

Stop pulling my strings — what telomeres taught us about the DNA damage response

Eros Lazzerini-Denchi¹ and Agnel Sfeir²

Abstract | Mammalian cells have evolved specialized mechanisms to sense and repair double-strand breaks (DSBs) to maintain genomic stability. However, in certain cases, the activity of these pathways can lead to aberrant DNA repair, genomic instability and tumorigenesis. One such case is DNA repair at the natural ends of linear chromosomes, known as telomeres, which can lead to chromosome-end fusions. Here, we review data obtained over the past decade and discuss the mechanisms that protect mammalian chromosome ends from the DNA damage response. We also discuss how telomere research has helped to uncover key steps in DSB repair. Last, we summarize how dysfunctional telomeres and the ensuing genomic instability drive the progression of cancer.

DNA damage response (DDR). A collection of pathways that sense, signal and repair DNA lesions.

Linear chromosomes pose a challenge to eukaryotic cells. This problem was first recognized by Barbara McClintock and Herman Muller^{1,2}, who postulated that specialized structures named ‘telomeres’ distinguish chromosome ends from sites of DNA double-strand breaks (DSBs). Since then, extensive research has revealed the composition, structure and function of telomeres (FIG. 1). Mammalian telomeres consist of arrays of TTAGGG repeats that range from 5 kb in human cells to 100 kb in mice, which are polymerized by telomerase, a specialized reverse transcriptase³. Telomeres end with a single-stranded G-rich overhang^{4,5} that can invade the preceding double-stranded region to generate a special lariat-like structure called the telomere loop or t-loop^{6,7}. Telomere DNA is transcribed by RNA polymerase II into a long non-coding telomeric repeat-containing RNA (TERRA)⁸. The function of TERRA is not fully understood, but the emerging view is that it functions as a molecular scaffold for proteins that assist in proper telomere function (for a review, see REF. 9).

Telomeres are bound by shelterin, a six-subunit protein complex that protects chromosome ends from aberrant activation of the DNA damage response (DDR)¹⁰ (FIG. 1). Shelterin recognizes TTAGGG repeats through the binding of its TRF1 (telomere repeat-binding factor 1; also known as TERF1)¹¹ and TRF2 (REFS 12, 13) subunits to duplex DNA. TRF1 and TRF2 co-interact with TIN2 (TRF1-interacting nuclear factor 2), which in turn binds the TPP1 (PTOP, PIP1 or TINT1)–POT1 (protection of telomere 1) heterodimer^{14–18}. POT1 is the third DNA-binding component within shelterin.

It is recruited to telomeres by interacting with TPP1 and coats the single-stranded part of the TTAGGG repeats with its oligonucleotide/oligosaccharide binding folds^{19,20}. Rodents express two POT1 paralogues — POT1A and POT1B — that are structurally similar yet functionally divergent^{21,22}. RAP1 (repressor activator protein 1) is the sixth and most conserved shelterin component; it is recruited to telomeres by interacting with TRF2 (REFS 23–25). The current view is that shelterin can form as a six-subunit complex as well as subcomplexes lacking TRF1 or TRF2–RAP1 (REFS 14, 15, 18, 26). The telomere proteome comprises additional telomere-associated proteins^{27–30}, including DNA damage factors (Ku, MRN (MRE11–RAD50–NBS1)), nucleases (structure-specific endonuclease subunit SLX4, Apollo), helicases (Bloom syndrome, RecQ helicase-like (BLM), Werner syndrome, RecQ helicase-like (WRN), regulator of telomere elongation helicase 1 (RTEL1)) and chromatin modifiers (α -thalassaemia/mental retardation syndrome X-linked (ATRAX)). A key complex that is central for telomere function is the trimeric CST complex, which is composed of the DNA polymerase α (Pol α)–primase accessory factors CTC1, STN1 and TEN1 (REFS 31, 32). Interestingly, recent data suggest that in mouse germ cells, a meiosis-specific telomere complex, composed of TERB1 (telomere repeats-binding bouquet formation protein 1), TERB2 and membrane-anchored junction protein (MAJIN), replaces the shelterin complex to facilitate the attachment of telomeres to the inner nuclear membrane^{33,34}.

¹Department of Molecular and Experimental Medicine, Scripps Research Institute, La Jolla, California 92037, USA.

²Skirball Institute of Biomolecular Medicine, Department of Cell Biology, NYU School of Medicine, New York, New York 10016, USA.

edenchi@scripps.edu; agnel.sfeir@med.nyu.edu

doi:10.1038/nrm.2016.43
Published online 11 May 2016

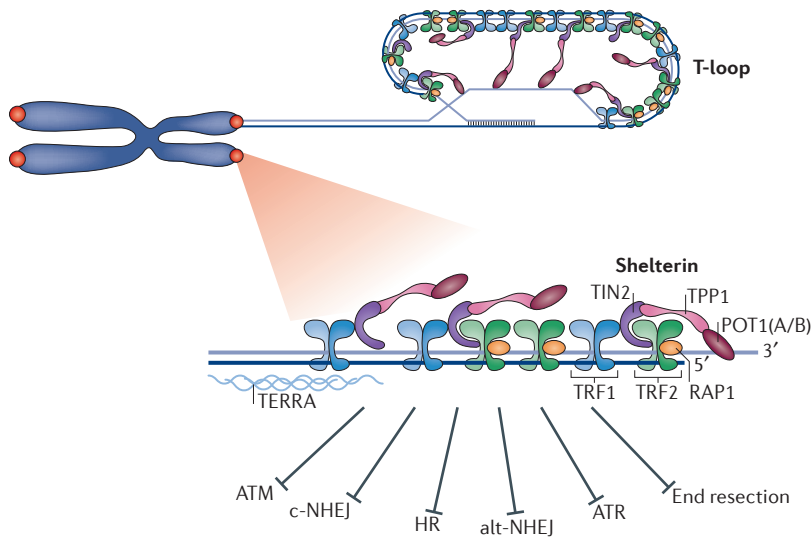


Figure 1 | Overview of telomere composition and function. Mammalian telomeres are composed of long stretches of TTAGGG repeats that range from 5 kb in human cells to 100 kb in mice and end with a single-stranded 3' overhang of up to a few hundred nucleotides in length^{4,5}. Telomeric DNA is bound by the specialized shelterin complex, transcribed into a long non-coding telomeric repeat-containing RNA (TERRA) and packaged into a t-loop (telomere loop) configuration. Shelterin subunits include TRF1 (telomere repeat-binding factor 1), TRF2, TIN2 (TRF1-interaction factor 2), RAP1 (repressor activator protein 1), TPP1 and POT1 (protection of telomere 1; POT1A and POT1B in mice). The six-subunit complex protects chromosome ends from DNA damage signalling by ATM (ataxia telangiectasia mutated) and ATR (ataxia telangiectasia and Rad3-related), and from DNA repair by c-NHEJ (classical non-homologous end joining), alt-NHEJ (alternative non-homologous end joining), HR (homologous recombination) and DNA end resection.

When shelterin function is compromised, the outcome is rapid telomere deprotection, activation of the DDR leading to cellular death (apoptosis) or irreversible cell cycle arrest (senescence), and, in certain cases, induction of genomic instability. Shelterin function is lost in cells with critically short telomeres, as chromosome ends in these cells lack sufficient binding sites for this protective complex. In other cases, loss of function of shelterin is caused by genetic alterations in components of the complex, leading to alteration in binding and/or function. In this Review, we discuss recent discoveries that have shed light on how chromosome ends are protected by the shelterin proteins to avoid genomic instability, and on how telomere deprotection has been used to identify novel DSB repair factors.

Peeling back layers of end protection

Cells detect DNA breaks with the help of two surveillance pathways, driven by the signalling kinases ATM (ataxia telangiectasia mutated) and ATR (ataxia telangiectasia and Rad3-related), which are activated in response to the formation of DSBs and single-strand breaks, respectively (for a review, see REF. 35). DNA damage signalling triggers cell cycle arrest, which allows cells to repair the breaks, if possible, or to undergo senescence or apoptosis. Repair of DSBs involves either the error-free pathway of homologous recombination, or one of the two error-prone, non-homologous end joining (NHEJ) pathways, classical NHEJ (c-NHEJ) or alternative NHEJ

(alt-NHEJ) (BOX 1). In addition, DSBs are subject to nucleolytic processing, which is a key step that can influence the choice of repair pathway. During the past two decades, genetic studies have helped to delineate how telomeres use shelterin to silence the DDR.

TRF2, the bouncer at the gate. DNA damage signalling by ATM is primarily repressed at telomeres by TRF2 (REFS 36,37). TRF2 inhibition activates ATM, which phosphorylates the downstream effectors Ser/Thr protein kinase CHK2 and p53 (REF. 24), thereby inducing the formation of telomere dysfunction-induced foci (TIFs). These are marked domains of telomere-associated DNA damage factors, including 53BP1 (p53-binding protein 1) and the histone variant H2AX³⁸. The underlying mechanism by which TRF2 inhibits ATM-dependent repair pathways is complex and not fully understood. The current view emphasizes two levels of control; the first involves the t-loop configuration, and the second consists of direct inhibition of the ATM signalling cascade (FIG. 2a). TRF2 binding to DNA *in vitro* stimulates strand invasion, forming structures that resemble t-loops³⁹. Furthermore, the frequency of t-loops *in vivo* is significantly reduced in cells lacking TRF2, implicating this shelterin subunit in their formation and/or stabilization⁷. When telomere ends are engaged in a t-loop configuration, they are unlikely to be detected by the MRN complex, which is essential for ATM activation. In addition, TRF2 inhibits ATM signalling directly by inhibiting the kinase itself³⁶, as well as crucial downstream effectors of the ATM pathway⁴⁰ (FIG. 2a). A motif within TRF2, termed the iDDR (inhibitor of the DDR pathway), inhibits the activity of the E3 ubiquitin ligase RNF168, thereby preventing the accumulation of 53BP1, which is a key effector of ATM⁴⁰. The two-step mechanism by which TRF2 inhibits ATM activation may be crucial for telomere protection during the S phase of the cell cycle. The progression of the replication fork through telomeric DNA triggers transient unwinding of t-loops and renders chromosome ends susceptible to the DDR. Direct inhibition of the ATM signalling pathway by TRF2 will therefore ensure end protection. Finally, biochemical analysis and atomic force microscopy has highlighted a topological mechanism of TRF2-mediated repression of ATM. Specifically, the wrapping of DNA around the TRFH domain of TRF2 was recently proposed to promote t-loop formation and inhibit ATM signalling^{39,41}.

The struggle to keep ends apart. The most deleterious outcome of telomere dysfunction is the formation of chromosome end-to-end fusions, resulting in dicentric chromosomes, which can lead to breakage–fusion–bridge cycles and induce extensive chromosomal instability. Mammalian cells have evolved sophisticated mechanisms to prevent such events from occurring. The major factor suppressing chromosome end-to-end fusions is TRF2. When telomeres are depleted of TRF2, they become substrates for c-NHEJ^{24,42}. Artificial tethering of TRF2 to non-telomere loci inhibits break repair, suggesting that TRF2 is both necessary and sufficient to suppress

Dicentric chromosomes
Aberrant chromosomes with two centromeres, resulting from the fusion of two chromosomes.

