TENT4A Non-Canonical Poly(A) Polymerase Regulates DNA-Damage Tolerance via Multiple Pathways That Are Mutated in Endometrial Cancer

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Abstract: TENT4A (PAPD7) is a non-canonical poly(A) polymerase, of which little is known. Here, we show that TENT4A regulates multiple biological pathways and focuses on its multilayer regulation of translesion DNA synthesis (TLS), in which error-prone DNA polymerases bypass unrepaired DNA lesions. We show that TENT4A regulates mRNA stability and/or translation of DNA polymerase η and RAD18 E3 ligase, which guides the polymerase to replication stalling sites and monoubiquitinates PCNA, thereby enabling recruitment of error-prone DNA polymerases to damaged DNA sites. Remarkably, in addition to the effect on RAD18 mRNA stability via controlling its poly(A) tail, TENT4A indirectly regulates RAD18 via the tumor suppressor CYLD and via the long non-coding antisense RNA PAXIP1-AS2, which had no known function. Knocking down the expression of TENT4A or CYLD, or overexpression of PAXIP1-AS2 led each to reduced amounts of the RAD18 protein and DNA polymerase η, leading to reduced TLS, highlighting PAXIP1-AS2 as a new TLS regulator. Bioinformatics analysis revealed that TLS error-prone DNA polymerase genes and their TENT4A-related regulators are frequently mutated in endometrial cancer genomes, suggesting that TLS is dysregulated in this cancer.

Keywords: TLS; DNA repair; poly(A) RNA polymerase; translesion; lesion bypass

1. Introduction

Maintaining genome integrity is critical for the proper functioning of cells and if compromised may lead to severe malfunction and a variety of diseases including cancer, immune malfunction, and neuronal disorders [1–6]. The majority of DNA lesions inflicted on DNA are removed by accurate DNA repair mechanisms that restore the original DNA sequence. However, the high rate at which DNA lesions are formed (estimated to be thousands/cell/day) and the difficulty in locating them in the genome, makes encounters with DNA lesions inevitable in essentially every cell cycle. Such encounters cause arrest of the replication fork and if not resolved may lead to fork collapse, double-strand breaks (DSBs) and subsequent chromosomal rearrangements, or even cell death. DNA damage tolerance (DDT) mechanisms function in these situations to bypass the
lesions, without removing them from DNA, a process that preserved the double-stranded continuity of DNA, later giving the chance for an accurate excision repair mechanism to remove the lesion lesion [7–9].

Translesion DNA synthesis (TLS), is a DDT mechanism in which specialized low-fidelity DNA polymerases synthesize across the lesion, a process that is inherently mutagenic [10]. Despite its mutagenic nature, TLS is surprisingly accurate when replicating across several common DNA lesions such as the sunlight-induced cyclobutane pyrimidine dimers (CPD) and the tobacco-smoke induced (+)-trans-BPDE-N2-dG (BP-G) adduct. To explain this, we have previously proposed that the existence of multiple TLS DNA polymerases enables specialization of certain polymerases to their cognate lesions (meaning more effective and more accurate bypass). When subjected to the appropriate regulation to ensure the activity of the suitable polymerase at the right lesion and the right time, this will lead to more accurate TLS and a lower mutation burden [11–13]. We also reported, that p53 and its target gene p21 (acting via its interaction with PCNA), are needed to maintain accurate TLS, partially via their effect on monoubiqutitnation of PCNA [14]. The latter is an important regulatory step, which functions to recruit TLS DNA polymerases to the damaged site in DNA [15–17]. The molecular deciphering of the regulatory mechanisms of TLS is critical to the understanding of mutation formation and is the focus of the current study.

TENT4A (terminal nucleotidyltransferase 4A; formerly P APD7), is a gene encoding a non-canonical poly(A) RNA polymerase, which we identified in an siRNA-based screen to be required for efficient TLS [18]. TRF4, its S. cerevisiae homolog, was thought to encode a novel DNA polymerase, termed POLS (or erroneously, POLK), involved in sister chromatid cohesion [19,20]. However, further studies showed that the TRF4 and TRF5 genes encode non-canonical poly(A) RNA polymerases [20]. Enzymes of this family are critical in the regulation and quality control of gene expression. There is scarce information about the mammalian TENT4A. However, it was reported that its main isoform is 94kDa, considerably larger than the yeast Trf4 and Trf5 proteins (66 and 74 kDa, respectively) and larger than its previously reported size of 62 kDa [21]. Recently TENT4A was shown along with its paralog TENT4B (PAPD5) to synthesize mixed poly(A) tails that contain other inserted nucleotides, primarily guanines, that protect mRNA from deadenylation [22]. Still, the functional involvement of TENT4A in biological regulatory pathways is largely unknown [23].

Here, we report that TENT4A regulates a broad range of genes belonging to various pathways. Focusing on its involvement in TLS, we found that TENT4A regulates mRNA stability and/or translation of DNA polymerase η (polη; POLH) and of the RAD18 E3 ligase. Remarkably, in addition to the effect on RAD18-mRNA stability via controlling its poly(A) tail, TENT4A indirectly regulates RAD18 and polη via the tumor suppressor CYLD and the long non-coding antisense RNA PAXIP1-AS2, which had no known function, highlighting it as a novel TLS regulator. We also partially purified TENT4A and show that it is a poly(A) RNA polymerase, with no detected DNA polymerase activity. Finally, we report that bioinformatics analysis of mutations in the genomes of 33 cancer types from the TCGA database revealed that components of TENT4A-regulated TLS are frequently mutated in endometrial cancer, suggesting an involvement of dysregulated TLS in the development of this type of cancer.

2. Results
2.1. TENT4A Is Needed for Effective TLS

A screen performed in our lab, identified TENT4A (PAPD7; POLS) as a novel TLS gene [18]. To further confirm its effect on TLS, we used the gap-lesion plasmid-based assay, with DNA constructs each containing a single defined lesion lesion [17,18,24] and two cell lines: The human osteosarcoma U2OS cell line and the MCF-7 breast cancer cell line. As can be seen in Figure 1A–C, Supplementary Materials Figure S1 and Tables S1 and S4, knocking down the expression of TENT4A in these cells, but not of TENT4B, led to a decrease in TLS across four types of DNA lesions: a BP-G adduct, a major lesion caused by tobacco smoke; a thymine-thymine cyclobutane pyrimidine dimer (TT-CPD) and a thymine-thymine 6–4...
photoproduct (TT 6-4PP), the most common and the second most common type of UV light-induced DNA damage, respectively; and a cisplatin guanine-guanine (cisPt-GG), a major intrastrand crosslink formed by the chemotherapeutic drug cisplatin. DNA sequence analysis showed that the error frequency of the residual TLS under TENT4A knockdown did not substantially change (Supplementary Materials Tables S2, S3, S5 and S6). The broad DNA damage spectrum of the effect suggests that TENT4A regulation of TLS is not directed to a specific TLS DNA polymerase, but instead functions in a rather more general TLS regulatory step that exerts a global effect.

