How Structural Features Define Biogenesis and Function of Human Telomerase RNA Primary Transcript

Maria Rubtsova $^{1,2,*}$ and Olga Dontsova $^{1,2,3}$

$^1$ Department of Chemistry, A.N. Belozersky Institute of Physico Chemical Biology, Lomonosov Moscow State University, 119991 Moscow, Russia; olga.a.dontsova@gmail.com

$^2$ Shemyakin-Ovchinnikov Institute of Bioorganic Chemistry of the Russian Academy of Sciences, 117997 Moscow, Russia

$^3$ Center of Life Sciences, Skolkovo Institute of Science and Technology, Skolkovo, 121205 Moscow, Russia

* Correspondence: mprubtsova@gmail.com

Abstract: Telomerase RNA has been uncovered as a component of the telomerase enzyme, which acts as a reverse transcriptase and maintains the length of telomeres in proliferated eukaryotic cells. Telomerase RNA is considered to have major functions as a template for telomeric repeat synthesis and as a structural scaffold for telomerase. However, investigations of its biogenesis and turnover, as well as structural data, have provided evidence of functions of telomerase RNA that are not associated with telomerase activity. The primary transcript produced from the human telomerase RNA gene encodes for the hTERP protein, which presents regulatory functions related to autophagy, cellular proliferation, and metabolism. This review focuses on the specific features relating to the biogenesis and structure of human telomerase RNA that support the existence of an isoform suitable for functioning as an mRNA. We believe that further investigation into human telomerase RNA biogenesis mechanisms will provide more levels for manipulating cellular homeostasis, survival, and transformation mechanisms, and may contribute to a deeper understanding of the mechanisms of aging.

Keywords: telomerase RNA; RNA biogenesis; RNA processing; localization; alternative function; transport; hTERP

1. Introduction

Telomerase is a ribonucleoprotein complex that is essential for telomere maintenance [1,2]. Telomeres are special structures located at the ends of linear eukaryotic chromosomes which are shortened in each round of cellular division due to the end replication problem and nuclease action. Critically short telomeres appear after a particular number of cellular divisions (50–70), activating the cellular senescence mechanism, which results in death in a majority of cells; however, some cells may overcome this crisis, acquire the ability of unlimited proliferation associated with cancer phenotypes, and activate telomerase as a major mechanism to maintain telomere length. Telomerase is active in limited types of cells, and its functioning has been associated with the proliferative status of cells [3,4]. For example, embryonic, stem, germ, cancer cells, activated immune cells, cells participating in regenerative processes, and other cells with a high proliferation level present telomerase activity. The decreased capacity to activate telomerase during regeneration and the immune response are associated with the aging of organisms. In contrast, the activation of telomerase in cells with uncontrolled proliferation stimulates oncogenesis.

Telomerase contains two core components—telomerase reverse transcriptase (TERT) and the telomerase RNA component (TERC)—which are necessary for telomere synthesis [5]. To present telomerase activity, cells should express both of these core components of telomerase (i.e., TERT and TERC) [6–8]. hTERT expression is tightly regulated and, as a rule, the expression level of hTERT determines the level of telomerase activity. A majority
of somatic cells lose telomerase activity through the silencing of hTERT gene expression during embryogenesis [9,10]; however, the expression of the hTERT gene remains constant in the majority of somatic cells [6]. It is not silenced and may be increased to facilitate telomerase activity in proliferated cells. The non-canonical non-telomeric functions of both core components of human telomerase have recently been reviewed [11]. The hTERT protein coding potential in the synthesis of human telomerase RNA protein (hTERP) has recently been uncovered [12]. hTERP is important for cell survival under stress conditions [12] and has been shown to be involved in autophagy regulation [13]. Although the existence of hTERP has been demonstrated, the canonical tertiary structure of hTERC described for the telomerase complex is not convenient for ribosome binding and translation initiation. Unfortunately, hTERP properties and functions have not been deeply investigated at this time and we do not know the correlation of hTERP appearance with the expression of hTERT and hTERC. However, we hypothesize that hTERP is synthesized when hTERT is absent in cells. hTERP may be constitutively expressed, like hTERC, and the presence of hTERP may be important for alternative telomere maintenance mechanism realization. The expression profile of hTERP and its correlation with the proliferative status of cells as well as with metabolism should be investigated in future. In this review, we discuss the alternative structural features of transcripts derived from the hTERC gene, which must define whether telomerase RNA acts as a telomerase component or as mRNA encoded for hTERP.