Breakage–fusion–bridge
A mechanism producing chromosomal instability, triggered by the fusion of deprotected telomeres, which leads to repeating cycles of chromosome breakage and fusion.

c-NHEJ⁴³. TRF2 forms a stable 1:1 complex with its interacting partner RAP1 (REFS 44,45), and RAP1 protein is rapidly destabilized upon deletion of *TRF2* (REF. 24). Interestingly, tethering of RAP1 to telomere DNA in cells lacking TRF2 was reported to reduce the frequency of telomere fusions⁴⁶. However, depletion of RAP1 from human and mouse telomeres is not sufficient to trigger c-NHEJ^{25,47,48}. A recent study provides a plausible explanation for the contrasting data by suggesting that RAP1 provides a redundant mechanism to block c-NHEJ when the function of TRF2 is partially compromised⁴¹.

Paradoxically, components of the c-NHEJ pathway, most notably the Ku70–Ku80 complex, are constitutively present at telomeres⁴⁹. Ku has a crucial role in protecting telomeres in human cells, and its depletion leads to rapid deletion of telomeric repeats⁵⁰. By contrast, deleting Ku in mice does not trigger major telomeric defects^{51,52}, hinting at alternative solutions to achieve telomere protection in rodents. Nevertheless, the strong impact of Ku depletion on telomere stability in human cells raises the obvious question of how TRF2 is able to disengage Ku without compromising telomere stability (FIG. 2a). A possible mechanism invokes a recently described interaction between TRF2 and the $\alpha 5$ region of Ku70, which prevents Ku70–Ku80 heterotetramerization⁵³. The TRF2–Ku interaction is thought to block the ability of Ku70–Ku80 to tether opposite DNA ends, which could explain why Ku70–Ku80 association with functional telomeres does not unleash c-NHEJ.

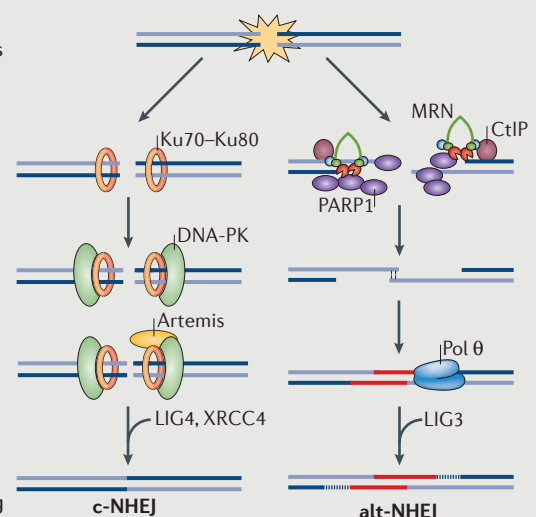
Intriguingly, recent reports suggest that during mitosis, telomeres are highly susceptible to c-NHEJ-mediated fusion^{54–56}. Mitotic cells attenuate c-NHEJ globally by preventing the phosphorylation of RNF8 and 53BP1 by mitotic kinases such as Polo-like kinase 1 (PLK1) and cyclin-dependent kinase 1 (CDK1)⁵⁵.

When this regulation is bypassed, telomeres are fused in an Aurora kinase B-dependent manner. These data corroborate a previous report showing that prolonged mitotic arrest leads to telomere uncapping following the eviction of TRF2 from telomeres, in a process that is dependent on Aurora kinase B⁵⁴. Future work is likely to shed light on telomere protection in mitosis and reveal why mammalian cells opt for a global shutdown of c-NHEJ, as opposed to simply configuring an extra layer of protection at telomeres.

Blocking ATR signalling. The activity of ATR at telomeres is primarily repressed by POT1. Deleting POT1, or blocking its recruitment to telomeres by inhibiting TPP1 or TIN2, induces the formation of ATR-dependent TIFs^{37,57–60}. POT1 binds to telomeric single-stranded DNA (ssDNA), thus preventing the recruitment of RPA (replication protein A), which is a crucial factor for the activation of ATR⁶¹ (FIG. 2b). Although POT1 affinity for ssDNA does not exceed the binding affinity of RPA, it has been proposed that the increased local concentration of POT1 at telomeres excludes RPA binding^{59,60}. An alternative model for POT1-mediated inhibition of ATR invokes a cell cycle-regulated RPA-to-POT1 switch mediated by hnRNPA (heterogeneous nuclear ribonucleoprotein A) and TERRA⁶². According to this model, accumulation of hnRNPA at replicated telomeres promotes the displacement of RPA and the subsequent recruitment of POT1. ATR activation in S phase is also inhibited, by TRF1, which is dedicated to counteracting replication defects at telomeres⁶³ (FIG. 2c). Notably, tethering of TPP1–POT1 to telomeres lacking TRF1 is sufficient to inhibit the TIF response, suggesting that ATR inhibition by TRF1 is dependent on the recruitment of TPP1 and POT1 to telomere DNA⁶⁴.

Box 1 | Classical non-homologous end joining (c-NHEJ) versus alternative NHEJ (alt-NHEJ)

Cells use two mechanistically distinct end-joining pathways to repair DNA double-strand breaks (DSBs)^{194,195}. C-NHEJ leads to minimal sequence alterations at the repair junctions, whereas alt-NHEJ (also known as microhomology-mediated end joining (MMEJ)) causes extensive deletions (as well as insertions) that scar the break sites following repair (see the figure). C-NHEJ is active throughout the cell cycle and is initiated when the Ku70–Ku80 heterodimer binds to DNA ends with high affinity. Ku then recruits the Ser/Thr kinase DNA-PKs (DNA-dependent protein kinase catalytic subunit) to phosphorylate a number of downstream targets, including the terminal end-processing enzyme Artemis that cleaves single-stranded overhangs, and DNA ligase 4 (LIG4) and the scaffold protein XRCC4, which catalyse the ligation of DNA ends. Alt-NHEJ, which is most active in the S and G2 phases of the cell cycle, is dependent on signalling by poly(ADP-ribose) polymerase 1 (PARP1) and relies on 5'–3' resection of DNA by MRN (MRE11–RAD50–NBS1) and CtBP-interacting protein (CtIP). Base pairing at the resected ends drives their annealing to promote synapsis of opposite ends of a DSB. Annealed ends are subject to fill-in synthesis by the low-fidelity DNA polymerase θ (Pol θ), which stabilizes the annealed intermediates and promotes end joining, primarily by LIG3. Alt-NHEJ introduces deletions and insertions that scar the break sites following repair. The deletions are caused by extended nucleolytic processing, whereas the insertions result from the activity of Pol θ .



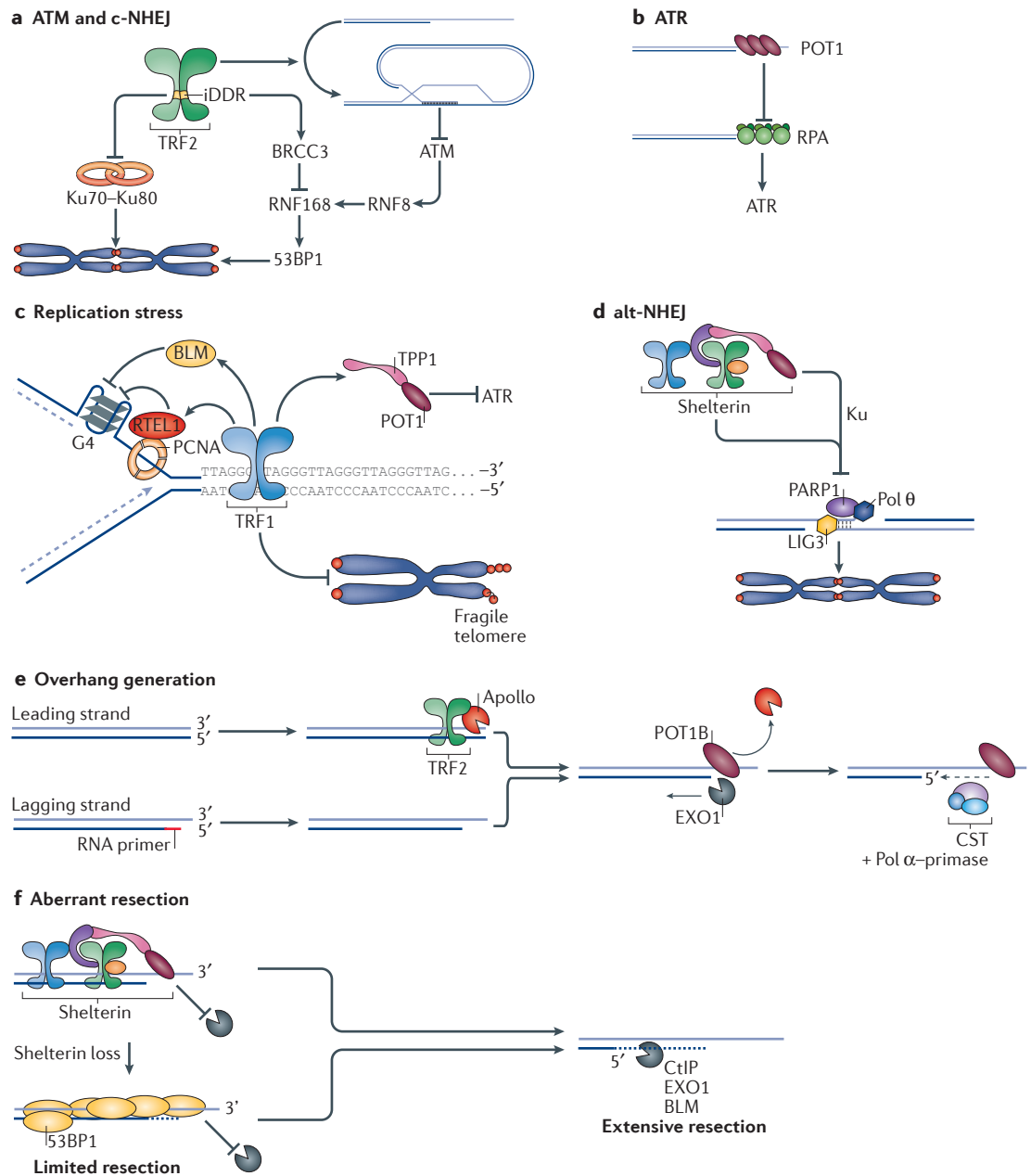


Figure 2 | How shelterin protects telomeres. **a** | TRF2 (telomere repeat-binding factor 2) represses ATM (ataxia telangiectasia mutated) signalling and classical non-homologous end joining (c-NHEJ). TRF2 promotes the formation of the protective telomere loop (t-loop) structure, which hides chromosome ends from ATM and c-NHEJ. In addition, TRF2 inhibits 53BP1 (p53-binding protein 1) accumulation by blocking RNF168-mediated ubiquitylation by activating the deubiquitylase BRCC3 (BRCA1/BRCA2-containing complex, subunit 3). Last, TRF2 blocks the dimerization of the Ku complex, thereby preventing the activation of c-NHEJ. **b** | POT1 (protection of telomere 1) represses ATR (ataxia telangiectasia and Rad3-related) signalling by competing with RPA (replication protein A) for single-stranded DNA (ssDNA) binding at telomeres. **c** | TRF1 inhibits ATR activity during telomere replication with the help of TPP1–POT1. TRF1 also counteracts replication fork stalling at telomeric secondary DNA structures (such as quadruplex DNA (G4)) with the help of RTEL1 (regulator of telomere elongation helicase 1) and BLM (Bloom syndrome, RecQ helicase-like), thereby protecting against telomere fragility. RTEL1 is recruited to replicating telomeres by interacting with PCNA (proliferating cell nuclear antigen). **d** | Alt-NHEJ (alternative-NHEJ), which is dependent on DNA ligase 3 (LIG3), PARP1 (poly(ADP-ribose) polymerase 1) and DNA polymerase θ (Pol θ), is repressed in a redundant manner by shelterin and the Ku70–Ku80 complex. **e** | The generation of telomere 3' overhang involves TRF2-dependent recruitment of the nuclease Apollo to resect double-stranded ends. Leading and lagging ends are then resected by EXO1 (exonuclease 1) to generate long single-stranded overhangs, which are subsequently filled in by Pol α -primase and the CST (CTC1–STN1–TEN1) complex. **f** | Aberrant resection of uncapped telomeres is carried out by the enzymatic machinery that processes double-strand breaks (DSBs) — the nucleases CtIP (CtBP-interacting protein) and EXO1 and the helicase BLM — and is repressed redundantly by shelterin and 53BP1. iDDR, inhibitor of the DNA damage response.