Figure 1. Involvement of TENT4A in TLS and its poly(A) RNA polymerase enzyme activity. (A) TLS across a TT 6-4PP and BP-G in U2OS cells. TENT4A expression was knocked down using lentivirus-mediated shRNA combined with siRNA against TENT4A. The results are presented as the mean ± SEM of three independent experiments (see Supplementary Materials Table S1 for details). Statistical analysis was performed using the two-tailed Student’s t-test (**p < 0.01). (B,C) TLS across a cisPt-GG and TT-CPD, respectively, in MCF-7 cells in which the expression of TENT4A and/or TENT4B genes was knocked down using specific siRNAs. The results are presented as the mean ± SEM of six independent experiments for cisPt-GG and three independent experiments for TT-CPD (see Supplementary Materials Table S4 for details). Statistical analysis was performed using the two-tailed Student’s t-test (****p < 0.0001, ***p < 0.001, **p < 0.01, *p < 0.05). (D) Poly(A) RNA
polymerase activity measured by the extension of an oligo (A)$_{20}$ substrate. Poly(A) polymerase assays were performed with the indicated amount of partially purified recombinant human 8xHis-MBP tagged TENT4A (lanes 1–3 and 9–11) or the TENT4A mutant D277A, D279A (lanes 4–6 and 12–14), using 5$^{32}$P-labeled oligo(A)$_{20}$ and 1 mM ATP in the presence of 5 mM MgCl$_2$ (lanes 1–6) or 1 mM MnCl$_2$ (lanes 9–14). For positive and negative controls, parallel reactions were also carried out with 0.5 units of E. coli poly(A) polymerase (lanes 8 and 16) or no added protein (lanes 7 and 15), respectively. The 5$^{32}$P-labeled RNA oligo sequence is shown below the image. Reaction products were resolved on a 15% polyacrylamide gel containing 8 M urea and analyzed by Phosphorimaging. (E) Poly(A) RNA polymerase assays with an oligo RNA. The assays were performed as described in panel (D), except that the oligo RNA shown underneath the gel image was used, as well as the four NTPs (1 mM each). (F) Ribonucleotide specificity of TENT4A. The $^{32}$P-labeled RNA oligo(A)$_{20}$ substrate was incubated with the TENT4A protein (lanes 1–5) or its mutant (lanes 6–10) in the presence of each of the four NTPs (1 mM each). (G) Assays of TENT4A DNA polymerase activity. Assays were performed with the indicated amount of partially purified recombinant human 8xHis-MBP tagged TENT4A (lanes 2–4 and 10–12) and mutant (lanes 5–7 and 13–15), using 0.5 pmol of $^{32}$P-labeled 15/60-nt primer/template in the presence of the four dNTPs (100 µM each), 5 mM MgCl$_2$ (lanes 2–7) or 1 mM MnCl$_2$ (lanes 10–15). For positive and negative controls, parallel reactions were carried out with the Klenow fragment of Pol I (lanes 1 and 9), or no added protein (lanes 8 and 16), respectively. The sequence of the primer/template DNA substrate is shown below the image.

2.2. Purified TENT4A Is a Non-Canonical Poly(A) RNA Polymerase, Which Also Incorporates Other Ribonucleotides

We partially purified overexpressed full-length human TENT4A as an 8xHis-MBP-tagged protein, (Supplementary Materials Figure S2) and assayed its potential activities as a poly(A) RNA polymerase and DNA polymerase. In parallel, we purified a TENT4A variant that carries the D277A and D279A mutations (Supplementary Materials Figure S2), which were expected to inactivate the coordination of the divalent metal ion as a co-substrate. As can be seen in Figure 1D, TENT4A used ATP in the presence of Mg$^{++}$ to extend the oligo (A)$_{20}$ substrate. In contrast, the TENT4A mutant was inactive. A similar activity was observed with the oligo 5$^{\prime}$r(AGAGUUGAUCCUGGCUCGA)-3$^{\prime}$ (Figure 1E). Unlike E. coli poly(A) RNA polymerase, which was used as a positive control and performed extensive extension of the substrate, TENT4A added a limited number of AMP residues, which is typical of a non-canonical poly(A) RNA polymerase activity (Figure 1D,E). The polyadenylation activity of TENT4A was stimulated when Mn$^{++}$ was used instead of Mg$^{++}$, whereas the mutant TENT4A remained unaffected (Figure 1D,E). TENT4A preferentially polymerizes AMP residues, although it can use other rNTPs to some extent (Figure 1F), consistent with a previous report [22]. Of note, the shorter form of TENT4A, previously believed to be the full-length enzyme, had negligible poly(A) RNA polymerase activity (not shown). We also examined the DNA polymerase activity of the recombinant tagged TENT4A and found no activity (Figure 1G). We conclude that TENT4A is a poly(A) RNA polymerase and as far as we can tell, it has no detectable DNA polymerase activity.

2.3. Effect of TENT4A on mRNA of Genes Involved in TLS

Poly(A) RNA polymerases are typically involved in regulating mRNA via 3'$'$ polyadenylation and therefore to start addressing the mechanism by which TENT4A regulates TLS, we examined whether the mRNA of TLS-related genes bind TENT4A. To that end, we used RNA immunoprecipitation-qPCR (RIP-qPCR). We overexpressed FLAG-TENT4A, then immunoprecipitated the protein using an anti-FLAG antibody and extracted the RNA from protein-RNA complexes. The amount of specific mRNA was determined by preparing cDNA and performing qPCR using gene-specific primers. As can be seen in Figure 2A, while the mRNAs of POLH, POLK and POLI showed no significant binding, the mRNAs of REV3L encoding the catalytic subunit of DNA polymerase ζ and of REV1, encoding a TLS scaffold protein/polymerase, as well as RAD18 and CYLD, showed significant preferential binding. RAD18 is the E3 ligase that monoubiquitinates PCNA [25], a central signaling event in TLS.
and CYLD is a deubiquitinase involved in cancer and required for efficient TLS, as we have previously reported [18].

Figure 2. Effect of TENT4A on binding and stability of TLS-related mRNAs. (A) FLAG-tagged TENT4A expressed in HEK293T cells was immunoprecipitated with an anti-FLAG antibody and RNA bound to it was analyzed by qPCR. The results are shown as fold-change relative to empty vector. The error bars indicate SEM (two independent experiments). (B–D) Representative gel images of RAD18, CYLD and POLH poly(A) tail are shown, each with the poly(A) tail densitometric trace underneath, using Image J. The position of TVN is indicated by an asterisk that predominantly recognizes A12, serving as an internal control. (E–H), TENT4A and/or TENT4B expression was knocked down in MCF-7 cells for 48 h, after which the cells were treated with 5 µg/mL Actinomycin D for up to 8 h. The expression levels of RAD18, CYLD, POLH and POLD3 genes were each determined by qPCR relative to the expression level at time 0. Half-life was calculated by using non-linear fit one-phase decay curve equation. The significance of the differences was calculated using Student’s t-test (one-tailed); Data are presented as mean ± SEM from three independent experiments. ns, non-significant; p > 0.05.

To examine whether TENT4A affects the length of the poly(A) tail of these genes, we used the ePAT (extension poly(A) test; [26]). We examined the TLS regulators RAD18 and CYLD, which bound TENT4A and also POLH, which did not bind TENT4A. As shown in Figure 2, knocking down the expression of TENT4A caused a shift to shorter amplified tail fragments of RAD18 and similar results were obtained for the tail fragments
of CYLD (Figure 2B,C). Interestingly, knocking down the expression of TENT4A caused a shift to shorter amplified tail fragments of POLH too, despite its lack of binding to TENT4A, suggesting indirect regulation. Knocking down both TENT4A and TENT4B caused a similar shift (Figure 2B–D). Analysis of the GAPDH mRNA as a control with a long poly(A) tail, showed marginal, if any, effect of TENT4A and TENT4B knockdown (Supplementary Materials Figure S3).

As the length of the poly(A) tail may affect mRNA stability, we examined the effect of TENT4A on the stability of the mRNA of these TLS-related genes by measuring their half-life in the presence of the transcription inhibitor Actinomycin D. As can be seen in Figure 2E, knocking down the expression of TENT4A, but not TENT4B, caused a decrease of about 33% in the half-life of RAD18 mRNA, whereas knocking down the expression of both caused a slightly more substantial decrease of 43%. The half-life of CYLD mRNA was decreased by 53% when the expression of TENT4A was knocked down, with a smaller effect of TENT4B knockdown (Figure 2F). Interestingly, for POLH mRNA, knocking down the expression of TENT4A had a decrease in half-life (29%), slightly lower than the effect of TENT4B knockdown (39% decrease) (Figure 2G). The stability of a control POLD3 mRNA was essentially unaffected by knocking down either TENT4A, TENT4B or both (Figure 2H), as was the stable RNA18S rRNA (Supplementary Materials Figure S4). These results suggest that TENT4A directly regulates the stability of RAD18 and CYLD mRNA and indirectly the stability of POLH mRNA via the length of their poly(A) tails.