2. Diverse Forms of Human Telomerase RNA Are Processed from the Nascent Transcript

Telomerase RNA is synthesized as a long precursor in Integrator-dependent mode [14], which is processed to obtain the mature RNA component of telomerase (451 nucleotides in length; see Figure 1A) [9]. Transcripts elongated up to 1000 nucleotides (Figure 1A) have been identified using next-generation sequencing (NGS) [15] and reverse transcription–polymerase chain reaction (RT-PCR) [16] techniques. Pathways of TERC processing (Figure 1) have also been reviewed in detail [17]. Oligoadenylated hTERC premature transcripts are trimmed by exosomes attracted through the interaction with adaptor proteins [15,18]. In the nucleoplasm, the nuclear exosome-targeting complex (NEXT) participates in the recruitment of exosomes to early unprocessed hTERC transcripts, resulting in degradation (Figure 1B). The shorter hTERC transcripts, containing several additional nucleotides at the 3′-end, are oligoadenylated by non-canonical poly(A)polymerase PAPD5 [19,20] and processed by poly(A)ribonuclease PARN [21,22], or by the TRAMP complex (Trf4/Air2/Mtr4p polyadenylation complex) [23] in the nucleolus or nucleoplasm, followed by transport to Cajal bodies [24]. Oligoadenylated hTERC precursors may be directly transported in Cajal bodies, where they are processed by PARN and TOE1 exonucleases [25] (Figure 1B). Finally, the matured hTERC is localized in Cajal bodies. The depletion of PABPN1 (poly(A)-binding protein nuclear 1) results in a decreased level of total hTERC, which is rescued when NEXT exosome components are co-depleted [18,26]. The depletion of the PAXT (poly(A) tail exosome targeting) complex involved in the degradation of polyadenylated RNAs through interaction with PABPN1 results in the accumulation of the unprocessed 3′-extended form of hTERC [26,27]. Polyadenylation is necessary for processing hTERC to the mature component of the telomerase complex. The impairment of this stage leads to the accumulation of unprocessed transcripts in the cytoplasm, where translation resulting in the synthesis of hTERP occurs (Figure 1B). It has been proposed that the gradually lengthened poly(A) tail acts as a sensor, and PABPN1 is preferentially recruited to hTERC precursors with longer poly(A) tails in order to prioritize their maturation [17]. However, PABPN1 can be substituted with PABP (poly(A)-binding protein) [26] to direct the export of long hTERC polyadenylated transcripts into the cytoplasm for translation. hTERC cytoplasmic localization in normal conditions has recently been determined [12,16]. Initially, the cytoplasmic localization of hTERC was observed when dyskerin or PARN were depleted and accumulation or defects in processing occurred. The co-depletion of exosome subunit RRP6 and cytoplasm decapping factor DCP2 restores the nuclear localization of
hTERC under the condition of decreased levels of dyskerin or PARN [22]. The enzyme responsible for the polyadenylation of hTERC localized in cytoplasm has not yet been identified. Interestingly, the depletion of canonical PAPs decreases the hTERC level, providing an argument for the mRNA properties of hTERC (Figure 1B).

Co-transcriptional monomethylguanosine (MMG) modification occurs at the 5'-ends of transcripts synthesized by RNA-polymerase II. The MMG cap on precursor snRNAs recruits the cap-binding complex (CBC), the PHAX adaptor protein, and CRM1 for transport to the cytoplasm [28]. After hypermethylation by the trimethylguanosine synthase 1 (TGS1) of the MMG cap, snRNAs are reimported to the nucleus. Conversely, PHAX directs MMG-capped C/D-box snoRNAs to Cajal bodies where TGS1 hypermethylates the cap. The obtained trimethylguanosine (TMG)-capped C/D-box snoRNA recruits CRM1 in a CBC-independent manner, which is necessary for nucleolar localization [28]. The modification of the cap and the nuclear transport and export of hTERC is less well-investigated; however, it is known that TGS1 is responsible for the hypermethylation of MMG-capped hTERC (Figure 1B) and that the depletion of TGS1 leads to the cytoplasmic accumulation of hTERC (Figure 1B), where it is not associated with hTERT. Notably, the total level of hTERC transcript was increased in TGS1-depleted cells, without the redistribution of portions of 3'-end-extended and oligoadenylated forms of hTERC [16].

**Figure 1.** Biogenesis pathways of human telomerase RNA transcripts (hTERC): (A) two isoforms of hTERC presented in cells (https://genome.ucsc.edu accessed on 9 June 2022); (B) scheme illustrating biogenesis mechanisms of products transcribed from hTERC gene. Integrator (Int), DSIF, NEFL, and cap-binding complex (CBC) assist RNA polymerase II to synthesize the primary hTERC transcript [14,27]. ARS2 substitutes NELF and coordinates the distinct, mutually exclusive complex assembly to process the primary transcript [29–32]. During processing, monomethylated (MMG), polyadenylated hTERP mRNA is protected by PABPN1, which is substituted by PABP and exported to the cytoplasm, where it is translated [18,27]. Oligoadenylated hTERC transcripts may be degraded by RNA exosomes attracted by the PAXT complex [27], or may be trimmed by PARN followed by degradation through NEXT-attracted RNA-exosomes [21], or transported by PHAX to the nucleolus [28] where it is hypermethylated (TMG) by TGS1 [16] and matured by TRAMP-mediated RNA exosome trimming [15,18] for assembly into the active telomerase complex.
It has recently been demonstrated [27] that PHAX deficiency does not influence the level of mature hTERC. The co-depletion of the PAXT complex decreased the amount of mature and 3′-extended hTERC transcripts, and the co-depletion of TRAMP stimulated the accumulation of 3′-extended hTR transcripts, especially in the cytoplasmic fraction. We hypothesize that, in the absence of both the PHAX transport factor and TRAMP complex, the 3′-extended hTERC transcripts accumulate and are redirected to an alternative transport pathway which is inherent to mRNA.