A joint effort to suppress alt-NHEJ at chromosome ends. Early evidence for the activation of the alt-NHEJ pathway at mammalian telomeres emerged from the analysis of telomerase-deficient mice. Specifically, chromosome end-to-end fusions following telomere attrition were evident even when core components of the c-NHEJ pathway (DNA ligase 4 and DNA-dependent protein kinase catalytic subunit (DNA-PKcs)) were deleted⁶⁵. These experiments hinted that alt-NHEJ could be responsible for processing dysfunctional telomeres in the early stages of tumorigenesis. Analysis of telomere fusion junctions in human tumours revealed hallmarks of alt-NHEJ repair — frequent microhomologies and extensive deletions^{66–68} — further implicating this error-prone repair pathway as operating at dysfunctional telomeres. Genetic manipulation of shelterin in mouse cells indicated that alt-NHEJ is repressed in a redundant manner^{69–71}. Specifically, ligase 3-mediated telomere fusions were maximally observed when the shelterin complex was completely depleted in Ku70–Ku80 deficient mouse cells⁶⁹ (FIG. 2d). The mechanism by which redundant suppression of alt-NHEJ is achieved has not been fully established. It is possible that the activity of alt-NHEJ is dependent on signalling by both ATM and ATR, which manifests when the entire shelterin complex is lost. In agreement with this idea, co-depletion of TRF2 and TPP1 — which activate the two kinases, respectively — is sufficient to trigger efficient alt-NHEJ activity⁷⁰. Similarly, the mechanism by which Ku inhibits alt-NHEJ remains unknown. It has been proposed that Ku has a higher binding affinity to DSBs than PARP1 (poly(ADP-ribose) polymerase 1) and could therefore block alt-NHEJ by repressing PARP1-mediated signalling^{72,73}. Alternatively, Ku might exert its effect by inhibiting 5' end resection^{74,75}, which is a prerequisite for alt-NHEJ-mediated repair⁷⁶.

Polishing the end: the art of making overhangs. Telomere ends are subject to two forms of nucleolytic processing, each of which is carried out by independent machineries and regulated differently (FIG. 2e). Following telomere replication, physiological processing of telomere ends generates a 3' overhang, a crucial structure for telomere protection. This is mediated by a number of factors, including the shelterin subunits TRF2 and POT1B (in mice)^{21,22,77,78}. TRF2 recruits the Apollo nuclease to resect blunt leading-strand ends and create short overhangs, whereas lagging-strand overhangs result from the removal of the RNA primer from the terminal Okazaki fragment^{78,79}. Subsequently, a long-range resection step is carried out by EXO1 (exonuclease 1), which acts on both leading and lagging strands and transiently elongates the overhang⁷⁷. Finally, overhang length is fine-tuned to an optimal length (~50–300 nucleotides⁸⁰) with the help of the CST complex, which, in the case of mouse telomeres, is recruited by POT1B to assist during fill-in synthesis⁷⁷. It is important to note that the genetic analysis of 3' overhang generation was primarily carried out in mouse cells, and whether human POT1 functions similarly to mouse POT1B remains to be determined.

In addition to the aforementioned physiological processing of telomeres, dysfunctional telomeres are subject

to aberrant degradation. Hyper-resection of uncapped telomeres is inhibited by shelterin and by 53BP1, which is a general repressor of DNA end resection at DSBs. Deletion of *TRF2* in 53BP1-null cells leads to an extended telomere overhang, mediated by ATM and dependent on the endonuclease CtIP (CtBP-interacting protein)⁸¹. A more substantial resection takes place following the deletion of both *TRF1* and *TRF2* and the creation of shelterin-free telomeres in 53BP1-deficient cells⁶⁹ (FIG. 2f). The unmitigated resection of shelterin-free telomeres is executed by CtIP and EXO1 and is aided by the helicase BLM⁶⁹.

How the replication machinery navigates TTAGGG repeats. TTAGGG repeats are prone to forming stable secondary structures (including quadruplex (G4) DNA)⁸² that challenge the replication machinery as it progresses through telomeric DNA. Among the various shelterin subunits, TRF1 has a major role in assisting the semi-conservative replication of telomeres^{63,83} (FIG. 2c), and its function is similar to that of the fission yeast orthologue, Taz1 (REF. 84). Deletion of TRF1 from mouse cells induces the formation of fragile telomeres⁶³, a phenomenon that is reminiscent of fragile sites and attributed to DNA replication defects. The protective function of TRF1 is achieved with the help of two helicases, RTEL1 and BLM, which unwind spurious secondary structures and allow faithful duplication of telomeres⁶³. The function of RTEL1 during telomere replication is mediated by an interaction with the replication clamp PCNA (proliferating cell nuclear antigen). Inhibiting the RTEL1–PCNA interaction increases the incidence of replication fork stalling and telomere fragility⁸⁵. BLM is recruited to telomeres by direct binding to TRF1 and suppresses telomere fragility^{64,86}. Studies have also implicated WRN, another RecQ helicase, in facilitating lagging-strand telomere synthesis^{87,88}, although WRN does not function in the same pathway as TRF1 (REF. 63). A parallel pathway that assists in the replication of telomeres is executed by CST. Inhibition of individual CST components compromises replication fork restart and leads to telomere fragility^{89,90}. The activity of CST is independent of TRF1 (REF. 89), and the complex functions by assisting Pol α -primase activity at telomeres. Last, TRF2 is also thought to facilitate telomere replication by relieving topological constraints that would otherwise hinder replication-fork progression⁹¹.

We currently know many of the players that assist telomere replication and counteract telomere fragility, but the dynamic interplay between the different factors assisting the replisome to progress through telomere DNA is unknown. In addition, the nature of the fragile aberrancy itself, and whether it is the result of altered packaging of the chromatin or actual DNA breaks and chromatin gaps, remains a mystery that in the future may be solved by super-resolution microscopy.

Homologous recombination at telomeres: keeping up with the neighbours. The activity of the homologous recombination pathway at telomeres may seem to be less harmful than that of NHEJ, but it can affect cellular

Fragile telomeres

Breaks or gaps at telomeres of metaphase chromosomes, caused by replication stress.

Fragile sites

Genomic regions that appear as gaps or breaks on metaphase chromosomes when DNA replication is partially inhibited.

survival when it alters telomere length. Homologous recombination at telomeres manifests in three major forms: exchange of sequence between sister (chromatid) telomeres (telomere sister chromatid exchange (T-SCE)), aberrant excision of t-loops (t-loop homologous recombination), and recombination that leads to alternative lengthening of telomeres (ALT).

T-SCE has detrimental consequences when an unequal exchange happens, in which case a daughter cell inherits a short telomere and suffers the deleterious impact of telomere uncapping. Loss-of-function analysis in mouse cells revealed that shelterin contributes to the repression of T-SCE together with the Ku complex, which is a general repressor of recombination in mammalian

cells⁹² (FIG. 3a). Depletion of TRF2, RAP1 or POT1 in the context of Ku70–Ku80 complex deficiency stimulates exchange of sequences between telomeres on sister chromatids^{69,93,94}. The mechanism by which these factors inhibit T-SCE is unknown. With regards to POT1, it is conceivable that its binding to telomere ssDNA counteracts the loading of homologous recombination factors — RPA and, subsequently, RAD51 — thereby inhibiting recombination. RAP1 and TRF2 possibly stabilize the telomeric double-stranded DNA (dsDNA)–ssDNA junction and block strand invasion during homologous recombination. Consistent with this idea, *in vitro* studies indicate that TRF2 has a greater preference for binding to junction sites when bound to RAP1 (REF. 95).