2.4. TENT4A Affects the Amounts of RAD18, CYLD and POLH Proteins via Different Mechanisms

Since effects on mRNA are often, but not always, manifested in the levels of the encoded proteins, we next analyzed whether TENT4A also affects the protein products of the RAD18, POLH and CYLD genes. As can be seen in Figure 3A, upon knocking down the expression of TENT4A, there was a decrease in the amount of RAD18, observed in both unirradiated and UV-irradiated cells. Knocking down TENT4A expression also led to a decrease in the amounts of the CYLD protein (Figure 3B) and polη (POLH gene product) (Figure 3B).

To explore whether this reduction in the amounts of the RAD18, CYLD and POLH proteins resulted from reduced translation efficiency, we performed polysome profiling under a condition in which TENT4A expression was knocked down. As shown in Figure 3C, there was an increase in the 80S monosome and a decrease in the polysomal fractions when TENT4A expression was knocked down, indicating an inhibition of translation efficiency. Analysis of individual genes revealed a shift from the heavy to the light fractions of CYLD and POLH mRNA (Figure 3E,F), but not of RAD18 mRNA (Figure 3D). Translation of the control gene POLD3 (Figure 3H) was unaffected by knocking down the expression of TENT4A. Thus, the reduction in CYLD and POLH protein amount following TENT4A knockdown is due to both their reduced mRNA stability and its reduced translation efficiency. However, the decreased amount of RAD18 protein following TENT4A knockdown is due to reduced mRNA stability, as the translation efficiency was unaffected.
Figure 3. Effect of TENT4A on levels and translation of TLS proteins. (A) MCF-7 cells were transfected with TENT4A-targeted siRNA (siTENT4A) or non-targeting control siRNA (siControl). At 65 h post-transfection, the cells were UV-irradiated at 20 J/m² and harvested 1 or 6 h post-irradiation. Whole-cell extracts were fractionated by SDS-PAGE followed by Western blot with indicated antibodies. Protein levels were normalized to GAPDH and presented each relative to its level in unirradiated siControl-treated cells. (B) Effect of cell treatment (72 h) with siTENT4A or siControl on the levels of the CYLD and POLH proteins in MCF-7 cells. Whole-cell extracts were fractionated by SDS-PAGE and analyzed by Western blot with indicated antibodies. Protein levels were normalized to GAPDH and presented each relative to its level in siControl-treated cells. (C) Polysome profile analysis in MCF-7 cells in which TENT4A expression was knocked down. Absorbance trace of polysome fractionation on a sucrose gradient. (D) qPCR analysis of mRNA levels of TLS-related genes in each fraction was expressed as a percentage of total levels summed over all fractions. The results are presented as the mean ± SEM of two independent experiments. (E–G) Fraction of polysome-associated mRNA of the genes CYLD, POLH and GAPDH, respectively. The values were taken from (D). (H) Fraction of polysome-associated mRNA of the POLD3 control gene. The results are presented as the mean ± SEM of two independent experiments. Statistical significance was determined using Student’s t-test (one-tailed); * p < 0.05, ns, non-significant; p > 0.05.

2.5. Regulation of PCNA Monoubiquitination by TENT4A

RAD18 is the key E3 ligase that monoubiquitinates PCNA and we therefore examined whether the TENT4A effect of reducing the amounts of RAD18 is also manifested in the level of mUb-PCNA. As shown in Figure 4, upon TENT4A knockdown the amount of mUb-PCNA was mildly decreased in UV-irradiated cells (Figure 4, lane 10) and strongly decreased in
unirradiated cells (Figure 4, lane 2). The amount of mUb-PCNA was calculated as the mUb-PCNA/PCNA ratio and as the amount (independent of PCNA) normalized to the loading control GAPDH, generally showing a similar behavior upon TENT4A knockdown.

Figure 4. Interrelationship among TENT4A, RAD18 and DTL in determining amounts of RAD18 and DTL and the extent of PCNA monoubiquitination. MCF-7 cells were transfected with siRNA against TENT4A, RAD18 and DTL or their double or triple combinations. At 65 h post-transfection, the cells were UV-irradiated at 20 J/m² and harvested 6 h post-irradiation. Whole-cell extracts were fractionated by SDS-PAGE followed by Western blot with indicated antibodies. Protein levels were normalized to GAPDH and presented each relative to its level in unirradiated or UV-irradiated siControl-treated cells.

PCNA monoubiquitination in untreated cells was reported to be carried out by the DTL (CTD2) E3 ligase [27] and we therefore examined the effect of TENT4A knockdown on DTL. As can be seen in Figure 4 lanes 2 and 10, knocking down the expression of TENT4A caused a strong reduction in the amount of the DTL protein, in both unirradiated and UV-irradiated cells. In parallel, also the amount of the RAD18 protein was decreased, consistent with the results shown above. We then examined whether RAD18 and DTL affect each other. Interestingly, when the expression of DTL was knocked down, the amount of the RAD18 protein was also reduced in both unirradiated (Figure 4, lane 4) and UV-irradiated cells (Figure 4, lane 12). On the other hand, when the expression of RAD18 alone was knocked down, the amount of DTL was increased, in both unirradiated (Figure 4, lane 3) and UV-irradiated cells (Figure 4, lane 11). When both TENT4A and RAD18 were knocked down, DTL was still decreased, but to an extent lesser than with TENT4A knockdown alone, indicating an additive effect (Figure 4, lanes 5 and 13). On the other hand, the TENT4A and DTL double knockdown had a similar effect to the single gene knockdown, indicating the two are epistatic (Figure 4, lanes 6 and 14).

How do these effects on RAD18 and DTL translate into levels of mUb-PCNA? In UV-irradiated cells RAD18 knockdown caused a strong reduction in mUb-PCNA, despite the increase in DTL, consistent with RAD18 being the major E3 ligase that monoubiquitinates PCNA in UV-irradiated cells, with a similar effect observed with the TENT4A and RAD18 double knockdown (Figure 4, lanes 11 and 13). Interestingly, knocking down DTL caused a mild decrease in mUb-PCNA in UV-irradiated cells (Figure 4, lane 12), likely because, under these conditions RAD18 was also decreased. In unirradiated cells, knocking down the expression of DTL caused a strong decrease in mUb-PCNA, consistent with previous work [27], but also knocking down RAD18 caused a decrease in mUb-PCNA (Figure 4, lane 3), despite a slight increase in the amount of DTL. Similar effects were observed when both
TENT4A and DTL were knocked down (Figure 4, lane 6). When both TENT4A and RAD18 were knocked down, mUb-PCNA was strongly reduced (Figure 4, lane 5), as under these conditions the two E3 ligases were reduced.

2.6. TENT4A-Regulated CYLD Regulates RAD18, DTL and POLH

CYLD, previously identified by us in a screen as a TLS regulator, acts downstream to TENT4A and is regulated by TENT4A at the mRNA stability and translation levels as shown above. To examine its own effect on TLS, we knocked down CYLD expression and examined levels of RAD18, mUb-PCNA and POLH. As can be seen in Figure 5A, knocking down CYLD expression caused a strong three-to-five-fold decrease in the amount of RAD18, clearly visible in both unirradiated and UV-irradiated cells. Similarly, POLH protein decreased about 3-fold (Figure 5A). It thus appears that in addition to its direct effect on RAD18, TENT4A has an indirect effect on both RAD18 and POLH proteins, mediated via its target gene CYLD.

Figure 5. Effect of CYLD on RAD18, DTL, POLH and PCNA monoubiquitination. (A) MCF-7 cells were transfected with CYLD-targeted siRNA (siCYLD) or siControl. At 65 h post-transfection, the cells were irradiated at 20 J/m² UV and harvested 1 or 6 h post-irradiation. (B) MCF-7 cells were transfected with siRNA against CYLD, RAD18 and DTL or their double or triple combinations. At 65 h post-transfection, the cells were UV-irradiated at 20 J/m² and harvested 6 h post-irradiation. Whole-cell extracts were fractionated by SDS-PAGE and analyzed by Western blot with indicated antibodies. Protein levels were normalized to GAPDH and presented each relative to its level in unirradiated or UV-irradiated siControl-treated cells.
An analysis of mUb-PCNA in cells in which CYLD was knocked down revealed that despite the reduction in RAD18 protein, UV-induced mUb-PCNA levels were little affected (Figure 5A). Assuming that DTL might be backing up RAD18 under these conditions, we knocked down the expression of CYLD, RAD18 or DTL individually, or in combinations of two or three genes. As shown in Figure 5B, knocking down CYLD caused a strong decrease of both RAD18 and DTL, in both unirradiated (Figure 5B, lane 2) and UV irradiated cells (Figure 5B, lane 10). Of note, a decrease in RAD18 protein was not always paralleled by a decrease in mUb-PCNA (Figure 5B).