Taking the details of processing and the localization of hTERC together, we can conclude that the hTERC gene produces two distinct types of molecules: the TMG-capped RNA component (451 nucleotides in length) of telomerase, which localizes in the nucleus and facilitates telomere synthesis in complex with hTERT in actively proliferated cells, and MMG-capped 3′-end-extended mRNA localized in the cytoplasm, which directs the synthesis of hTERP under unidentified conditions (Figure 1).

3. Structural Features of hTERC Responsible for Telomerase Activity

Considering the evidence that distinct forms of hTERC are present in cells, the structure of the mature telomerase component has been investigated in detail, where the object of investigation was TERC as a component of the ribonucleoprotein complex that provides the structural frame for telomerase. The telomerase complex structure has recently been reviewed in detail [33]. hTERC contains several structural domains essential for telomerase activity. The pseudoknot and H/ACA domains may reconstitute telomerase activity, even if they are expressed separately [34,35] (Figure 2A). Pseudoknot is important for binding hTERT and contains regions which possess a template for telomere synthesis (Figure 2B). The H/ACA domain, along with the NOP10, NHP2, dyskerin, and GAR1 proteins, forms the H/ACA complex [36], which blocks excessive degradation by exosomes and PARN, which trims the nascent transcript [18,21] (Figure 1A). The formation of a triple helix at the 3′-end of hTERC prevents its excessive degradation prior to binding the H/ACA complex [37]. The CAB-box of the H/ACA domain contains the signal for localization in Cajal bodies, as assisted by the TCAB1 protein [38] (Figure 2B). The association of the TCAB1 protein with the CAB-box promotes the formation of a tertiary structure in the CR4/5 region of hTERC, which is preferable for association with hTERT [39]. TCAB1 dysfunction prevents telomerase complex formation, leading to a decrease in telomerase activity and the shortening of telomeres in human embryonic stem cells [40]. Conservative regions 4/5 (CR4/5), consisting of P5, P6, and P6.1, are responsible for association with hTERT, and the P6.1 stem is cooperatively shaped by TERT and histone H2A-H2B, the presence of which has recently been revealed in the active telomerase complex [41] (see Figure 2).

Figure 2. Schematic presentation of human telomerase structure: (A) secondary structure of mature human telomerase RNA hTERC; and (B) schematic view of human telomerase holoenzyme complex.
4. Human Telomerase RNA Structural Elements Form during Telomerase Complex Assembly

Structural investigations have mainly focused on the telomerase complex and telomerase RNA in the context of telomerase. Several investigations have been performed in vivo where the TERC structure was chemically probed in cellular lysates [42,43]. Much effort has been put into the investigation of the pseudoknot domain of hTERC by NMR [44,45]. Finally, cryo-EM structures of human telomerase and its complex with substrate have recently been reported [36,41]. Significant information has also been obtained from biochemical investigations of telomerase activity using mutant variants of the enzyme component.

The search for genes which, through their dysfunction, are responsible for diseases characterized by short telomeres and senescence phenotypes—known as telomeropathies—has revealed various telomerase components and components involved in telomerase biogenesis. Such mutations predominantly affect the steady-state accumulation of hTERC [46]. Dyskerin mutations lead to the accumulation of hTERC in the cytoplasm or excessive degradation by exosomes during biogenesis (Figure 1); however, mutations in the hTERC sequence may affect telomerase activity without presenting an apparent impact on hTERC steady-state accumulation. The dysfunction of TCAB1 or mutations in the CR4/5 domain of hTERC disturb TCAB1-dependent tertiary structure formation, blocking the interaction with hTERT but not influencing the stability of hTERC [39] (see Figure 2B). Mutations in hTERC associated with the dyskeratosis congenita (DKC) disease include a shift in the equilibrium toward the hairpin structure due to pseudoknot destabilization (Figure 3A) [44]. Critical evaluations of the disease-relevant impact of hTERC variants not compromised for accumulation in vivo have been performed in several works [47,48], which revealed that disturbances to almost all stems presented in the hTERC structure impair telomerase activity. Substitutions that replace a canonical base pair with a wobble pair have a less severe effect than the most disruptive substitutions. Each mutation affecting the structure of the P2b and P3 stems of the pseudoknot domain (Figure 2A) caused the dramatic inhibition of telomerase activity; however, mutations in other regions of the pseudoknot domains did not affect telomerase activity significantly. Mutations in the CR4/5 domain, which is responsible for hTERT binding, also affected telomerase activity. The most dramatic influence was observed when mutations disrupted the structure of the P6.1 stem (Figure 2A) [47,48].