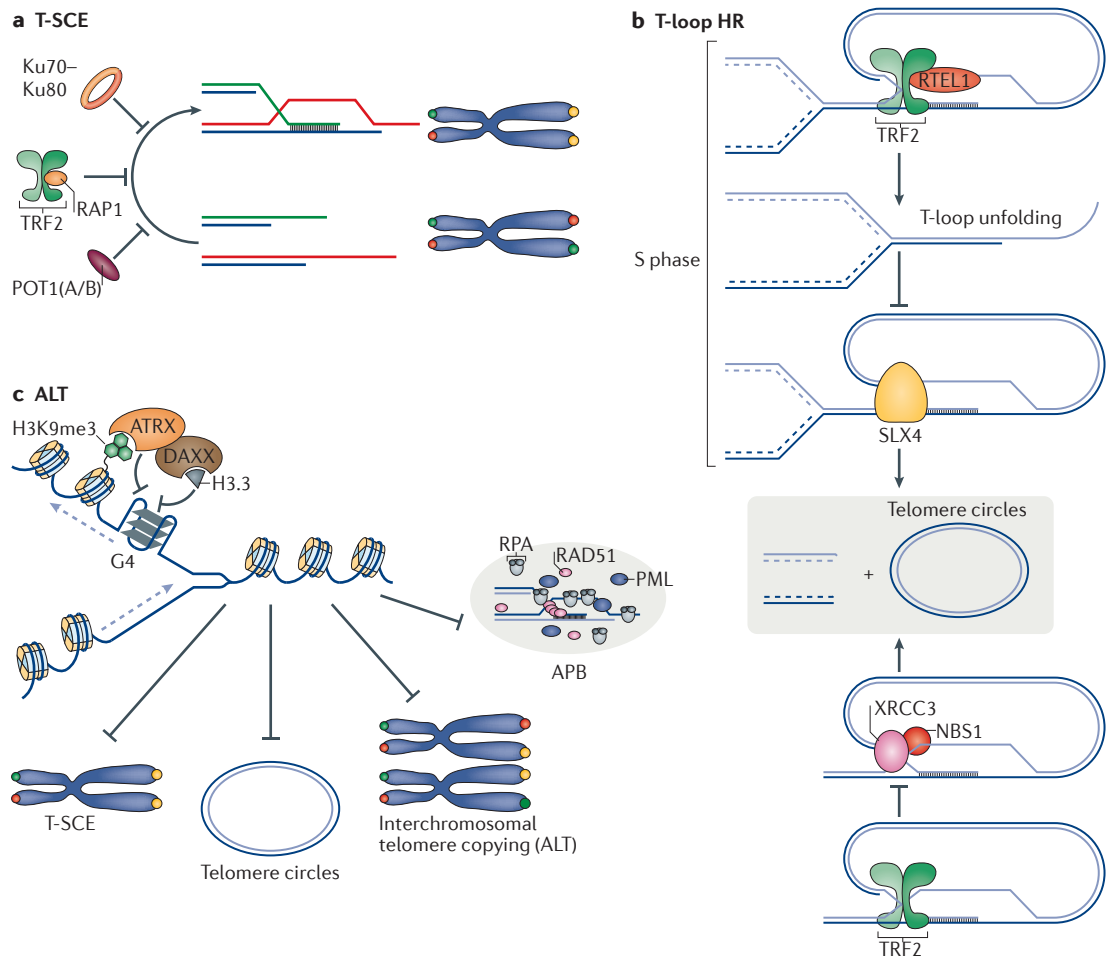


Figure 3 | The three facets of telomere homologous recombination: T-SCE (telomere sister chromatid exchange), t-loop (telomere loop) homologous recombination and ALT (alternative lengthening of telomeres). **a** | Exchange of sequence between sister chromatid telomeres (marked in red and green) is inhibited by RAP1 (repressor activator protein 1), POT1 (protection of telomere 1) and Ku70–Ku80. **b** | T-loop homologous recombination is blocked by TRF2 (telomere repeat-binding factor 2). TRF2 recruits RTEL1 (regulator of telomere elongation helicase 1) during S phase to unwind the t-loop and therefore protect it from being cleaved by structure-specific endonuclease subunit SLX4. In addition, TRF2 inhibits t-loop excision by inhibiting the activity of NBS1–XRCC3 (Nijmegen breakage syndrome 1–X-ray repair complementing defective repair in Chinese hamster cells 3). **c** | Telomere repeats have the propensity to form stable quadruplex (G4) DNA structures, which would impede replication fork progression. It has been proposed that ATRX (α -thalassaemia/mental retardation syndrome X-linked) unwind G4 DNA, enabling the deposition of histone H3.3 and ultimately assisting replication fork progression. The activity of ATRX at telomeres inhibits various ALT (alternative lengthening of telomeres) phenotypes including T-SCEs, formation of telomere circles, intrachromosomal telomere recombination and formation of APBs (ALT-associated promyelocytic leukaemia nuclear bodies). DAXX, death domain-associated protein; HR, homologous recombination; PML, promyelocytic leukaemia; RPA, replication protein A.

Aberrant telomere homologous recombination leads to t-loop excision owing to the activity of DNA repair protein XRCC3, NBS1 and SLX4 (FIG. 3b). The t-loop configuration poses a challenge to the replication machinery, as it needs to be unfolded for the replication fork to progress through telomeres, otherwise it could be excised. In addition to promoting t-loop formation, TRF2 protects t-loops from illegitimate homologous recombination. TRF2 recruits RTEL1 to unwind t-loops in S phase⁹⁶. Deleting RTEL1 or inhibiting its interaction with TRF2 allows SLX4-mediated t-loop excision, resulting in the formation of double-stranded telomere circles (t-circles) and rapid telomere loss^{96,97}. TRF2 also protects the t-loop from XRCC3- and NBS1-mediated cleavage^{98,99}. Notably, telomere trimming by XRCC3 occurs in normal cells that possess long telomeres, including in the male germ line^{100,101}. This mechanism has been proposed to provide an additional layer of telomere length regulation, although how it is kept in check to avoid rampant telomere shortening remains elusive.

ALT is activated in a subset of human tumours that lack telomerase activity¹⁰² to maintain the length of the telomere repeats¹⁰³. Telomeres maintained by ALT typically cluster in ALT-associated PML bodies (APBs)¹⁰⁴, have increased expression of TERRA¹⁰⁵ and display elevated levels of T-SCE¹⁰⁶. A hallmark of ALT is recombination between telomeres on separate chromosomes, which was demonstrated experimentally by interchromosomal copying of a telomere-embedded neomycin tag¹⁰³. ALT telomeres are highly heterogeneous in length¹⁰⁷ and are littered with non-canonical repeats, probably owing to recombination with subtelomeric regions¹⁰⁸. Survival of ALT cells is compromised when homologous recombination factors (RAD51, MRN, RAD9, RAD17, RPA and others) are inhibited, confirming their dependency on ALT for telomere maintenance (for a review, see REF. 109).

What triggers ALT and why it is activated in a subset of tumours remains unknown. A strong candidate is the histone chaperone ATRX, which is mutated in a large majority of cells and tumours that exploit the ALT pathway^{110,111}. Reintroducing ATRX into ALT cells suppresses T-SCE, APB and c-circle formation, and interchromosomal telomeric recombination^{112,113} (FIG. 3c). ATRX is part of the SWI/SNF family of ATP-dependent helicases¹¹⁴ and associates with chromatin by binding to sites of histone H3 Lys9 trimethylation (H3K9me3), which are enriched at telomeric DNA¹¹⁵. ATRX also interacts with the histone chaperone DAXX (death domain-associated protein), allowing the deposition of the histone variant H3.3 at repetitive sequences of telomeres and pericentromeres^{116,117}. The mechanism by which ATRX protects TTAGGG repeats from aberrant recombination remains unclear. However, several lines of evidence suggest that it may relate to telomere replication. First, *in vitro* studies indicate that ATRX binds to and unwinds G4 DNA¹¹⁸. Second, ATRX-deficient ALT cells accumulate increased levels of RPA at telomeres^{117,119}. Last, re-expression of ATRX in ALT cell lines reduces the frequency of replication fork stalling¹¹². Taken together, these studies suggest that both the helicase-unwinding

activity of ATRX and the histone chaperone properties of the ATRX–DAXX complex are likely to counteract telomere recombination by resolving stable secondary structures that would otherwise impede fork progression. It is important to emphasize that ATRX depletion by itself is not sufficient to induce ALT^{112,113}, indicating that additional genetic alteration(s) are necessary. Interestingly, deletion of the gene encoding the histone chaperone ASF1A is sufficient to trigger ALT-like phenotypes in telomerase-positive cells¹²⁰, and binding of the nucleosome-remodelling deacetylase (NuRD) complex to variant repeats found at telomeres in ALT cells creates a permissive environment for recombination¹²¹. Such observations reinforce the notion that alteration in chromatin status renders telomeres conducive to homologous recombination. In addition to chromatin environment, the presence of TERRA–telomeric DNA hybrids was proposed to affect ALT — reduced levels of these RNA–DNA hybrids leads to a significant reduction in homologous recombination-mediated telomere elongation¹⁰⁵.

In order for ALT telomeres to engage in interchromosomal recombination, they must first disengage from their cohered sister chromatid and move across the nucleus to meet a telomere on another chromosome. ATRX was proposed to regulate this choice between inter- and intratelomeric recombination. In the absence of ATRX, cohesion between sister telomeres persists, prompting an increase in T-SCE¹²². In addition, HOP2 (homologous-pairing protein 2 homologue), a protein that is normally required for synapsis of meiotic chromosomes, was recently shown to promote rapid and directional movement of telomeres over micrometre distances before their synapsis with recipient telomeres during ALT¹²³.

Using telomeres to discover DDR genes

The realization that inhibition of shelterin activity marks telomeres as sites of DSBs provides an opportunity to interrogate various aspects of the DDR using telomeres as an experimental system. As discussed above, removal of particular shelterin components activates specific DNA damage signalling pathways and repair mechanisms. Accordingly, shelterin manipulation provides a tractable system that has been used to gain insight into the mechanistic basis of DSB repair in mammalian cells.

Ingredients to make sticky ends. Different approaches have been used over the years to isolate factors that bind to functional as well as dysfunctional telomeres. A quantitative telomeric chromatin isolation protocol (QTIP) was applied to identify differences in telomeric chromatin composition between cells with functional telomeres and cells with telomeres depleted of TRF2 and POT1 (REF. 28). In a similar approach, proteomics of isolated chromatin segments (PICH)²⁷ was used to compare functional telomeres to those rendered dysfunctional following the removal of TRF2 (REF. 29). These approaches confirmed that dysfunctional telomeres are recognized as site of DNA damage and recruit the same repair factors that are associated with DSBs at other sites in the genome. Moreover, these methods and others helped

ALT-associated PML bodies (APBs). Promyelocytic leukaemia (PML) bodies are dynamic protein aggregates within the nuclei of some cells that contain the PML protein. Alternative lengthening of telomeres (ALT)-associated PML bodies are found exclusively in cancer cells, which rely on the ALT pathway to maintain telomeres.

Quantitative telomeric chromatin isolation protocol (QTIP). A telomere-protein purification method used to quantify changes in the content of telomeric chromatin.

Proteomics of isolated chromatin segments (PICH). A method to identify proteins associated with specific genomic loci that are rich in repetitive DNA.

to identify novel factors that contribute to the cellular response to telomere dysfunction. For instance, the Polycomb protein Ring1b was found to promote NHEJ at TRF2-depleted telomeres²⁹; a genome-wide RNA interference (RNAi) screen identified factors (such as RNF8) that mediate the response to telomere dysfunction¹²⁴; and isolation of TERRA-interacting proteins led to the identification of a Lys-specific demethylase (LSD1) as a factor that binds to dysfunctional telomeres and stimulates the nuclease activity of MRE11 (REF. 125).

Resecting DSBs — the way it's done at telomeres. DSB end resection is a crucial processing step that influences DSB repair pathway choice and can commit cells to repairing the break by homologous recombination. The underlying basis of end resection is a subject of intense investigation, and experiments carried out using dysfunctional telomeres have contributed to our understanding of key regulatory steps. 53BP1 is central to the process of DSB resection in mammalian cells. A seminal publication in 2010 reported that the loss of 53BP1 rescues embryonic lethality associated with loss of BRCA1 (breast cancer type 1 susceptibility protein)^{126,127}. This phenotype was due to the reactivation of homologous recombination, which was in turn attributed to the restoration of 5' end resection upon co-depletion of 53BP1 and BRCA1. DNA end resection at genome-wide DSBs is typically assessed by measuring the accumulation of RAD51 and phosphorylated RPA¹²⁶. A more accurate measurement can be achieved by analysing telomeres. In that respect, the role of 53BP1 in end resection was confirmed in a direct manner by analysing dysfunctional telomeres, in which the length of ssDNA can be accurately quantified using native gels (FIG. 4a). Specifically, depleting 53BP1 in TRF2-deficient cells blocks telomeres fusions by c-NHEJ and leads to increased end resection^{81,128}. The identification of 53BP1 as a key regulator for resection triggered a race to highlight downstream effectors, and the first to be identified was the 53BP1 partner, RIF1 (RAP1-interacting factor 1)¹²⁹. Loss of RIF1 in TRF2-null cells lead to an increase in overhang length at telomere termini, and epistasis analysis confirmed that RIF1 and 53BP1 function in the same pathway¹²⁹ (FIG. 4b). Interestingly loss of RIF1 in the context of shelterin depletion yielded less resection than 53BP1 loss, implying that RIF1 is unlikely to be the only downstream effector of 53BP1 (REF. 129). Subsequently, PAX-interacting protein 1 (PTIP) was identified as a second 53BP1-binding partner and a potential regulator of 5' end resection. Deleting PTIP, or inhibiting its binding to 53BP1, delays the appearance of fusions in TRF2-null cells¹³⁰, presumably owing to increased telomere end resection¹³⁰. Uncovering the role of RIF1 (REFS 129, 131–133) and PTIP¹³⁰ in DSB processing brings us a step closer to understanding the molecular basis of end resection, and further investigation will uncover downstream factors. The first hints were provided by an RNAi screen for genes that mediate the response to TRF2 depletion¹³⁴. This approach revealed REV7 (also known as MAD2L2) as a key inhibitor of end resection downstream of RIF1 (REF. 134).