2.7. TENT4A Regulates Multiple Genes and Pathways, Including Long Non-Coding RNAs and Antisense Transcripts

As TENT4A, being a poly(A) RNA polymerase, is expected to exert its effect via RNA, we conducted an RNA-seq analysis, comparing genes in control cells to the same cells in which TENT4A was knocked down. The experiments were conducted in triplicates in two different human cell lines, in which the expression of TENT4A was knocked down with shRNA plus siRNA (U2OS), or siRNA (XP12RO). Overall, the expression of many genes was altered (Figure 6A and Supplementary Materials Data S1), with an intersection of 313 genes downregulated and 149 genes upregulated in the two cell lines (Figure 6A). A heatmap showed that the overall expression pattern of the shared genes was similar in the two cell lines (Figure 6B).

Gene set enrichment analysis (GSEA; Broad Institute) revealed that the significantly downregulated genes in TENT4A cells, corresponded to 44 gene sets that were enriched in both cell lines with an FDR of <0.05 (Supplementary Materials Data S2). No gene sets with upregulated genes were common for both cell lines. Of note, Pathway Studio analysis revealed among genes whose expression was upregulated, also antisense transcripts and long untranslated RNA (Figure 6C). Overall, the biological pathways with genes that were downregulated can be categorized to five main processes, namely (1) Cell division; (2) Membranes, vesicles and Golgi; (3) Metabolism; (4) Proteolysis; (5) RNA processing; (Figure 6D). Thus, TENT4A appears to regulate many genes involved in diverse biological processes, including mRNA degradation, as expected from a poly(A) RNA polymerase and consistent with its effect on the half-life of several TLS-related mRNAs (Figure 2). Of note, Gene Ontology defined DNA repair pathways were downregulated upon treatment with siRNA against TENT4A (Supplementary Materials Data S2).
Figure 6. RNA-seq analysis of cells in which the expression of TENT4A was knocked down. (A) Venn diagram illustrating the number of significantly differentially expressed genes (FDR ≤ 0.05, absolute fold change ≥2 and a count of at least 30 at least in one of the samples) in TENT4A knocked down U2OS and XP12RO cells. The numbers of upregulated, downregulated and overlapping genes are shown (see detailed RNA-seq results in Supplementary Materials Data S1). (B) Heatmap representation of the shared significantly differentially expressed genes in U2OS and XP12RO experiments (a total of 475 genes). The response of most genes is the same in both cell lines (149 are upregulated and 313 are downregulated). For each experiment, the log2 normalized counts were standardized to have for each gene zero mean and unit standard deviation. K-mean clustering of the two data sets using Euclidean distance measure is presented. The expression profile is accompanied by a colored bar indicating the standardized log2 normalized counts. (C) Pathway Studio ontology of differentially expressed upregulated genes that are common to U2OS and XP12RO analyzed by Pathway Studio. Ontology details are in Supplementary Materials Data S2. (D) Gene sets enriched in downregulated genes in TENT4A knocked down U2OS and XP12RO cells. The mRNA degradation pathway was from the Pathway Studio analysis. See details data in Supplementary Materials Data S2.

2.8. A TENT4A-Regulated Long Non-Coding Antisense RNA to the PAXIP1 (PTIP) Gene Regulates RAD18, POLH and mUb-PCNA

As mentioned above, the RNA-seq analysis revealed that knocking down the expression of TENT4A causes upregulation of long non-coding RNA and antisense transcripts. One of these is PAXIP1-AS2, which is a long non-coding RNA, antisense to the PAXIP1 gene. As PAXIP1 was reported to promote PCNA monoubiquitination [28], we studied the involvement in TLS of PAXIP1-AS2, for which no biological function was yet assigned.

Knocking down the expression of TENT4A caused a significant increase in the amount of the PAXIP1-AS2 transcript (Figure 7A), as expected from the RNA-seq analysis. This effect was not observed when TENT4B was knocked down, suggesting that it is specific to TENT4A (Figure 7A). We next overexpressed PAXIP1-AS2 and examined its effect on TLS. As shown in Figure 7B and Supplementary Materials Tables S7 and S8, TLS across cisPt-GG
was strongly suppressed by overexpressing PAXIP1-AS2. This effect was not mediated via the amount or stability of PAXIP1 mRNA, which remained essentially unchanged upon overexpression of the antisense RNA (Figure 7C,D); However, the amount of PAXIP1 protein was diminished (Figure 7E). Directly knocking down the expression of PAXIP1 also inhibited TLS across a TT-CPD lesion (Figure 7F and Supplementary Materials Tables S9 and S10), consistent with the effect of the antisense RNA.

Figure 7. Involvement of PAXIP1-AS2 and PAXIP1 in TLS. (A) Expression of TENT4A and/or TENT4B was knocked down for 48 h, after which the level of PAXIP1-AS2 transcript was measured by qPCR, normalized to GAPDH, and compared to cells treated with siControl. The results are presented as the mean ± SEM of three independent experiments. (B) TLS across a cisPt-GG in MCF-7 cells in which PAXIP1-AS2 antisense RNA was overexpressed (See Supplementary Materials Tables S7 and S8). The results are presented as the mean ± SEM of two independent experiments. Statistical analysis was performed using the two-tailed Student’s t-test (** p < 0.01). (C) PAXIP1-AS2 antisense RNA was overexpressed in XP12RO or MCF-7 cells for 48 h, after which the amount of PAXIP1 mRNA was measured by qPCR, normalized to GAPDH and presented relative to cells transfected with an empty vector (EV). The results are presented as the mean ± SEM of three independent experiments. (D) PAXIP1-AS2 antisense transcript was overexpressed for 48 h in MCF-7 cells, after which PAXIP1 mRNA levels were determined as described in Figure 2. GAPDH was used as normalized control.
The results are presented as the mean ± SEM of two independent experiments. (E) Effect of overexpression of PAXIP1-AS2 antisense RNA on PAXIP1 protein. Cells were harvested after 48 h and whole-cell extracts were analyzed by SDS-PAGE followed by Western blot with indicated antibodies. (F) TLS across a TT-CPD lesion in MCF-7 cells in which the expression of PAXIP1 was knocked down for 68 h. TT-CPD TLS assay was performed for 4 h. See Supplementary Materials Tables S9 and S10 for details. The results are presented as the mean ± SEM of two independent experiments. Statistical analysis was performed using the two-tailed Student’s t-test (*p < 0.05). (G) Effect of overexpression of PAXIP-AS2 on TLS-related proteins in UV-irradiated cells. PAXIP1-AS2 was overexpressed in XP12RO cells for 48 h after which cells were UV-irradiated at 10 J/m² UV and harvested 1 or 6 h post-irradiation. Whole-cell extracts were analyzed by SDS-PAGE followed by Western blot with indicated antibodies. Protein levels are presented relative to those in extracts of unirradiated cells transfected with empty vector. (H) Effect of PAXIP1 knockdown on TLS proteins. MCF-7 cells were knocked down by siRNA against PAXIP1 for 72 h, after which cells were UV-irradiated at 20 J/m² UV and harvested 1 or 6 h post-irradiation. Whole-cell extracts were analyzed by SDS-PAGE and detected with indicated antibodies. Amounts are presented relative to those in extracts of unirradiated cells treated with siControl.

We next examined the effects of PAXIP1 and PAXIP1-AS2 on POLH, RAD18 and mUb-PCNA. As shown in Figure 7G,H, PAXIP1-AS2 overexpression and similarly knockdown of PAXIP1, each caused a decrease in the amount of POLH, whereas the effect on RAD18 and mUb-PCNA was marginal.