The telomerase pseudoknot domain may form a phylogenetically conserved hairpin, which is stabilized by a unique uridine helix (Figure 3A). A functionally important interconversion between the hairpin and pseudoknot conformations has been proposed [44]. The existence of the hairpin, formed as an alternative structure to the pseudoknot, was initially determined by NMR [44,49]. Several years later, functional mutational analysis of the pseudoknot structure in human telomerase RNA confirmed the importance of the pseudoknot structure for telomerase activity, both in vitro and in vivo [50]. Indeed, it has been demonstrated that mutations which abolish the P3 helix have an inhibitory effect on telomerase activity, which was reversed through compensatory mutations. It was concluded that the alternative hairpin structure does not maintain telomerase function, as destabilization of the intraloop J2b/3 base pairing did not affect the enzyme activity. Unfortunately, the effect of stabilization of J2b/3 intraloop, pairing on telomerase activity was not investigated [50]. The existence of alternative structures of the pseudoknot domain of telomerase RNA in the ciliate Tetrahymena and yeast Saccharomyces cerevisiae strengthen the hypothesis about its evolutionary origin. An idea regarding the regulatory role of the P3 stem in active telomerase complex formation has recently been proposed and checked [51], and the data obtained suggest that the nascent hTERC transcript initially folds as an alternative hairpin, which is then remodeled into the more stable pseudoknot structure with time (Figure 3A). hTERT interaction could promote the refolding of hTERC. Interestingly, in this model of the alternative hairpin, the template region of hTERC is included in the stem stabilizing this structure (Figure 3A). The authors hypothesized that such a structure protects the single-stranded template region until the complex with hTERT is formed. Moreover, it was suggested that the order of the pseudoknot folding and interaction with hTERT is
important for proper complex formation. According to this model, hTERT should bind the CR4/5 domain first, then promote the pseudoknot folding associated with correct template accommodation in the catalytic center of hTERT to produce the active enzyme; meanwhile, the interaction of hTERT with the pre-folded pseudoknot leads to incorrect template positioning in the enzyme active center, producing a non-functional telomerase complex.

Mutations disrupting the structure of the P6.1 stem of the CR4/5 domain of hTERC result in telomerase dysfunction and diseases related to telomeropathies. The P6.1 stem is an evolutionarily conserved structural feature of telomerase RNA that forms a three-way junction fold. Careful analysis of the structure of this region, performed using different techniques such as chemical probing, NMR, cryo-EM, and FRET, has demonstrated its structural heterogeneity in free hTERC and hTERC bound with hTERT [39,42,52–54] (see Figure 3B). The association of hTERT with the CR4/5 domain depends on the structure formed by the P6.1 stem, and mutations which destabilize its three-way junction structure prevent hTERT binding.

Taken together, these structural data and mutational analysis may allow for the proposal of a model of telomerase complex association (Figure 3C). The native transcript is presented in several steady-state conformations [44,50,51,55]. The H/ACA-domain should be folded first, which occurs co-transcriptionally (Figure 1). The binding of H/ACA proteins protects hTERC from excessive exosome-driven 3′-end processing; however, the folding of hTERC requires specific interactions. TCAB1 binding is necessary for the P6.1 stem folding responsible for the association of the CR4/5 domain with hTERT [39]. Finally, the pseudoknot structure is formed after hTERT binding, which facilitates the positioning of the template region at the active center of the enzyme [42] (Figure 3C).

**Figure 3.** Biogenesis-driven human telomerase RNA structure formation: (A) alternative structures of pseudoknot domain of hTERC; (B) alternative structures of CR4/5 domain of hTERC; and (C) scheme illustrating hTERC structure formation during telomerase complex assembly. The primary synthesized hTERC transcript may be folded into two alternative structures [45,49]. Binding of TCAB1 facilitates folding of the P6.1 stem and CR4/Cr5 domain and Cajal body localization [39], where TERT binding promotes pseudoknot formation associated with proper template region positioning with respect to the enzyme active center [42].
5. hTERC Structure and Localization Determine Its Function

Intuitively, it is expected that the mutations in hTERC—especially in domains involved in its association with hTERT—will affect telomerase activity. Many efforts have been undertaken to determine the intracellular localization of hTERC; however, the data obtained in different works show no unity, and the question regarding the natural localization of hTERC (and, moreover, that of its different isoforms) remains open. The localization of hTERC has been explored mainly in cancer cells and under conditions of the overexpression of mature hTERC capable of forming the complex with hTERT. It has been demonstrated that hTERC localizes in Cajal bodies only in the presence of hTERT [56], where the active telomerase complex assembles and interacts with telomeres. Telomerase RNA has been found to be localized in the nucleoplasm, nucleolus, Cajal bodies, cytoplasm, and mitochondria [12,16,22,56–58]. The functionality of the non-nuclear localizations of telomerase RNA is not yet clear. Nucleolar localization is necessary for hTERC processing or turnover, and it is thought that Cajal body localization promotes telomerase complex assembly and association with telomeres. However, it has been demonstrated that telomerase yields proper telomere synthesis even in cells with disrupted Cajal bodies [24].