A promiscuous polymerase for a sloppy repair pathway.

The robust fusions observed in shelterin-free and Ku70–Ku80-deficient cells enabled investigation of the basis of the increased mutagenicity of alt-NHEJ. Using deep sequencing, non-TTAGGG nucleotide insertions were identified at the junction between two fused telomeres¹³⁵ (FIG. 5). Such random insertions had previously been identified in other cells in which alt-NHEJ is active¹³⁶ and provide a molecular signature for this repair pathway. To identify the enzymatic activity responsible for these insertions, a number of low-fidelity DNA polymerases were inhibited in shelterin-free, Ku80-null cells. This led to the identification of the translesion DNA polymerase Pol θ ¹³⁷ as a key alt-NHEJ factor that catalyses the addition of random nucleotides at fusion junctions¹³⁵ and stimulates the joining of opposing ends of a broken DNA¹³⁸ (FIG. 5). Depletion of mammalian Pol θ hinders alt-NHEJ at uncapped telomeres, blocks non-reciprocal chromosomal translocations in mouse embryonic stem cells¹³⁵ and inhibits repair of endonuclease-induced DNA breaks^{138–140}. The function of this translesion polymerase is conserved in *Drosophila melanogaster*¹⁴¹ and *Caenorhabditis elegans*¹⁴². Notably, Pol θ inhibition in mammalian cells was marked by an increase in homologous recombination^{135,139}, indicating that the erroneous polymerase potentially influences the choice of DSB repair pathway in S phase, when both homologous recombination and alt-NHEJ are most active⁷⁶.

Pol θ is overexpressed in several human cancers^{143,144}, especially those with homologous recombination deficiency¹³⁹. Interestingly, depletion of Pol θ in BRCA-mutant tumour cells resulted in significant accumulation of chromosomal aberrations and unrepaired breaks, and compromised cellular survival^{135,139}. Given the increased mutagenicity of alt-NHEJ, it is tempting to speculate that this compensatory mechanism shapes the genome of homologous recombination-defective cancers and therefore influences tumour progression and resistance to therapy.

Telomeres gone bad

Alterations in the activity of telomere-associated proteins are important factors in the onset of human diseases. Dyskeratosis congenita, the prototypical telomere biology disorder (TBD), is caused by mutations in genes involved in telomere length regulation. To date, known mutations causing dyskeratosis congenita have been found in telomerase genes (telomerase RNA template component (*TERC*) and telomerase reverse transcriptase (*TERT*)), the *TERC*-regulating gene *H/ACA* ribonucleoprotein complex subunit 4 (also known as dyskerin), and the genes encoding the shelterin component TIN2 and the helicase RTEL1, as well as in *TCAB1* (also known as *WRAP53*), *NOP10* and *NHP2* (REFS 145–154). Dyskeratosis congenita patients have critically short telomeres and display a plethora of symptoms that range from impaired tissue regeneration capacity to cognitive defects. Severe variants of dyskeratosis congenita include Hoyeraal–Hreidarsson syndrome and Revesz syndrome. Less clinically severe variants, such as subsets of apparently isolated aplastic anaemia or pulmonary fibrosis,

Telomere biology disorder (TBD). One of a set of pathologies that are defined by the presence of short telomeres.

have also been recognized as TBDs. The genetic basis of TBDs, as well their clinical manifestations and implications, have recently been discussed in an excellent review¹⁵⁵. Here, we focus exclusively on how alterations in telomere length and in telomere-associated proteins affect genomic stability and have an impact on cancer development.

The good the bad and the ugly: telomeres and cancer.

In certain types of cancer, telomere dysfunction is considered to be a key trigger for chromosomal instability and a promoter of tumorigenesis (FIG. 6a). Rapid proliferation of pre-neoplastic cells leads to gradual telomere shortening, which ultimately triggers a DDR, inducing

cellular senescence and/or apoptosis. This illustrates the tumour suppressor function of telomere shortening that limits the proliferative potential of cancer cells^{156–158}. It is estimated that the accrual of five dysfunctional telomeres in a cell is sufficient to elicit a DDR and induce senescence¹⁵⁹. Despite losing their end protection, telomeres in senescent cells remain non-fusogenic, and it has been reasoned that this is due to the retention of few molecules of TRF2 that block end joining¹⁶⁰. Inactivation of p53 and/or RB pathways allows cells to bypass senescence, leading to telomere attrition and formation of dicentric chromosomes. This is known as telomere crisis, and it is estimated that ~50% of chromosome end-to-end fusions are completely devoid of TTAGGG

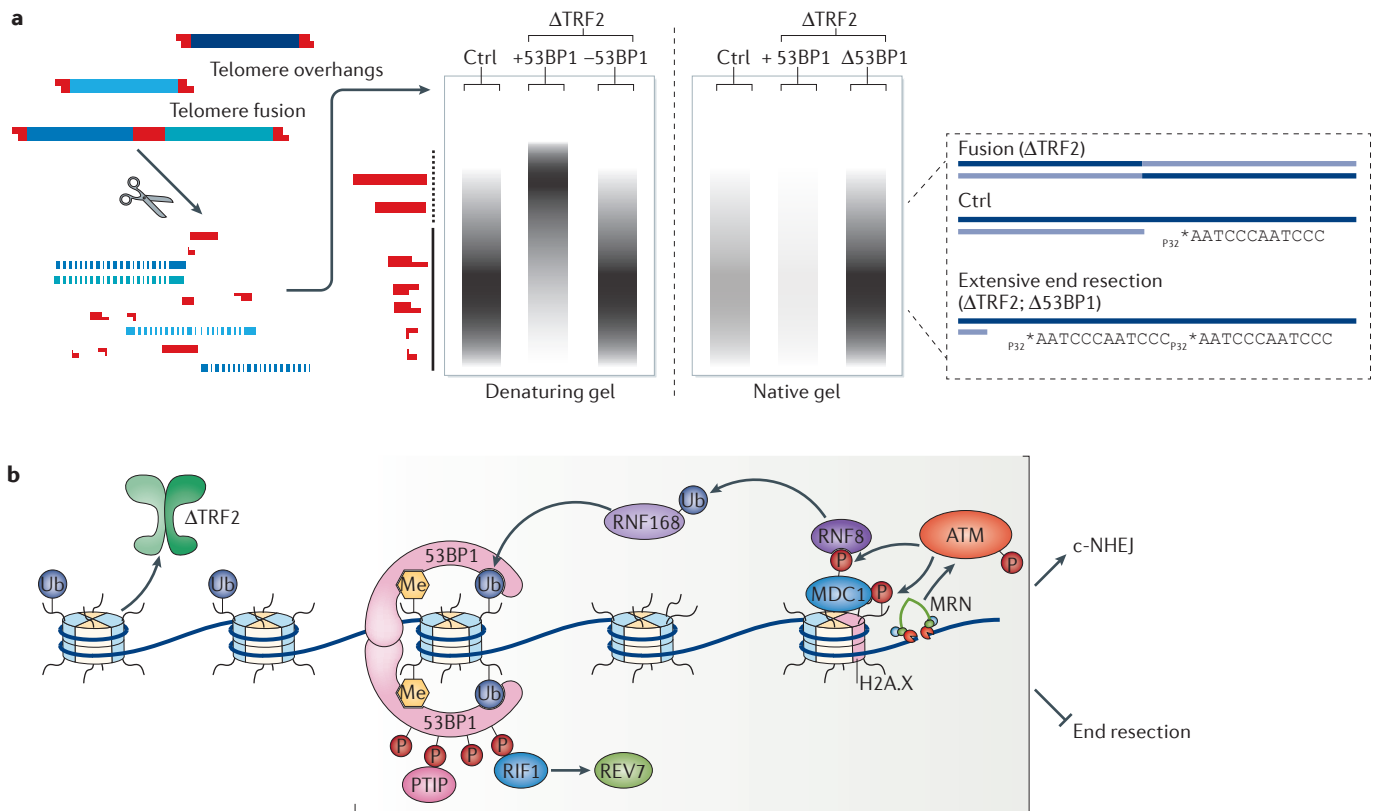


Figure 4 | Telomeres as a tool to investigate DNA end resection and classical non-homologous end joining (c-NHEJ). **a** | An overview of the assay to monitor c-NHEJ and DNA end resection at telomeres. Southern blot analysis allows the visualization of telomere fusion events. Genomic DNA is cleaved with frequently cutting restriction enzymes, resolved on a denaturing gel and hybridized with a radiolabelled telomere probe. As TTAGGG repeats are not cut by restriction enzymes, they are resolved according to their length (in range of the solid vertical line). Telomere fusions that occur following telomere repeat-binding factor 2 (TRF2) depletion are delineated as slow-migrating restriction fragments (dotted vertical line). Inhibition of factors that promote end joining, one example being p53-binding protein 1 (53BP1), prevents the accumulation of these long restriction fragments. This assay can be adjusted to quantify the length of the 3' overhang, which is generated by 5' end resection. Specifically, in-gel hybridization is carried out using a radiolabelled telomere probe in native conditions, in which the probe only hybridizes to the terminal single-stranded DNA (ssDNA) telomere overhang. The overhang signal in the native gel is quantified and normalized to the total telomeric DNA. Depletion of

factors, including 53BP1 and RAP1-interacting factor 1 (RIF1), that block end resection will lead to excess overhang signal and can be readily examined with this assay. **b** | A schematic representing key players that promote c-NHEJ and block DNA end resection, focusing on factors that were studied in the context of dysfunctional telomeres in TRF2-deficient cells. Double-strand break (DSB) sensing by the MRE11–RAD50–NBS1 (MRN)^{196–198} complex triggers a signalling cascade by recruiting autophosphorylated ataxia telangiectasia mutated (ATM)³⁷. ATM then phosphorylates the histone variant H2A.X at Ser139 (REF. 197), which recruits MDC1 (mediator of DNA damage checkpoint protein 1) to sites of breaks¹⁹⁹. The phosphorylation of MDC1 by ATM leads to the sequential recruitment of the E3 ubiquitin ligases RNF8 (REF. 124) and RNF168. One substrate of RNF168 is H2AK15 (histone 2A Lys15), which, together with mono- and dimethylated H4K20, serves as a platform to recruit 53BP1 (REFS 128,200), which then recruits the effector proteins RIF1 (RAP1-interacting factor 1)¹²⁹ and PTIP (Pax transactivation domain-interacting protein)¹³⁰, both of which bind to phosphorylated 53BP1. RIF1 functions in part by recruiting REV7 (also known as MAD2L2) to sites of breaks, where it inhibits end resection¹³⁴.

repeats¹⁶¹. Although most cells succumb to telomere crisis, rare survivors reactivate telomerase (or engage ALT) to replenish telomere repeats and proliferate indefinitely.