As described above, TENT4A did not bind POLH mRNA, yet it affected its stability, raising the possibility that the effect was mediated via PAXIP1. Knocking down the expression of PAXIP1 reduced the half-life of POLH mRNA to an extent similar to knocking down the expression of TENT4A (Figure 8A). Knocking down both TENT4A and PAXIP1 caused a decrease of POLH mRNA stability similar to that of each gene alone, suggesting that the two are epistatic and that possibly the effect of TENT4A on POLH mRNA is mainly mediated via PAXIP1. Western blot analysis revealed that the decrease in polη amount was more pronounced under PAXIP1 than TENT4A knockdown (Figure 8C). Thus, while the effects of the two seem to be epistatic at the mRNA level, additional regulation affects the polη protein level by mechanisms that are yet to be explored, e.g., via a positive regulation by PAXIP1, or a branch of negative regulation by TENT4A. A similar epistatic analysis was performed with TENT4A and PAXIP1 for RAD18 mRNA, which binds TENT4A. Knocking down the expression of TENT4A had a bigger effect than PAXIP1 of decreasing the half-life of RAD18 mRNA, while knocking down the two had an intermediate effect (Figure 8B). A similar effect was observed also at the protein level, in both unirradiated and UV-irradiated cells (Figure 8C). Thus, the direct effect of TENT4A on RAD18 mRNA appears to be dominant over the PAXIP1 axis.
Figure 8. Effect of TENT4A and PAXIP1 on the stability of TLS-related mRNAs and level of TLS-related proteins. (A, B) TENT4A and/or PAXIP1 expression was knocked down in MCF-7 cells for 48 h, after which the cells were treated with 5 µg/mL Actinomycin D for up to 8 h. The expression levels of POLH and RAD18 were each determined by qPCR relative to the expression level at time 0. Half-life was calculated using a non-linear fit one-phase decay curve equation. The significance of the differences was calculated using Student’s t-test (one-tailed); Data are presented as mean ± SEM from three independent experiments. ns, non-significant; p > 0.05. (C) MCF-7 cells were transfected with TENT4A and/or PAXIP1 or non-targeting control siRNA for a final combined concentration of 100 nM. At 65 h post-transfection, the cells were UV-irradiated at 20 J/m² and harvested 1 or 6 h post-irradiation. Whole-cell extracts were fractionated by SDS-PAGE followed by Western blot with indicated antibodies. Amounts are presented relative to those in extracts of unirradiated cells treated with siControl and the values are shown in the corresponding position of the blots.

2.9. TENT4A and TLS Genes Are Frequently Mutated in Endometrial Cancer

The importance of TLS in tolerating DNA damage and affecting genetic stability prompted us to examine whether mutations in genes of the TLS pathway are over-represented in particular cancer types. To that end we used the TCGA database and first analyzed, for each cancer type, the percentage of the samples that contained mutations in at least one of the genes that we defined as a TLS-related genes group, containing TENT4A, CYLD, NPM1, TENT4B, PAXIP1, PCNA, POLH, POLI, POLK, PRIMPOL, RAD18, REV1, REV3L and USP1. As can be seen in Supplementary Materials Table S11, there is a big variation in the percent of samples with mutations in the TLS genes group, ranging from about 1–2% (e.g., Thyroid Carcinoma; THCA) up to 37% for Uterine Corpus Endometrial Carcinoma (UCEC; endometrial cancer). Cancer types with a higher overall number of mutations tend to exhibit a higher percentage of samples with mutations in TLS-related genes, as expected (e.g., Skin Cutaneous Melanoma, SKCM; Colon Adeno-
cancer, COAD; Lung Squamous Cell Carcinoma; LUSC; Supplementary Materials Table S11). However, endometrial cancer stands out with the highest occurrence of samples with mutations in the TLS genes group, despite a higher median overall number of mutations in several other cancers (e.g., Bladder Cancer, BLCA; COAD; Lung Adenocarcinoma, LUAD; Supplementary Materials Table S11).

To probe the statistical significance of the high percentage of samples with mutations in the TLS genes group, we estimated the probability that this high percentage was obtained by chance. To that end we chose a control gene set of 14 random genes (the size of the TLS-related gene group) and examined in each cancer type the percentage of samples with mutations in at least one gene of this control gene set. This was repeated 1000 times and used to plot the chance distribution and calculate the probability of the TLS genes group (Supplementary Materials Figure S5 and Table S11). Supplementary Materials Figure S5 also shows the average number of samples in each distribution and the fraction of samples with TLS-related genes group in each cancer type. The fraction of the 1000 random sets that yielded a number of samples with mutations higher than the number of TLS mutations represents the probability that the fraction of TLS-related genes group was obtained by chance and was termed the p-total value. Only three cancer types passed this statistical test with p-total < 0.05, namely acute myeloid leukemia (LAML), thymoma (THYM) and UCEC (Supplementary Materials Table S11 and Table 1A). For these three we tested the statistical significance of the difference between the number of samples with TLS mutations and the average number of samples with random mutation sets, using the Chi-squared test. Only LAML and UCEC passed this additional test with p < 0.05 (Table 1A). Analysis of the fraction of samples with mutations in each TLS gene (Table 1B), shows that UCEC has a considerable percentage of mutations indeed in each of the TLS-related genes, including TENT4A (Table 1B), whereas in the LAML, the majority of samples contained mutations in a single gene, namely NPM1 (Table 1B).

**Table 1.** Analysis of the frequency of TLS-related gene mutations in cancer genomes.

(A) Number of samples with mutations in TLS-related vs. random genes in top-ranking cancers.

| Cancer type b | Samples, n c | Samples with mutations, n | TLS-related genes d | Random genes, Ave e | P-total f | p-value g |
|--------------|--------------|-----------------|------------------|--------------------|----------|----------|
| LAML         | 144          | 21              | 5.03             | <0.001             | <0.001   |
| THYM         | 123          | 9               | 3.30             | 0.001              | 0.08     |
| UCEC         | 530          | 196             | 153.92           | 0.029              | 0.006    |

(B) Fraction of cancer samples with mutations in TLS-related genes.

| Cancer type b: | LAML | UCEC | Cancer type b: | LAML | UCEC |
|---------------|------|------|---------------|------|------|
| Gene          |      |      | Gene          |      |      |
| CYLD          | 1%   | 11%  | PRIMPOL       | 1%   | 6%   |
| NPMI          | 13%  | 3%   | RAD18         | 0%   | 6%   |
| PAXIP1        | 1%   | 11%  | REV1          | 0%   | 11%  |
| PCNA          | 0%   | 3%   | REV3L         | 2%   | 18%  |
| POLH          | 0%   | 6%   | TENT4A        | 0%   | 11%  |
| POLI          | 1%   | 7%   | TENT4B        | 1%   | 15%  |
| POLK          | 0%   | 8%   | USP1          | 0%   | 7%   |

a All mutations included. b Cancer type according to TCGA cancer acronyms. LAML, Acute Myeloid Leukemia; THYM, Thymoma; UCEC, Uterine Corpus Endometrial Carcinoma. c Number of samples for which sequence information is available in the TCGA database. d Number of samples with a mutation in at least one of 14 TLS-related genes listed in Table 1B. e Number of samples with a mutation in at least one of 14 random genes; Average of 1000 random sets. f p-value of the difference between the TLS-related genes and the average random genes (d and e). g First the average number of samples with mutations in random genes was rounded and then the statistical significance of the difference between the two groups (TLS and random) was estimated using the chi-squared test. For example, for LAML the groups are: 21 samples with TLS mutations and 123 without mutations in these genes, compared to 5 samples with mutations in a set of random genes and 139 samples without mutations in these genes.
3. Discussion

DNA damage tolerance by TLS must be carefully regulated to balance between the beneficial effect of enabling overcoming replication obstacles, with a minimal adverse effect on genetic stability. Indeed, no less than 17 genes were previously identified in our lab as new regulators of TLS [18], amongst them TENT4A (PAPD7). While initially this gene was proposed to encode in yeast a DNA polymerase, it was subsequently shown to encode a poly(A) RNA polymerase in both yeast [20] and human cells [22], with no detectable DNA polymerase activity, as confirmed in this study.