Evidence for hTERC function outside of the telomerase complex has recently been reviewed [11]. It has been found that the mutant form of hTERC which is unable to form telomerase enzyme protects cells under an apoptosis-inducing treatment [59]. Analysis of the hTERC sequence has revealed that it contains the open reading frame that encodes for the protein named hTERP [12]. The existence of hTERP has been demonstrated clearly, and it has been shown that it protects cells under stress conditions and is involved in the regulation of autophagy and proliferation signaling pathways [13]. However, the position of the start AUG codon in the tertiary structure of hTERC is not convenient for ribosome binding and efficient translation (Figure 1). Data confirming the existence of multiple forms of native hTERC transcripts and alternative folding of the naked hTERC transcript provide additional arguments in favor of hTERC protein coding capacity. We hypothesize that the alternative structure of hTERC, distinct from the pseudoknot, may provide ribosome scanning and translation initiation abilities; the existence of alternative structures of hTERC are necessary to facilitate hTERP synthesis when hTERT is absent, and cells do not require the synthesis of telomere repeats.

We hypothesize that auxiliary factors regulate the biogenesis, structure formation, and function of native hTERC transcripts. The binding of additional RNA chaperones should stimulate specific structure stabilization and transcript localization. We speculate that the structure of hTERC should be subject to regulation by some yet-undetermined factors or intracellular conditions (e.g., pH, ionic strength) in order to determine the fate of particular transcripts—either to become part of the telomerase complex and function in the nucleus, to be exported to the cytoplasm to meet the ribosomes, to be transported to mitochondria, or to be degraded.

6. Conclusions

Taken together, the available data concerning the assembly of the telomerase complex and the multiple mechanisms involved in TERC processing suggest that the function of TERC transcripts depends on their structure, bound proteins, and the exact intracellular localization of the final product of biogenesis. Novel data have uncovered new forms of hTERC transcripts and other non-telomerase functions of hTERC, opening perspectives for further investigations of the role of telomerase RNA transcripts in cellular functioning, survival, metabolism switching, and other aspects of cellular homeostasis.

Author Contributions: Conceptualization, M.R. and O.D.; writing—original draft preparation, M.R.; writing—review and editing, O.D.; funding acquisition, O.D. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by Russian Science Foundation, grant number 21-64-00006.

Institutional Review Board Statement: Not applicable.
Informed Consent Statement: Not applicable.