It was first proposed by McClintock¹ that dicentric chromosomes undergo repeated cycles of breakage–fusion–bridge leading to chromosomal rearrangements. Almost eight decades later, the fate of human dicentric chromosomes derived from telomere fusions¹⁶² was traced using live-cell imaging. Surprisingly, dicentric

chromosomes form anaphase bridges that persist through mitosis and are processed by the cytoplasmic nuclease TREX1 (three prime repair exonuclease 1). Clones that survive this crisis stage display chromothripsis and kataegis, which are localized hypermutation events often found in cancer genomes. Interestingly, previous work suggested a crucial role for DNA ligase 3 in the survival of cells undergoing telomere dysfunction¹⁶³. It is therefore possible that DNA ligase 3 is required for processing of TREX1-generated DNA breaks, leading to hypermutagenesis.

This paradigm of telomere dysfunction and cancer has been tested *in vivo* using the telomerase-knockout mouse. When combined with p53 mutations, later generations of telomerase-null mice display accelerated tumour formation and a shift in the tumour spectrum towards mostly carcinomas^{164,165}, characterized by non-reciprocal translocations and chromosome fusions¹⁶⁴. One caveat with the telomerase-knockout mouse model is that constitutive telomerase deficiency constrains tumour progression. In order to firmly establish the function of telomere dysfunction in malignancy and metastasis, an inducible telomerase allele was studied in the context of a PTEN mouse model of prostate cancer. Reactivation of telomerase in tumour cells that have already experienced telomere dysfunction was sufficient to suppress DNA damage signalling, and, importantly, resulted in the formation of highly metastatic tumours that invade the bone¹⁶⁶.

In addition to these animal studies, evidence in support of a role for telomere dysfunction during tumorigenesis came from the analysis of telomeres in cells derived from cancer patients at different stages of the disease^{167–169}. Specifically, telomere fusions, which were detected molecularly using a PCR-based method, were evident in chronic lymphocytic leukaemia (CLL) and breast cancers, and were found to be predictive of poor prognosis^{66,67}.

The reactivation of human telomerase is crucial for malignant progression. Telomerase activity has been detected in ~90% of human cancers¹⁷⁰, and its inhibition limits the survival of human cancer cells¹⁷¹. Mutations in the promoter region of *TERT* are among the most prevalent mutations in cancers. The first mutations identified in a genome-wide association study of melanoma patients are in close proximity to the *TERT* transcription start site and create a binding motif for the ternary complex factor (TCF) and E-twenty-six (ETS)-domain transcription factors^{172,173}. Subsequent sequence analysis identified similar point mutations in the *TERT* promoter in a wide range of cancers^{172–182}; in glioblastomas, the mutations facilitate the recruitment of the multimeric GA-binding protein (GABP) transcription factor to the *TERT* promoter region¹⁸² (FIG. 6b). In many cases, the mutations correlate with increased *TERT* transcription and enhanced telomerase activity^{174,180}. The *TERT* promoter mutations were recently engineered into human embryonic stem cells (hES cells) using the CRISPR–Cas9 editing system. In pluripotent cells, which already express *TERT*, the mutations did not increase telomerase activity, but they prevented telomerase silencing upon

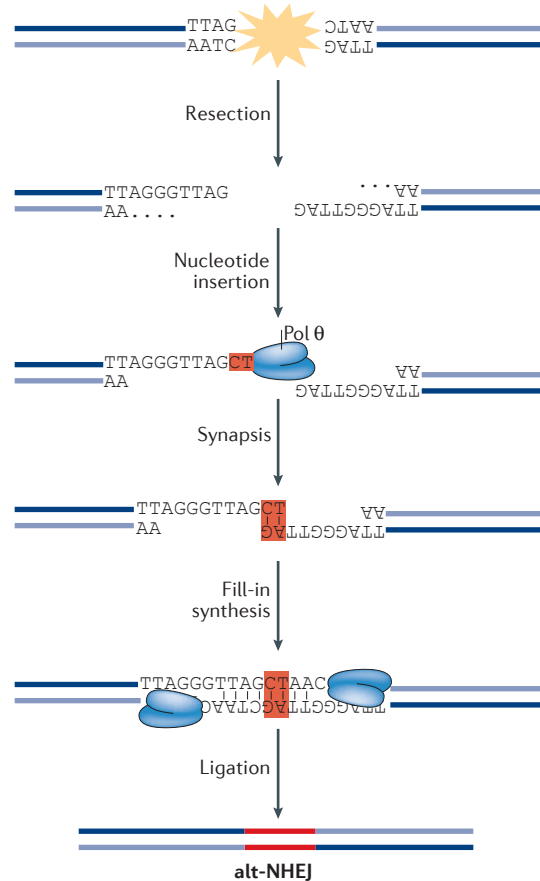


Figure 5 | The mechanism by which DNA polymerase θ (Pol θ) promotes alternative non-homologous end joining (alt-NHEJ). Sequence analysis of shelterin-free telomeres in Ku-deficient cells identified random nucleotide insertions at telomere fusion junctions. Subsequent genetic studies identified Pol θ as a key alt-NHEJ factor that promotes the joining of dysfunctional telomeres. Following double-strand break (DSB) formation or telomere uncapping, DNA ends are resected to create short 3' overhangs. On the basis of *in vitro* experiments, genetic studies and sequence analysis of fusion junctions, Pol θ seems to be capable of extending the 3' single-stranded DNA (ssDNA) using a combination of template-dependent as well as template-independent activities, the latter potentially mediated through a snap-back intermediate. The incorporation of random nucleotides as sites of breaks is predicted to increase the level of microhomology, thereby promoting the synapsis of opposite ends of a DSB. Annealed intermediates are then subject to fill-in synthesis by Pol θ , a step that would stabilize the duplexed DNA. Ultimately, the DNA is joined by DNA ligase 3.

Chromothripsis

A mutational phenomenon that involves catastrophic shattering and rebuilding of chromosomes, leading to multiple clustered chromosomal rearrangements.

Kataegis

Clustered point mutations that localize to particular regions of certain cancer genomes.

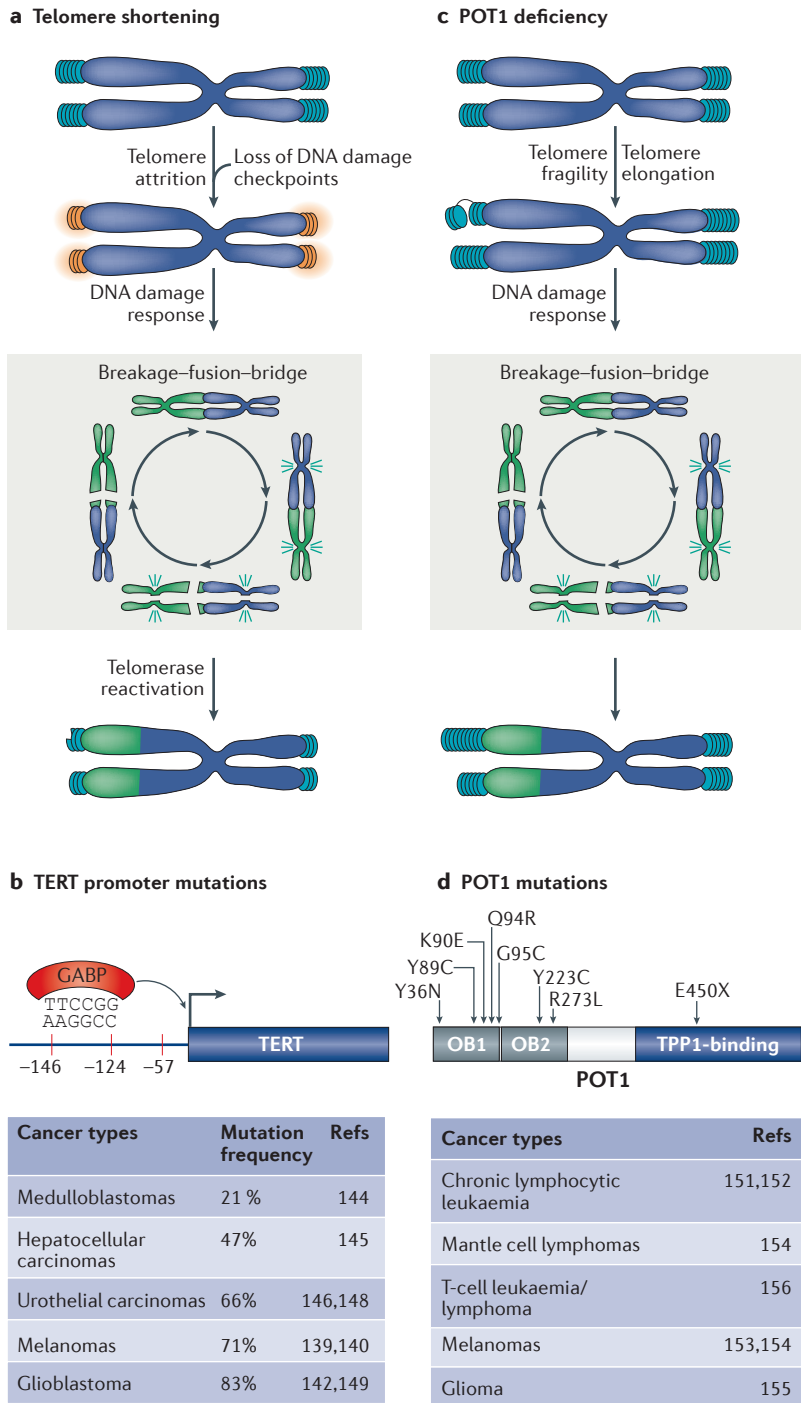


Figure 6 | Two independent pathways trigger telomere dysfunction in cancer. **a** | Telomere attrition induces telomere dysfunction and promotes gross chromosomal rearrangement due to breakage–fusion–bridge cycles (a non-reciprocal translocation is shown). Telomerase reactivation is a key event that stabilizes chromosome ends and supports the proliferation of tumours. **b** | Recurrent mutations in the *TERT* promoter are common in many cancers and seem to create a *de novo* binding site for the transcription factor GABP (GA-binding protein transcription factor). **c** | Deficiency in the shelterin subunit POT1 (protection of telomere 1) represent a novel mechanism that triggers telomere dysfunction in cancer. POT1 mutations induce telomere fragility and are associated with considerable telomere elongation. POT1 mutations also manifest in a mild chromosome fusion phenotype, which is predicted to induce chromosomal instability and augment tumour progression. **d** | POT1 mutations cluster primarily in its oligonucleotide-binding (OB) fold domains and are widespread across many tumour types.

differentiation of hES cells¹⁸³. Whether such mutations can lead to telomerase reactivation in somatic cells remains to be addressed. It is worth noting that a significant number of telomerase-positive tumours do not carry mutations in the *TERT* promoter, suggesting that additional *TERT*-activating pathway(s) may exist and are yet to be determined.