While little is known about the role of TENT4A in various biological processes, the current study shows that it has broad effects and is involved in multiple pathways. These include, most notably, RNA processing and mRNA degradation, membrane-related pathways such as membrane budding, vesicle-mediated transport and vesicle organization, metabolism, proteolysis, and cell division-related processes (Figure 6). The finding that TENT4A is involved in TLS, a pathway that has been studied in our lab over many years, prompted us to further explore the molecular mechanisms that underlie this involvement. Interestingly, the TENT4A-dependent regulation of TLS uncovered in the current study spans several major components of the TLS machinery, i.e., RAD18, DTL, POLH, and mUb-PCNA, in a complex interrelationship using both direct and indirect mechanisms (Figure 9). Interestingly, the CYLD tumor suppressor gene, previously identified as a TLS regulator [18], turned out to be acting downstream from TENT4A and was directly regulated by it at the mRNA stability and translation levels and PAXIP1-AS2 was identified as a new significant TENT4A-regulated TLS regulator. Of note, PAXIP1, which PAXIP1-AS2 regulates, is a paired box (PAX) gene with six BRCT domains, which is involved in the development of [29,30] and responses to DNA damage [31,32]. It is a suggested lung cancer tumor suppressor [33] and linked to breast and ovarian cancer and response to chemotherapy [34].

![Figure 9. A scheme representing the three branches of regulation of TLS by TENT4A. See text for details.](image-url)
While TENTA directly regulates RAD18 and CYLD via binding their mRNAs and controlling the length of their poly(A) tails and stability, consistent with its activity as non-canonical poly(A) RNA polymerase, POLH mRNA regulation appears to be indirect, since no binding of POLH mRNA to TENT4A protein was observed. It is likely mediated via the PAXIP1 protein (see below), which showed an effect similar and epistatic to TENT4A. Interestingly, TENT4A knockdown also had a global effect on translation, as indicated by the shift in ribosome profile and specifically manifested in the inhibited translation of the POLH and CYLD genes, but not RAD18.

Knocking down the expression of TENT4A or its downstream effector CYLD reduced the level of RAD18 and was therefore expected to be manifested in a decrease in mUb-PCNA. Unexpectedly, such a decrease was not necessarily observed, leading to the unraveling of an interesting relationship between TENT4A, RAD18 and DTL. First, in addition to RAD18, DTL is also regulated by TENT4A, whose knockdown strongly decreased the level of DTL protein. Second, knocking down the expression of DTL caused a decrease in the amount of the RAD18 protein, in both unirradiated and UV-irradiated cells. This is not due to cross targeting of the siRNAs, since they are gene-specific, raising the possibility that DTL acts upstream to RAD18. As knocking down the expression of CYLD also reduced the amount of DTL, the CYLD branch regulated by TENT4A may operate in the order of TENT4A ⇒ CYLD ⇒ DTL ⇒ RAD18.

Interestingly, knocking down the expression of RAD18 led to an increase in the amount of DTL protein, as if to compensate for the absence of this activity, possibly acting via a feedback loop. The fact that a reduction in the amount of RAD18 is not always manifested in a significant decrease in mUb-PCNA, may have been due to the remaining amounts of the RAD18, which were sufficient to carry out significant PCNA monoubiquitination. In this context, it should be pointed out that knocking down the expression of TENT4A caused a 40–80% decrease in the amount of RAD18 protein, depending on the experiment. This decrease was likely caused by post-transcriptional regulation, as the RNA-seq results showed up to a 20% decrease in the steady-state level of RAD18 mRNA (Supplementary Materials Data S1). Additionally, there might exist a third, as yet unknown E3 ligase that monoubiquitinates PCNA.

As expected from its sequence complementarity, we found that PAXIP1-AS2 is indeed a negative regulator of PAXIP1 (Figure 7E). However, somewhat unexpectedly, this regulatory effect was not mediated via an effect on the amount or stability of PAXIP1 mRNA (Figure 7C,D). It was previously reported that antisense long non-coding RNA could control translation [35] and a similar effect may explain the effect of PAXIP1-AS2.

PAXIP1-AS2 overexpression, PAXIP1 or CYLD knockdown each reduced the levels of POLH and CYLD knockdown also reduced the level of RAD18, causing a significant decrease in TLS, despite little change in the level of mUb-PCNA. This decreased TLS was likely caused by the decreased level of POLH, which performs actual TLS reactions, with possible participation of RAD18, which in addition to the monoubiquitination of PCNA has a non-catalytic function in TLS, involving an interaction with POLH and guiding it to replication stalling sites [25,36,37]. The phenomenon, in which TLS was reduced but mUb-PCNA remained essentially unchanged, indicates that the formation of mUb-PCNA cannot be used as a surrogate for TLS, as is sometimes done and may lead to erroneous conclusions.

A limitation of this study is that TENT4A expression was reduced using only siRNA or shRNA knockdown. Future studies using CRISPR knockout of TENT4A are expected to be instrumental in further strengthening the effects caused by depletion of TENT4A. Another limitation is the lack of experiments demonstrating that the expression of TENT4A can complement the deficiencies caused by TENT4A knockdown. Such experiments were attempted, but were unsuccessful due to difficulties in ectopic expression of the TENT4A protein. Future research in improving expression will help to overcome this difficulty.

TLS is involved in various biological processes, most notably DNA damage tolerance [9] and the generation of somatic hypermutation in the immune system [38]. Dys-regulation of TLS can cause an altered load of point mutations. If TLS is inhibited, it can increase chromosomal aberrations due to the activity of alternative recombinational and
repair events [39]. Of the 33 cancer types examined for the occurrence of mutations in TLS-related genes, endometrial cancer is the only one that exhibits TLS mutations in a considerable fraction of samples, spread over all 14 genes, including TENT4A (Table 1). This suggests that dysregulated or downregulated TLS is involved in endometrial carcinogenesis [40,41]. It is yet unclear why this should be specific to endometrial cancer; however, dysregulation or inactivation of discrete DNA repair mechanisms is generally cancer type-specific, for reasons that are not fully understood [1–6]. Of note, the loss of TLS might sensitize cancer cells to DNA damaging chemotherapy, providing potential prognostic and therapeutic value [42,43]. To the best of our knowledge, this is the first report on such extensive somatic mutations in genes of the TLS pathway in sporadic cancers.

4. Materials and Methods

4.1. Cell Cultures and Transfections

Osteosarcoma U2OS, breast cancer MCF-7, human embryonic kidney HEK 293T and 293FT epithelial cells were cultured in DMEM (Gibco) supplemented with 2 mM L-Alanyl-L-Glutamine, 100 units/mL of penicillin, 100 µg/mL of streptomycin, 1 mM sodium pyruvate (Biological Industries, Beit Haemek, Israel) and 10% fetal bovine serum (HyClone). XPA (XP12RO) human fibroblasts derived from xeroderma pigmentosum patients were a gift from A. R. Lehmann (University of Sussex, Brighton, U.K.). XP12RO cells were cultured in MEM Eagle (Biological Industries, Beit Haemek, Israel) supplemented with 2 mM L-Alanyl-L-Glutamine, 100 units/mL of penicillin, 100 µg/mL of streptomycin and 15% fetal bovine serum (HyClone, Thermo Fisher Scientific, MA, USA). The cells were incubated at 37 °C in a humidified incubator with 5% atmosphere and periodically examined for mycoplasma contaminations by EZ-PCR test kit (Biological Industries, Beit Haemek, Israel). For plasmid transfection, cells were transfected with Lipofectamine 2000 (Invitrogen Life Technologies, Carlsbad, CA, USA) or JetPRIME reagent (Polyplus-transfection S.A, Illkirch, France) according to the manufacturer’s instructions. For siRNA transfection, the siGENOME SMART pool siRNA oligonucleotides, siGENOME Non-Targeting Control siRNA #5 (D-001210–05) and ON-TARGETplus SMART pool siRNA oligonucleotides and ON-TARGETplus NON-targeting pool (Dharmacon, Lafayette, CO, USA) were transfected with 25 or 50 nM siRNA by Lipofectamine RNAimax (Invitrogen Life Technologies, Carlsbad, CA, USA) for 48 to 72 h following the manufacturer’s protocol. For combinatorial knockdown, an equal amount of siRNAs against each gene was mixed to have a final combined concentration of 100 nM for the double knockdown and 150 nM for the triple knockdown, unless otherwise stated. siRNAs used in this study are listed in Supplementary Materials Table S12.