Data Availability Statement: Not applicable.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Greider, C.W.; Blackburn, E.H. Identification of a Specific Telomere Terminal Transferase Activity in Tetrahymena Extracts. Cell 1985, 43, 405–413. [CrossRef]
2. Morin, G.B. The Human Telomere Terminal Transferase Enzyme Is a Ribonucleoprotein That Synthesizes TTAGGG Repeats. Cell 1989, 59, 521–529. [CrossRef]
3. Holt, S.E.; Shay, J.W. Role of Telomerase in Cellular Proliferation and Cancer. J. Cell. Physiol. 1999, 180, 10–18. [CrossRef]
4. Bodnar, A.G.; Ouellette, M.; Frolikis, M.; Holt, S.E.; Chiu, C.P.; Morin, G.B.; Harley, C.B.; Shay, J.W.; Lichtsteiner, S.; Wright, W.E. Extension of Life-Span by Introduction of Telomerase into Normal Human Cells. Science 1998, 279, 349–352. [CrossRef] [PubMed]
5. Lingner, J.; Cooper, J.; Chell, T. Telomerase and DNA End Replication: No Longer a Lagging Strand Problem? Science 1995, 269, 1533–1534. [CrossRef] [PubMed]
6. Yashima, K.; Maitra, A.; Rogers, B.B.; Timmons, C.F.; Rathi, A.; Pinar, H.; Wright, W.E.; Shay, J.W.; Gazdar, A.F. Expression of the RNA Component of Telomerase during Human Development and Differentiation. Cell Growth Differ. 1998, 9, 805–813.
7. Blasco, M.A.; Rizen, M.; Greider, C.W.; Hanahan, D. Differential Regulation of Telomerase Activity and Telomerase RNA during Multi-Stage Tumorigenesis. Nat. Genet. 1996, 12, 200–204. [CrossRef]
8. Cayuela, M.L.; Flores, J.M.; Blasco, M.A. The Telomerase RNA Component Terc Is Required for the Tumour-Promoting Effects of Tert Overexpression. EMBO Rep. 2005, 6, 268–274. [CrossRef]
9. Feng, J.; Funk, W.D.; Wang, S.S.; Weinrich, S.L.; Avilion, A.A.; Chiu, C.P.; Adams, R.R.; Chang, E.; Allsopp, R.C.; Yu, J. The RNA Component of Human Telomerase. Science 1995, 269, 1236–1241. [CrossRef]
10. Roake, C.M.; Artandi, S.E. Regulation of Human Telomerase in Homeostasis and Disease. Nat. Rev. Mol. Cell Biol. 2020, 21, 384–397. [CrossRef]
11. Rubtsova, M.; Dontsova, O. Human Telomerase RNA: Telomerase Component or More? Biomolecules 2020, 10, 873. [CrossRef]
12. Rubtsova, M.; Naraykin, Y.; Vasilkova, D.; Meerson, M.; Zvereva, M.; Prassolov, V.; Lazarev, V.; Manuvera, V.; Kovalchuk, S.; Anikanov, N.; et al. Protein Encoded in Human Telomerase RNA Is Involved in Cell Protective Pathways. Nucleic Acids Res. 2018, 46, 8966–8977. [CrossRef] [PubMed]
13. Shliapina, V.; Koriagina, M.; Vasilkova, D.; Govorun, V.; Dontsova, O.; Rubtsova, M. Human Telomerase RNA Protein Encoded by TERC Is Involved in Cell Protective Pathways. Front. Cell Dev. Biol. 2021, 9, 754611. [CrossRef] [PubMed]
14. Rubtsova, M.P.; Vasilkova, D.P.; Moshareva, M.A.; Malynko, A.N.; Meerson, M.B.; Zatsepin, T.S.; Naraykin, Y.V.; Beletsky, A.V.; Ravin, N.V.; Dontsova, O.A. Integrator Is a Key Component of Human Telomerase RNA Biogenesis. Sci. Rep. 2019, 9, 1701. [CrossRef] [PubMed]
15. Tseng, C.-K.; Wang, H.-F.; Burns, A.M.; Schroeder, M.R.; Gaspari, M.; Baumann, P. Human Telomerase RNA Processing and Quality Control. Cell Rep. 2015, 13, 2223–2243. [CrossRef] [PubMed]
16. Chen, L.; Roake, C.M.; Galati, A.; Bavasso, F.; Micheli, E.; Saggio, I.; Schoetfner, S.; Cacchione, S.; Gatti, M.; Artandi, S.E.; et al. Loss of Human TGS1 Hypermethylase Promotes Increased Telomerase RNA and Telomere Elongation. Cell Rep. 2020, 30, 1358–1372.e5. [CrossRef]
17. Qin, J.; Autexier, C. Regulation of Human Telomerase RNA Biogenesis and Localization. RNA Biol. 2021, 18, 305–315. [CrossRef]
18. Nguyen, D.; Grenier St-Sauveur, V.; Bergeron, D.; Dupuis-Sandoval, F.; Scott, M.S.; Bachand, F. A Polyadenylation-Dependent 3' End Maturation Pathway Is Required for the Synthesis of the Human Telomerase RNA. Cell Rep. 2015, 13, 2244–2257. [CrossRef]
19. Fok, W.C.; Shukla, S.; Vessoni, A.T.; Brenner, K.A.; Parker, R.; Sturgeon, C.M.; Batista, L.F.Z. Posttranscriptional Modulation of TERC by PAPD5 Inhibition Rescues Hematopoietic Development in Dyskeratosis Congenita. Blood 2019, 133, 1308–1312. [CrossRef]
20. Shukla, S.; Jeong, H.-C.; Sturgeon, C.M.; Parker, R.; Batista, L.F.Z. Chemical Inhibition of PAPD5/7 Rescues Telomerase Function and Hepatopoiesis in Dyskeratosis Congenita. Blood Adv. 2020, 4, 2717–2722. [CrossRef]
21. Moon, D.H.; Segal, D.H.; Boyraz, B.; Guinan, E.; Hofmann, I.; Cahan, P.; Tai, A.K.; Agarwal, S. Poly(A)-Specific Ribonuclease (PARN) Mediates 3'-End Maturation of the Telomerase RNA Component. Nat. Genet. 2015, 47, 1482–1488. [CrossRef] [PubMed]
22. Shukla, S.; Schmidt, J.C.; Goldfarb, K.C.; Cech, T.R.; Parker, R. Inhibition of Telomerase RNA Decay Enables Telomerase Deficiency Caused by Dyskerin or PARN Defects. Nat. Struct. Mol. Biol. 2016, 23, 286–292. [CrossRef] [PubMed]
23. Jia, H.; Wang, X.; Liu, F.; Guenther, U.-P.; Srinivasan, S.; Anderson, J.T.; Jankowsky, E. The RNA Helicase Mtr4p Modulates Polyadenylation in the TRAMP Complex. Cell 2011, 145, 890–901. [CrossRef] [PubMed]
24. Chen, Y.; Deng, Z.; Jiang, S.; Hu, Q.; Liu, H.; Songyang, Z.; Ma, W.; Chen, S.; Zhao, Y. Human Cells Lacking Coilin and Cajal Bodies Are Proficient in Telomerase Assembly, Trafficking and Telomere Maintenance. Nucleic Acids Res. 2015, 43, 385–395. [CrossRef]
25. Deng, T.; Huang, Y.; Weng, K.; Lin, S.; Li, Y.; Shi, G.; Chen, Y.; Huang, J.; Liu, D.; Ma, W.; et al. TOE1 Acts as a 3' Exonuclease for Telomerase RNA and Regulates Telomere Maintenance. Nucleic Acids Res. 2019, 47, 391–405. [CrossRef]
26. Meola, N.; Jensen, T.H. Targeting the Nuclear RNA Exosome: Poly (A) Binding Proteins Enter the Stage. RNA Biol. 2017, 14, 820–826. [CrossRef]
27. Pakhomova, T.; Moshareva, M.; Vasilkova, D.; Zatsepin, T.; Dontsova, O.; Rubtsova, M. Role of RNA Biogenesis Factors in the Processing and Transport of Human Telomerase RNA. Biomedicines 2022, 10, 1275. [CrossRef]