A novel way in which cancer cells do business. The sequencing of cancer genomes highlighted a potentially novel mechanism capable of inducing telomere dysfunction and promoting genomic instability in tumours (FIG. 6c). Acquired mutations in *POT1* were noted in CLL (chronic lymphocytic leukaemia)^{184,185}, and shortly thereafter, mis-sense variants of *POT1* were identified in familial melanoma^{186,187}, gliomas¹⁸⁸, mantle cell lymphomas¹⁸⁷, and T-cell leukaemia/lymphoma¹⁸⁹ (FIG. 6d). Interestingly, analysis of the clonal evolution of several mutations in CLL patients suggested that *POT1* mutations arise early in CLL development and are likely to contribute to disease progression¹⁹⁰. Despite their prevalence among many different cancer types, the mechanism by which *POT1* mutations induce telomere dysfunction and influence tumour progression is not fully understood. Limited functional analyses indicate that *POT1* mutations lead to telomere elongation, increased telomere fragility and mild telomere fusion phenotype^{185,191}. Chromosome end-to-end fusions can instigate breakage–fusion–bridge cycles, leading to increased genomic instability in *POT1*-mutated tumours. The observed telomere fragility suggests that replication defects triggered by *POT1* alterations constitute a novel type of tumour-promoting mechanism. Future experiments are necessary to reveal how the identified cancer-associated *POT1* mutations induce telomere fragility.

The identification of *POT1* mutations raises the question of whether other shelterin subunits, especially ones that induce telomere fragility, are mutated in cancers. Nonsense mutations in *TPP1* and *RAP1* were recently detected in melanoma¹⁹². Although no *TRF1* mutations have been detected so far, it is noteworthy that mice with reduced *TRF1* levels have increased incidence of lymphoid tumours¹⁹³, and deletion of *TRF1* in p53-null keratinocytes leads to squamous cell carcinomas⁸³. In the same way, mice carrying a mutation in *RTEL1*, which affects telomere replication, display accelerated tumorigenesis⁸⁵.

But humans are not mice! A recurrent concern in evaluating mouse models of human diseases is that humans, after all, are not mice. Telomere biology is not an exception and there are significant differences between human and mouse telomeres that need to be taken into consideration when evaluating mouse models of telomere dysfunction. Two key differences are the length of telomeres and the regulation of telomerase. Mice have significantly longer telomeres compared with humans, and they express telomerase in most cell types. As a result, telomere shortening is not a limiting factor in the lifespan of a mouse cell or in murine

tumours. The composition of shelterin is also different in that mice have two POT1 orthologues, POT1A and POT1B. POT1A is more closely related to human POT1 in its ability to prevent DDR activation, whereas POT1B is mainly involved in preventing resection at chromosomes²². Additional differences in the composition of telomeric chromatin between mouse and human telomeres might exist. The use of CRISPR–Cas9-mediated gene editing is likely to highlight the function of individual telomere-associated proteins in human cells and may provide additional insights into the differences between mouse and human telomeres.

Conclusions

It is now clear that there is a great degree of specialization within the shelterin complex in suppressing different DDR pathways. However, we still do not have a full understanding of the molecular mechanisms by which the shelterin subunits silence the DDR. Furthermore,

how fragile telomeres lead to genomic instability and tumour formation remains unknown and merits further investigation. Thus far, manipulating shelterin to trigger telomere uncapping has been successfully used to identify novel factors involved in DSB processing and repair. Deprotected telomeres continue to be used to improve our understanding of the DDR; for instance, telomeres may provide a powerful tool in which the role of histone modifications and chromatin remodelling factors in the DDR can be addressed. Sequencing of cancer genomes has provided a wealth of information on recurrent mutations in genes involved in telomere maintenance and protection. Continued investigation into how these mutations promote tumour progression and how cancerous cells evade the detrimental effect of telomere dysfunction will provide a greater understanding of the role of telomere biology in cancer progression and, hopefully, will guide the development of new and better therapies for cancer.

- McClintock, B. The stability of broken ends of chromosomes in *Zea mays*. *Genetics* **26**, 234–282 (1941).
- Meier, R. & Muller, R. A new arrangement for the registration of diaphragm movements. *J. Physiol.* **94**, 227–231 (1938).
- Greider, C. W. Telomerase and telomere-length regulation: lessons from small eukaryotes to mammals. *Cold Spring Harb. Symp. Quant. Biol.* **58**, 719–723 (1993).
- Makarov, V. L., Hirose, Y. & Langmore, J. P. Long G tails at both ends of human chromosomes suggest a C strand degradation mechanism for telomere shortening. *Cell* **88**, 657–666 (1997).
- McElligott, R. & Wellinger, R. J. The terminal DNA structure of mammalian chromosomes. *EMBO J.* **16**, 3705–3714 (1997).
- Griffith, J. D. *et al.* Mammalian telomeres end in a large duplex loop. *Cell* **97**, 503–514 (1999).
- Doksani, Y., Wu, J. Y., de Lange, T. & Zhuang, X. Super-resolution fluorescence imaging of telomeres reveals TRF2-dependent T-loop formation. *Cell* **155**, 345–356 (2013).
- Azzalin, C. M., Reichenbach, P., Khoriatou, L., Giulotto, E. & Lingner, J. Telomeric repeat containing RNA and RNA surveillance factors at mammalian chromosome ends. *Science* **318**, 798–801 (2007).
- Azzalin, C. M. & Lingner, J. Telomere functions grounding on TERRA firma. *Trends Cell Biol.* **25**, 29–36 (2015).
- de Lange, T. Shelterin: the protein complex that shapes and safeguards human telomeres. *Genes Dev.* **19**, 2100–2110 (2005).
- Chong, L. *et al.* A human telomeric protein. *Science* **270**, 1663–1667 (1995).
- Broccoli, D., Smogorzewska, A., Chong, L. & de Lange, T. Human telomeres contain two distinct Myb-related proteins, TRF1 and TRF2. *Nat. Genet.* **17**, 231–235 (1997).
- Bilaud, T. *et al.* Telomeric localization of TRF2, a novel human telobox protein. *Nat. Genet.* **17**, 236–239 (1997).
- Ye, J. Z. & de Lange, T. TIN2 is a tankyrase 1 PARP modulator in the TRF1 telomere length control complex. *Nat. Genet.* **36**, 618–623 (2004).
- Ye, J. Z. *et al.* POT1-interacting protein PIP1: a telomere length regulator that recruits POT1 to the TIN2/TRF1 complex. *Genes Dev.* **18**, 1649–1654 (2004).
- Houghtaling, B. R., Cuttonaro, L., Chang, W. & Smith, S. A dynamic molecular link between the telomere length regulator TRF1 and the chromosome end protector TRF2. *Curr. Biol.* **14**, 1621–1631 (2004).
- Kim, S. H., Kaminker, P. & Campisi, J. TIN2, a new regulator of telomere length in human cells. *Nat. Genet.* **23**, 405–412 (1999).
- Liu, D. *et al.* PTPOT interacts with POT1 and regulates its localization to telomeres. *Nat. Cell Biol.* **6**, 673–680 (2004).
- Baumann, P. & Cech, T. R. Pot1, the putative telomere end-binding protein in fission yeast and humans. *Science* **292**, 1171–1175 (2001).
- Loayza, D. & de Lange, T. POT1 as a terminal transducer of TRF1 telomere length control. *Nature* **423**, 1013–1018 (2003).
- Wu, L. *et al.* Pot1 deficiency initiates DNA damage checkpoint activation and aberrant homologous recombination at telomeres. *Cell* **126**, 49–62 (2006).
- Hockemeyer, D., Daniels, J. P., Takai, H. & de Lange, T. Recent expansion of the telomeric complex in rodents: two distinct POT1 proteins protect mouse telomeres. *Cell* **126**, 63–77 (2006).
- Li, B., Oestreich, S. & de Lange, T. Identification of human Rap1: implications for telomere evolution. *Cell* **101**, 471–483 (2000).
- Celli, G. B. & de Lange, T. DNA processing is not required for ATM-mediated telomere damage response after TRF2 deletion. *Nat. Cell Biol.* **7**, 712–718 (2005).
- Sfeir, A., Kabir, S., van Overbeek, M., Celli, G. B. & de Lange, T. Loss of Rap1 induces telomere recombination in the absence of NHEJ or a DNA damage signal. *Science* **327**, 1657–1661 (2010).
- Ye, J. Z. *et al.* TIN2 binds TRF1 and TRF2 simultaneously and stabilizes the TRF2 complex on telomeres. *J. Biol. Chem.* **279**, 47264–47271 (2004).
- Dejardin, J. & Kingston, R. E. Purification of proteins associated with specific genomic loci. *Cell* **136**, 175–186 (2009).
- Grolimund, L. *et al.* A quantitative telomeric chromatin isolation protocol identifies different telomeric states. *Nat. Commun.* **4**, 2848 (2013).
- Bartocci, C. *et al.* Isolation of chromatin from dysfunctional telomeres reveals an important role for Ring1b in NHEJ-mediated chromosome fusions. *Cell Rep.* **7**, 1320–1332 (2014).
- Nittis, T. *et al.* Revealing novel telomere proteins using *in vivo* cross-linking, tandem affinity purification, and label-free quantitative LC-FTICR-MS. *Mol. Cell. Proteomics* **9**, 1144–1156 (2010).
- Miyake, Y. *et al.* RPA-like mammalian Ctc1–Stn1–Tcn1 complex binds to single-stranded DNA and protects telomeres independently of the Pot1 pathway. *Mol. Cell* **36**, 193–206 (2009).
- Surovtseva, Y. V. *et al.* Conserved telomere maintenance component 1 interacts with STN1 and maintains chromosome ends in higher eukaryotes. *Mol. Cell* **36**, 207–218 (2009).
- Shibuya, H. *et al.* MAJIN links telomeric DNA to the nuclear membrane by exchanging telomere cap. *Cell* **163**, 1252–1266 (2015).
- Shibuya, H., Ishiguro, K. & Watanabe, Y. The TRF1-binding protein TERB1 promotes chromosome movement and telomere rigidity in meiosis. *Nat. Cell Biol.* **16**, 145–156 (2014).
- Shiloh, Y. ATM and related protein kinases: safeguarding genome integrity. *Nat. Rev. Cancer* **3**, 155–168 (2003).
- Karlseder, J. *et al.* The telomeric protein TRF2 binds the ATM kinase and can inhibit the ATM-dependent DNA damage response. *PLoS Biol.* **2**, E240 (2004).
- Denchi, E. L. & de Lange, T. Protection of telomeres through independent control of ATM and ATR by TRF2 and POT1. *Nature* **448**, 1068–1071 (2007). **In this publication, the authors demonstrate that ATM and ATR signalling are repressed by TRF2 and POT1, respectively, and that efficient end joining of deprotected telomeres is dependent on DNA damage signalling.**
- Takai, H., Smogorzewska, A. & de Lange, T. DNA damage foci at dysfunctional telomeres. *Curr. Biol.* **13**, 1549–1556 (2003).
- Amiard, S. *et al.* A topological mechanism for TRF2-enhanced strand invasion. *Nat. Struct. Mol. Biol.* **14**, 147–154 (2007).
- Okamoto, K. *et al.* A two-step mechanism for TRF2-mediated chromosome-end protection. *Nature* **494**, 502–505 (2013). **This publication highlights a mechanism by which TRF2 inhibits ATM signalling and identifies the iDDR domain of TRF2 as important for RNF168 inhibition.**
- Benarroch-Popivker, D. *et al.* TRF2-mediated control of telomere DNA topology as a mechanism for chromosome-end protection. *Mol. Cell* **61**, 274–286 (2016).
- van Steensel, B., Smogorzewska, A. & de Lange, T. TRF2 protects human telomeres from end-to-end fusions. *Cell* **92**, 401–413 (1998). **This report demonstrates that deprotection of telomeres by inhibiting the function of human TRF2 leads to chromosome fusions.**
- Fumagalli, M. *et al.* Telomeric DNA damage is irreparable and causes persistent DNA-damage-response activation. *Nat. Cell Biol.* **14**, 355–365 (2012).
- Liu, D., O'Connor, M. S., Qin, J. & Songyang, Z. Telosome, a mammalian telomere-associated complex formed by multiple telomeric proteins. *J. Biol. Chem.* **279**, 51338–51342 (2004).
- Takai, K. K., Hooper, S., Blackwood, S., Gandhi, R. & de Lange, T. *In vivo* stoichiometry of shelterin components. *J. Biol. Chem.* **285**, 1457–1467 (2010).
- Sarthy, J., Bae, N. S., Scraftford, J. & Baumann, P. Human RAP1 inhibits non-homologous end joining at telomeres. *EMBO J.* **28**, 3390–3399 (2009).
- Kabir, S., Hockemeyer, D. & de Lange, T. TALEN gene knockouts reveal no requirement for the conserved human shelterin protein Rap1 in telomere protection and length regulation. *Cell Rep.* **9**, 1273–1280 (2014).
- Martinez, P. *et al.* Mammalian Rap1 controls telomere function and gene expression through binding to telomeric and extratelomeric sites. *Nat. Cell Biol.* **12**, 768–780 (2010).
- Hsu, H. L., Gilley, D., Blackburn, E. H. & Chen, D. J. Ku is associated with the telomere in mammals. *Proc. Natl Acad. Sci. USA* **96**, 12454–12458 (1999).