4.2. Generation of Lentiviral Stable Cell Lines

Lentiviral pLKO.1-puro empty vector control plasmid and human TENT4A shRNA pLKO.1-puro plasmid (clone TRCN0000053036, target sequence CCAACAATCAGACCAGGTTTA) were obtained from Sigma (Mission shRNA library). Lentiviruses were produced in 293FT cells by co-transfecting pLKO-derived plasmids and the second-generation packaging plasmids using JetPRIME. Lentiviral particles were harvested 48 h post-transfection. The resulting lentiviral particles were used to infect the U2OS cells. Exactly 48 h post-infection, 2 µg/mL of puromycin (InvivoGen, San Diego, CA, USA) was added to select for infected cells. After two weeks of selection, individual colonies were isolated and tested for knockdown of TENT4A by quantitative real-time PCR (qPCR).

4.3. TLS Assay in Cultured Mammalian Cells

The TLS assay was described earlier [24,44,45]. The TLS gap-plasmid transfection was performed by co-transfecting with a mixture containing 50 ng of a lesions plasmid (KanR), 50 ng of a gapped plasmid without a lesion (CmR) and 1900 ng of the carrier plasmid pUC18, using Lipofectamine 2000. The cells were incubated for 4 h (for the TTCPD gap-lesion plasmid), 20 h (TT 6-4PP), 16 h (BP-G), or 6 h (cisPt-GG) to allow TLS. The plasmids
were extracted using alkaline lysis conditions followed by renaturation, such that only covalently closed plasmids remained non-denatured. A fraction of the purified DNA was used to transform a TLS-defective *E. coli* recA strain JM109, which was then plated on LB-kan and LB-cm plates. The efficiency of gap repair was calculated by dividing the number of transformants obtained from the gap-lesion plasmid (number of colonies on LB-kan plates) by the number of corresponding transformants obtained with the control gapped plasmid GP20-cm (number of colonies on LB-cm plates). To obtain precise TLS extents, the plasmid repair extents were multiplied by the fraction of TLS events out of all plasmid repair events, based on the DNA sequence analysis of the plasmids from KanR colonies. To determine the DNA sequence changes that have occurred during plasmid repair, sequence analysis was carried using the TempliPhi DNA Sequencing Template Amplification Kit (GE Healthcare Life Sciences, Piscataway, NJ, USA) and the BigDye Terminator v1.1 Cycle Sequencing Kit (Applied Biosystems, Foster City, CA, USA). Reactions were analyzed by capillary electrophoresis on an ABI 3130XL Genetic Analyzer from Applied Biosystems.

4.4. RNA Isolation and qPCR

Total RNA was isolated using RNeasy plus mini kit (Qiagen, Hilden, Germany) according to the manufacturer’s instructions, including treatment with RNase-free DNase I (Qiagen, Hilden, Germany). cDNA was synthesized using a High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Foster City, CA, USA). qPCR was performed using qPCRBIO SyGreen Blue Mix (PCR Biosystems, London, UK) and run on a QuantStudio™ 6 Flex Real-Time PCR System (Applied Biosystems). Primer sequences used were pre-designed KiCqStart SYBR® Green primers (Sigma-Aldrich Israel Ltd, Rehovot, Israel) and are included in Supplementary Materials Table S13. The expression of the indicated transcript was normalized to endogenous reference control GAPDH according to the ∆∆Ct method using the DataAssist software v3.0 (Applied Biosystems).

4.5. mRNA Stability Assay

The expression of TENT4A, TENT4B, PAXIP1 or TENT4A plus TENT4B, or TENT4A plus PAXIP1 was knocked down for 48 h in MCF-7 cells. To measure mRNA stability, 5 µg/mL Actinomycin D (Sigma-Aldrich, St Louis, MO, USA) was added to the growth medium to inhibit transcription. Cells were harvested at the indicated time points and mRNA expression was measured by qPCR. The half-life of the mRNA was calculated by the non-linear fit, one-phase decay curve equation using GraphPad Prism 8 software (San Diego, CA, USA).

4.6. RNA-Immunoprecipitation (RNA-IP) and Extension Poly(A) Test (eP AT)

RNA-IP to assay TENT4A/RNA interactions was performed as described [46] and the ePAT assay was performed as described with some modifications [26]. The two are presented in the Supplementary Materials and Methods and Supplementary Materials Tables S13 and S14.

4.7. UV-Irradiation, Whole-Cell Extracts and Western Blotting

When indicated, cells were rinsed in Hanks’ buffer and irradiated in Hanks’ buffer with UV-C using a low-pressure mercury lamp (TUV 15W G15T8, Philips) at a dose rate of 0.2 J/m²/s. The UV dose rate was measured using a UVX Radiometer (UVP) equipped with a 254-nm detector. After irradiation, Hanks’ buffer was removed, and the cells were incubated in fresh growth medium for additional time before harvest.

Cells were resuspended in CelLytic M lysis buffer (Sigma-Aldrich) containing protease inhibitor cocktail, phosphatase inhibitor 2 and 3 cocktail (Sigma-Aldrich), 2.5 mM MgCl₂ and 50 units/mL Benzonase (Merck, Darmstadt, Germany), incubated on ice for 30 min followed by centrifugation. The supernatant was collected and the protein concentration was determined using the Bradford protein assay (Bio-Rad, Hercules, CA, USA). The extracts were fractionated on a 4–20% ExpressPlus™ PAGE gel (Genscript, Piscataway, NJ, USA) using SDS-MOPS
buffer. They were transferred onto nitrocellulose membrane (BioTrace™ NT Nitrocellulose transfer membrane, Pall Corporation), followed by blocking of membranes with Odyssey Blocking buffer (LI-COR Biosciences, Lincoln, NE, USA) for 1 h at room temperature. The membranes were incubated with primary antibodies diluted in Odyssey Blocking Buffer with 0.1% Tween-20 overnight at 4 °C. The membranes were washed 3 × 10 min with Tris-buffered saline with 0.1% Tween-20 (TBST) and further incubated for 1 h at room temperature in the IRDYE-680 or 800 conjugated secondary antibodies diluted in Odyssey Blocking Buffer with 0.1% Tween-20. Membranes were washed 3 × 10 min TBST and finally rinsed with TBS to remove residual Tween-20. The membranes were imaged on a LI-COR Odyssey Fc Imager (LI-COR Biosciences, Lincoln, NE, USA) and the bands were quantified by ImageStudio v 5.2.5 (LI-COR Biosciences) software.

4.8. Antibodies

Commercially available antibodies used were as follows: rabbit anti-RAD18 (Cell Signaling, Danvers, MA, USA, 9040S, dilution 1:2000); rabbit anti-ubiquityl-PCNA (Lys164) (Cell Signaling, 13439S, dilution 1:2000); rabbit anti-GAPDH (Cell Signaling, 5174S, dilution 1:10000); rabbit anti-POLH (Cell Signaling, 13848S, dilution 1:1000); rabbit anti-CYL1D (Cell Signaling, 8462S, dilution 1:1000); mouse anti-GAPDH (EMD Millipore Corporation, Temecula, CA, USA, MAB374, dilution 1:10000); mouse anti-PCNA (PC-10) (Santa Cruz Biotechnology, Santa Cruz, CA, USA, sc-56, dilution 1:1000); rabbit anti-TRF4 (H-172) (TENT4A) (Santa Cruz Biotechnology, sc-98490, dilution 1:500); mouse anti-FLAG M2 (Sigma-Aldrich, F1804, dilution 1:1000); rabbit anti-PTIP (PAXIP1) (Bethyl Laboratories, Montgomery, TX, USA, A300–370A, dilution 1:2000); rabbit anti-DTL (Bethyl Laboratories, A300–948A, dilution 1:1000); anti-MBP monoclonal antibody, HRP conjugated (New England Biolabs, Ipswich, MA, USA, E80385, dilution 1:10000); IRRDye 680RD goat anti-mouse IgG (LI-COR Biosciences, 926–68070, dilution 1:10000); IRRDye 800CW goat anti-mouse IgG (LI-COR Biosciences, 926–32210, dilution 1:10000); IRRDye 680RD goat anti-rabbit IgG (LI-COR Biosciences, 926–68071, dilution 1:10000); IRRDye 800CW goat anti-rabbit IgG (LI-COR Biosciences, 926–32211, dilution 1:10000).