28. Boulou, S.; Verheggen, C.; Jady, B.E.; Girard, C.; Pescia, C.; Paul, C.; Ospina, J.K.; Kiss, T.; Matera, A.G.; Bordonné, R.; et al. PHAX and CRM1 Are Required Sequentially to Transport U3 SnRNA to Nucleoli. Mol. Cell 2004, 16, 777–787. [CrossRef]
29. O’Sullivan, C.; Howard, P.L. The Diverse Requirements of ARS2 in Nuclear Cap-Binding Complex-Dependent RNA Processing. RNA Dis. 2017, 4, e1376. [CrossRef]
30. Narita, T.; Yung, T.M.C.; Yamamoto, J.; Tsuboi, Y.; Tanabe, H.; Tanaka, K.; Yamaguchi, Y.; Handa, H. NELF Interacts with CBC and Participates in 3’ End Processing of Replication-Dependent Histone MRNAs. Mol. Cell 2007, 26, 349–365. [CrossRef]
31. Hallais, M.; Pontvianne, F.; Andersen, P.R.; Clerici, M.; Lener, D.; Benbahouche, N.E.H.; Gostan, T.; Vandermoere, F.; Robert, M.-C.; Cusack, S.; et al. CBC–ARS2 Stimulates 3’-End Maturation of Multiple RNA Families and Favors Cap-Proximal Processing. Nat. Struct. Mol. Biol. 2013, 20, 1358–1366. [CrossRef] [PubMed]
32. Giacometti, S.; Benbahouche, N.E.H.; Domanski, M.; Robert, M.-C.; Meola, N.; Lubas, M.; Bukenborg, J.; Andersen, J.S.; Schulze, W.M.; Verheggen, C.; et al. Mutually Exclusive CBC-Containing Complexes Contribute to RNA Fate. Cell Rep. 2017, 18, 2635–2650. [CrossRef] [PubMed]
33. Nguyen, T.H.D. Structural Biology of Human Telomerase: Progress and Prospects. Biochem. Soc. Trans. 2021, 49, 1927–1939. [CrossRef] [PubMed]
34. Mitchell, J.R.; Collins, K. Human Telomerase Activation Requires Two Independent Interactions between Telomerase RNA and Telomerase Reverse Transcriptase. Mol. Cell 2000, 6, 361–371. [CrossRef]
35. Tesmer, V.M.; Ford, L.P.; Holt, S.E.; Frank, B.C.; Yi, X.; Aisner, D.L.; Ouellette, M.; Shay, J.W.; Wright, W.E. Two Inactive Fragments of the Integral RNA Cooperate To Assemble Active Telomerase with the Human Protein Catalytic Subunit (HTERT) In Vitro. Mol. Cell Biol. 1999, 19, 6207–6216. [CrossRef]
36. Nguyen, T.H.D.; Tam, J.; Wu, R.A.; Greber, B.J.; TosO, D.; Nogales, E.; Collins, K. Cryo-EM Structure of Substrate-Bound Human Telomerase Holoenzyme. Nature 2018, 557, 190–195. [CrossRef]
37. Tseng, C.-K.; Wang, H.-F.; Schroeder, M.R.; Baumann, P. The H/ACA Complex Disrupts Triplex in HTR Precursor to Permit Processing by RRP6 and PARN. Nat. Commun. 2018, 9, 5430. [CrossRef]
38. Vogan, J.M.; Zhang, X.; Youmans, D.T.; Regalado, S.G.; Johnson, J.Z.; Hockemeyer, D.; Collins, K. Minimized Human Telomerase Maintains Telomeres and Resolves Endogenous Roles of H/ACA Proteins, TCAB1, and Cajal Bodies. Elife 2016, 5, e18221. [CrossRef]
39. Chen, L.; Roake, C.M.; Freund, A.; Batista, P.J.; Tian, S.; Yin, Y.A.; Gajera, C.R.; Lin, S.; Lee, B.; Pech, M.F.; et al. An Activity Switch in Human Telomerase Based on RNA Conformation and Shaped by TCAB1. Cell 2018, 174, 218–230.e13. [CrossRef]
40. Huang, J.; Brown, A.F.; Wu, J.; Xue, J.; Bley, C.J.; Rand, D.P.; Wu, L.; Zhang, R.; Chen, J.J.-L.; Lei, M. Structural Basis for Protein-RNA Recognition in Telomerase. Nat. Struct. Mol. Biol. 2014, 21, 507–512. [CrossRef]
41. Ghanim, G.E.; Fountain, A.J.; van Roon, A.-M.M.; Rangan, R.; Das, R.; Collins, K.; Nguyen, T.H.D. Structure of Human Telomerase Holoenzyme with Bound Telomeric DNA. Nature 2021, 593, 449–453. [CrossRef] [PubMed]
42. Zemora, G.; Handl, S.; Waldisch, C. Human Telomerase Reverse Transcriptase Binds to a Pre-Organized HTR Template. Nucleic Acids Res. 2016, 44, 413–425. [CrossRef] [PubMed]
43. Niederer, R.O.; Zappulla, D.C. Refined Secondary-Structure Models of the Core of Yeast and Human Telomerase RNAs Directed by SHAPE. RNA 2015, 21, 254–261. [CrossRef] [PubMed]
44. Theimer, C.A.; Finger, L.D.; Trantirek, L.; Feigon, J. Mutations Linked to Dyskeratosis Congenita Cause Changes in the Structural Equilibrium in Telomerase RNA. Proc. Natl. Acad. Sci. USA 2003, 100, 449–454. [CrossRef]
45. Zhang, Q.; Kim, N.-K.; Peterson, R.D.; Wang, Z.; Feigon, J. Structurally Conserved Five Nucleotide Bulge Determines the Overall Topology of the Core Domain of Human Telomerase RNA. Proc. Natl. Acad. Sci. USA 2010, 107, 18761–18768. [CrossRef]
46. Roake, C.M.; Chen, L.; Chakravartthy, A.L.; Ferrrell, J.E.; Raffa, G.D.; Artandi, S.E. Disruption of Telomerase RNA Maturation Kinetics Precipitates Disease. Mol. Cell 2019, 74, 688–700.e3. [CrossRef]
47. Robart, A.R.; Collins, K. Investigation of Human Telomerase Holoenzyme Assembly, Activity, and Processivity Using Disease-Linked Subunit Variants. J. Biol. Chem. 2010, 285, 4375–4386. [CrossRef]
48. Errington, T.M.; Fu, D.; Wong, J.M.Y.; Collins, K. Disease-Associated Human Telomerase RNA Variants Show Loss of Function for Telomere Synthesis without Dominant-Negative Interference. Proc. Natl. Acad. Sci. USA 2010, 107, 6510–6520. [CrossRef]
49. Comolli, L.R.; Smirnov, I.; Xu, L.; Blackburn, E.H.; James, T.L. A Molecular Switch Underlies a Human Telomerase Disease. Proc. Natl. Acad. Sci. USA 2002, 99, 16998–17003. [CrossRef]
50. Chen, J.-L.; Greider, C.W. Functional Analysis of the Pseudoknot Structure in Human Telomerase RNA. Proc. Natl. Acad. Sci. USA 2005, 102, 8080–8085. [CrossRef]
51. Deshpande, A.P.; Collins, K. Mechanisms of Template Handling and Pseudoknot Folding in Human Telomerase and Their Manipulation to Expand the Sequence Repertoire of Procesive Repeat Synthesis. Nucleic Acids Res. 2018, 46, 7886–7901. [CrossRef] [PubMed]
52. Antal, M.; Boros, E.; Solymosy, F.; Kiss, T. Analysis of the Structure of Human Telomerase RNA in Vivo. Nucleic Acids Res. 2002, 30, 912–920. [CrossRef] [PubMed]
53. Leeper, T.C.; Varani, G. The Structure of an Enzyme-Activating Fragment of Human Telomerase RNA. *RNA* 2005, 11, 394–403. [CrossRef] [PubMed]

