard *Tetrahymena* telomerase reaction buffer containing 50 mM Tris acetate (pH 8.0), 2 mM MgCl<sub>2</sub>, 10 mM spermidine, and 5 mM  $\beta$ -mercaptoethanol. Reactions additionally contained 24 nM [ $\alpha$ -<sup>32</sup>P]dGTP, 300 nM dGTP, 200  $\mu$ M dTTP, and 200 nM DNA primer (GT<sub>2</sub>G<sub>3</sub>)<sub>3</sub>. Reactions were allowed to proceed at room temperature for 5 min for cell-assembled telomerase or 15 min for telomerase assembled in RRL, unless indicated otherwise. A 5′-labeled oligonucleotide DNA (the recovery control) was added to telomerase products before precipitation and denaturing PAGE. Electrophoretic mobility shift assays were performed as described previously (38). Binding affinities were calculated based on free probe signal using ImageQuant software.

**Telomerase Reconstitutions**—Recombinant telomerase was reconstituted by RRL expression of synthetic open reading frames for TERT, p50, p75, p45, p19, Teb1, Teb2, and/or Teb3 (34). Bacterially expressed, purified p65 (51) and *in vitro* transcribed TER were added at 25 nM each to the TERT RRL reaction before protein synthesis to assemble the RNP catalytic core. A separate RRL reaction was performed to produce the p75-p45-p19 CST complex or p50, which here was the p50N30 domain sufficient for p50 biological function (36). RRL-expressed telomerase subunits were combined, bound to anti-FLAG M2 affinity resin (Sigma), and washed into T2MG with 2 mM DTT. Afterward, purified proteins other than p65 were added to a final concentration of 200 nM (unless indicated otherwise) and allowed to bind for 20 min at room temperature, followed by an activity assay.

**Sequence Depositions**—GenBank™ accession numbers for the *Tetrahymena thermophila* protein sequences expressed recombinantly in this work are as follows: Rpa1, ADB03555.1; Rlp1, GU384877; Teb1 (encoded by gene *TAP82*), EU873081; Teb2, BK009378; Teb3, BK009379. Revised Rpa1 sequence has GenBank™ accession number ADB03555.2. Rlp2 sequence has GenBank™ accession number KX987301.

**Author Contributions**—H. E. U. and K. C. conceived and interpreted the experiments and wrote the paper. H. E. U. performed all experiments. H. C. and J. F. shared reagents and information from parallel studies. All authors revised and approved the final version of the manuscript.

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