4.9. RNA-Seq and Bioinformatics

The sources of RNA were U2OS cells in which the expression of TENT4A was knocked down using lentivirus-mediated shRNA combined with siRNA against TENT4A and human XP12RO cells transfected with 25 nM of siTENT4A and siControl by Lipofectamine RNAiMax for 48 h. RNA was extracted from three biological replicates, the quality assessed on the 2200 TapeStation system (Agilent Technologies, Santa Clara, CA, USA). Library preparation and sequencing were performed at the Crown Genomics Institute of the Nancy and Stephen Grand Israel National Center for Personalized Medicine at the Weizmann Institute of Science. Briefly, sequencing libraries were prepared using TruSeq® Stranded Total RNA (Illumina) (Cat # RS122–2301, RS-122–2302) for ribosomal-depletion libraries. After rRNA depletion from 500 ng total RNA with rRNA Ribo-Zero Gold removal mix, cDNA was prepared, followed by second-strand synthesis with dUTP instead of dTTP. Then, A base addition, adapter ligation, UDG treatment and PCR amplification steps were performed. Sequencing was performed on the Illumina HiSeq2500 machine. The median sequencing depth was 45 million reads. Poly-A/T stretches and Illumina adapters were trimmed from the reads using Cutadapt; resulting reads shorter than 30bp were discarded. Reads were mapped to the Homo Sapiens GRCh38 reference genome using STAR (version 2.4.2a), supplied with gene annotations downloaded from Ensembl (release 88). The alignEndsType was set to EndToEnd and outFilterMismatchNoverLmax was set to 0.04. Expression levels for each gene were quantified using htseq-count (the stranded option was set to reverse), using the gtf above. Differential expression analysis was performed using DESeq2 (version 1.10.1). The betaPrior, Cook’s distance cutoff and independent filtering parameters were set to False. Raw p values were adjusted for multiple testing using the procedure of Benjamini and Hochberg [47]. We defined significantly differentially
expressed genes in U2OS and XP12RO cells as those with FDR $\leq 0.05$, absolute fold change $\geq 2$ and a count of at least 30 at least in one of the samples.

4.10. Gene Set Enrichment Analysis (GSEA)

We performed GSEA using a pre-ranked algorithm with rank determined by $-\log_{10}(\text{FDR q-value}) \times \text{sign (fold change)}$. The Java GSEA Desktop Application version 3 from the Broad Institute was used and the enrichment statistic parameter set to classic but other parameters remained at their default values. Enrichment analysis was performed, and leading-edge gene sets were determined by using GO processes (c5.bp.v6.1.symbols.gmt) from Molecular Signature Database (MSigDB) v6.1. Gene sets were considered significantly enriched at FDR < 0.05. We further performed enrichment analysis in Pathway Studio MammalPlus V 12 (Elsevier B.V, Amsterdam, The Netherlands) by GSEA and Fisher’s Exact Test for genes of U2OS and XP12RO.

4.11. TCGA Database Mutation Analysis

The R package ‘TCGAbiolinks’ [48] was used to download all the MAF (Mutation Annotation Format) files related to the 33 cancer types from TCGA database. Loading and summarizing the MAF files for each cancer was done using the R package ‘maftools’ [49]. For the analysis of each cancer type, MAF files from all four pipeline analysis available in the TCGA database (“muse”, “varscan2”, “somaticsniper”, “mutect2”) were merged using the ‘merge_mafs’ function from ‘maftools’. The variant classifications, “Frame_Shift_Del”, “Frame_Shift_Ins”, “Splice_Site”, “Translation_Start_Site”, “Nonsense_Mutation”, “Non-stop_Mutation”, “In_Frame_Del”, “In_Frame_Ins”, “Missense_Mutation”, “3’UTR”, “5’UTR”, “3’Flank”, “Targeted_Region”, “Silent”, “Intron”, “RNA”, “IGR”, “Splice_Region”, “5’Flank”, “lincRNA”, “De_novo_Start_InFrame”, “De_novo_Start_OutOfFrame”, “Start_Codon_Ins”, “Start_Codon_SNP”, “Stop_Codon_Del”, were used in the analysis of mutations. Mutations frequencies were calculated as the number of mutated samples divided by the total number of samples for each gene per cancer.

4.12. Statistical Analysis

The statistical results were obtained from at least three independent biological replicates unless otherwise stated. All results were presented as mean ± SEM. P values were obtained via the Student’s $t$-test (two-tailed), unless otherwise stated, using GraphPad Prism 8.0 software. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/ijms22136957/s1, Supplementary Materials and Methods, Figure S1: Knockdown of TENT4A and/or TENT4B in MCF-7 cells, Figure S2: Partially purified 8xHis-MBP-tagged TENT4A (Wt) and mutant 8xHis-MBP-tagged TENT4A DD277,279AA (Mut), Figure S3: ePAT analysis of the poly(A) tail of GAPDH mRNA, Figure S4: Effect of TENT4A knockdown on the stability of RNA18S RNA, Figure S5: Distribution of the fraction of samples with mutations in sets of random 14 genes., Table S1: TLS across a site-specific TT 6-4PP or BP-G in TENT4A knocked down U2OS cells, Table S2: Analysis of mutations formed during TLS across a TT 6-4PP in TENT4A knocked down U2OS cells, Table S3: Analysis of mutations formed during TLS across a TT 6-4PP in TENT4A knocked down U2OS cells, Table S4: TLS across a site-specific cisPt-GG or TT-CPD lesion in TENT4A and/or TENT4B knocked down MCF-7 cells, Table S5: Analysis of mutations formed during TLS across a site-specific cisPt-GG lesion in TENT4A or/and TENT4B knocked down MCF-7 cells, Table S6: Analysis of mutations formed during TLS across a site-specific TT-CPD lesion in MCF-7 cells in which TENT4A expression was knocked down, Table S7: TLS across a site-specific cisPt-GG lesion in MCF-7 cells overexpressing PAXIP1-A52 long non-coding antisense RNA, Table S8: Analysis of mutations formed during TLS across a site-specific cisPt-GG lesion in MCF-7 cells overexpressing PAXIP1-A52 long non-coding antisense RNA, Table S9: TLS across a site-specific TT-CPD lesion in MCF-7 cells in which PAXIP1 expression was knocked down, Table S10: Analysis of mutations formed during TLS across a site-specific TT-CPD lesion in MCF-7 cells in which PAXIP1 expression was knocked down, Table S11: Number of samples with mutations in TLS-related vs random genes, Table S12: siRNA used for the knockdown studies, Table S13: List of qPCR primers used in this study, Table S14: List of primers used in the ePAT assay.
Author Contributions: U.S. (Umakanta Swain) designed, performed, and analyzed most experiments and participated in writing the manuscript. G.F. performed the RNAseq analysis and commented on the manuscript. U.S. (Umakanta Sehrawat) performed polysome profiling experiments, analyzed their data and commented on the manuscript. A.S.-P. and R.R. performed the mutational analysis based on the TCGA database. C.E., T.C. and N.E.G. provided oligonucleotides and commented on the manuscript. T.P.-E., participated in designing the experiments, analysis of the data, and writing the manuscript. R.D. provided advice, contributed to the analysis of the results, and commented on the manuscript. Z.L. conceived, devised, and supervised the study and wrote the manuscript. All authors have read and agreed to the published version of the manuscript.

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