54. Kim, N.-K.; Theimer, C.A.; Mitchell, J.R.; Collins, K.; Feigon, J. Effect of Pseudouridylation on the Structure and Activity of the Catalytically Essential P6.1 Hairpin in Human Telomerase RNA. *Nucleic Acids Res.* 2010, 38, 6746–6756. [CrossRef]

55. Palka, C.; Forino, N.M.; Hentschel, J.; Das, R.; Stone, M.D. Folding Heterogeneity in the Essential Human Telomerase RNA Three-Way Junction. *RNA* 2020, 26, 1787–1800. [CrossRef] [PubMed]

56. Cristofari, G.; Adolf, E.; Reichenbach, P.; Sikora, K.; Terns, R.M.; Terns, M.P.; Lingner, J. Human Telomerase RNA Accumulation in Cajal Bodies Facilitates Telomerase Recruitment to Telomeres and Telomere Elongation. *Mol. Cell* 2007, 27, 882–889. [CrossRef]

57. Zhu, Y.; Tomlinson, R.L.; Lukowiak, A.A.; Terns, R.M.; Terns, M.P. Telomerase RNA Accumulates in Cajal Bodies in Human Cancer Cells. *Mol. Biol. Cell* 2004, 15, 81–90. [CrossRef]

58. Cheng, Y.; Liu, P.; Zheng, Q.; Gao, G.; Yuan, J.; Wang, P.; Huang, J.; Xie, L.; Lu, X.; Tong, T.; et al. Mitochondrial Trafficking and Processing of Telomerase RNA TERC. *Cell Rep.* 2018, 24, 2589–2595. [CrossRef]

59. Gazzaniga, F.S.; Blackburn, E.H. An Antia apoptotic Role for Telomerase RNA in Human Immune Cells Independent of Telomere Integrity or Telomerase Enzymatic Activity. *Blood* 2014, 124, 3675–3684. [CrossRef]