50. Wang, Y., Ghosh, G. & Hendrickson, E. A. Ku86 represses lethal telomere deletion events in human somatic cells. *Proc. Natl Acad. Sci. USA* **106**, 12430–12435 (2009).
51. Gu, Y. *et al.* Growth retardation and leaky SCID phenotype of Ku70-deficient mice. *Immunity* **7**, 653–665 (1997).
52. Samper, E., Goytisolo, F. A., Slijepcevic, P., van Buul, P. P. & Blasco, M. A. Mammalian Ku86 protein prevents telomeric fusions independently of the length of TTAGGG repeats and the G-strand overhang. *EMBO Rep.* **1**, 244–252 (2000).
53. Ribes-Zamora, A., Indiviglio, S. M., Mihalek, I., Williams, C. L. & Bertuch, A. A. TRF2 interaction with Ku heterotetramerization interface gives insight into c-NHEJ prevention at human telomeres. *Cell Rep.* **5**, 194–206 (2013).
54. Hayashi, M. T., Cesare, A. J., Fitzpatrick, J. A., Lazzarini-Denchi, E. & Karlseder, J. A telomere-dependent DNA damage checkpoint induced by prolonged mitotic arrest. *Nat. Struct. Mol. Biol.* **19**, 387–394 (2012).
55. Orthwein, A. *et al.* Mitosis inhibits DNA double-strand break repair to guard against telomeric fusions. *Science* **344**, 189–193 (2014).
This publication identifies the molecular mechanism that allows mammalian cells to inhibit DSB repair during mitosis. The authors show that restoring mitotic DSB repair results in severe genomic instability owing to the accumulation of telomere fusions.
56. Hayashi, M. T., Cesare, A. J., Rivera, T. & Karlseder, J. Cell death during crisis is mediated by mitotic telomere depletion. *Nature* **522**, 492–496 (2015).
57. Guo, X. *et al.* Dysfunctional telomeres activate an ATM-ATR-dependent DNA damage response to suppress tumorigenesis. *EMBO J.* **26**, 4709–4719 (2007).
58. Kibe, T., Osawa, G. A., Keegan, C. E. & de Lange, T. Telomere protection by TPP1 is mediated by POT1a and POT1b. *Mol. Cell Biol.* **30**, 1059–1066 (2010).
59. Takai, K. K., Kibe, T., Donigian, J. R., Frescas, D. & de Lange, T. Telomere protection by TPP1/POT1 requires tethering to TIN2. *Mol. Cell* **44**, 647–659 (2011).
60. Gong, Y. & de Lange, T. A. Shld1-controlled POT1a provides support for repression of ATR signaling at telomeres through RPA exclusion. *Mol. Cell* **40**, 377–387 (2010).
61. Zou, L. & Elledge, S. J. Sensing DNA damage through ATRIP recognition of RPA-ssDNA complexes. *Science* **300**, 1542–1548 (2003).
62. Flynn, R. L. *et al.* TERRA and hnRNP1 orchestrate an RPA-to-POT1 switch on telomeric single-stranded DNA. *Nature* **471**, 532–536 (2011).
63. Sfeir, A. *et al.* Mammalian telomeres resemble fragile sites and require TRF1 for efficient replication. *Cell* **138**, 90–103 (2009).
This publication identifies telomere replication defects as the major consequence of TRF1 loss in mouse cells and defines telomere fragility as a hallmark of replication stress.
64. Zimmermann, M., Kibe, T., Kabir, S. & de Lange, T. TRF1 negotiates TTAGGG repeat-associated replication problems by recruiting the BLM helicase and the TPP1/POT1 repressor of ATR signaling. *Genes Dev.* **28**, 2477–2491 (2014).
65. Wong, K. K. *et al.* Diminished lifespan and acute stress-induced death in DNA-PKcs-deficient mice with limiting telomeres. *Oncogene* **26**, 2815–2821 (2007).
66. Lin, T. T. *et al.* Telomere dysfunction and fusion during the progression of chronic lymphocytic leukemia: evidence for a telomere crisis. *Blood* **116**, 1899–1907 (2010).
In this study, the authors carry out single-molecule telomere analysis in patients with chronic lymphocytic leukaemia and report the incidence of telomere erosion and, subsequently, telomere fusions in advanced stages of the disease.
67. Simpson, K. *et al.* Telomere fusion threshold identifies a poor prognostic subset of breast cancer patients. *Mol. Oncol.* **9**, 1186–1193 (2015).
68. Letsolo, B. T., Rowson, J. & Baird, D. M. Fusion of short telomeres in human cells is characterized by extensive deletion and microhomology, and can result in complex rearrangements. *Nucleic Acids Res.* **38**, 1841–1852 (2010).
69. Sfeir, A. & de Lange, T. Removal of shelterin reveals the telomere end-protection problem. *Science* **336**, 593–597 (2012).
This publication delineates the process of end protection by removing all six subunits of the shelterin complex and identifying the various DNA damage signalling and repair pathways activated at deprotected chromosome ends.
70. Rai, R. *et al.* The function of classical and alternative non-homologous end-joining pathways in the fusion of dysfunctional telomeres. *EMBO J.* **29**, 2598–2610 (2010).
71. Badie, S. *et al.* BRCA1 and CtIP promote alternative non-homologous end-joining at uncapped telomeres. *EMBO J.* **34**, 410–424 (2015).
72. Cheng, Q. *et al.* Ku counteracts mobilization of PARP1 and MRN in chromatin damaged with DNA double-strand breaks. *Nucleic Acids Res.* **39**, 9605–9619 (2011).
73. Wang, M. *et al.* PARP-1 and Ku compete for repair of DNA double strand breaks by distinct NHEJ pathways. *Nucleic Acids Res.* **34**, 6170–6182 (2006).
74. Clerici, M., Mantiero, D., Guerini, I., Lucchini, G. & Longhese, M. P. The Yku70-Yku80 complex contributes to regulate double-strand break processing and checkpoint activation during the cell cycle. *EMBO Rep.* **9**, 810–818 (2008).
75. Mimitou, E. P. & Symington, L. S. Ku prevents Exo1 and Sgs1-dependent resection of DNA ends in the absence of a functional MRX complex or Sae2. *EMBO J.* **29**, 3358–3369 (2010).
76. Truong, L. N. *et al.* Microhomology-mediated end joining and homologous recombination share the initial end resection step to repair DNA double-strand breaks in mammalian cells. *Proc. Natl Acad. Sci. USA* **110**, 7720–7725 (2013).
77. Wu, P., Takai, H. & de Lange, T. Telomeric 3' overhangs derive from resection by Exo1 and Apollo and fill-in by POT1b-associated CST. *Cell* **150**, 39–52 (2012).
78. Wu, P., van Overbeek, M., Rooney, S. & de Lange, T. Apollo contributes to G overhang maintenance and protects leading-end telomeres. *Mol. Cell* **39**, 606–617 (2010).
79. Lam, Y. C. *et al.* SNM1B/Apollo protects leading-strand telomeres against NHEJ-mediated repair. *EMBO J.* **29**, 2230–2241 (2010).
80. Chai, W., Shay, J. W. & Wright, W. E. Human telomeres maintain their overhang length at senescence. *Mol. Cell Biol.* **25**, 2158–2168 (2005).
81. Lotterberger, F., Bothmer, A., Robbiani, D. F., Nussenzweig, M. C. & de Lange, T. Role of 53BP1 oligomerization in regulating double-strand break repair. *Proc. Natl Acad. Sci. USA* **110**, 2146–2151 (2013).
82. Parkinson, G. N., Lee, M. P. & Neidle, S. Crystal structure of parallel quadruplexes from human telomeric DNA. *Nature* **417**, 876–880 (2002).
83. Martinez, P. *et al.* Increased telomere fragility and fusions resulting from TRF1 deficiency lead to degenerative pathologies and increased cancer in mice. *Genes Dev.* **23**, 2060–2075 (2009).
84. Miller, K. M., Rog, O. & Cooper, J. P. Semi-conservative DNA replication through telomeres requires Taz1. *Nature* **440**, 824–828 (2006).
85. Vannier, J. B. *et al.* RTEL1 is a replisome-associated helicase that promotes telomere and genome-wide replication. *Science* **342**, 239–242 (2013).
This publication describes an interaction between RTEL1 and PCNA. Blocking this interaction affects genome-wide replication, induces telomere fragility and accelerates the onset of tumorigenesis in p53-deficient mice.
86. Drosopoulos, W. C., Kosiyatrakul, S. T. & Schildkraut, C. L. BLM helicase facilitates telomere replication during leading strand synthesis of telomeres. *J. Cell Biol.* **210**, 191–208 (2015).
87. Crabbe, L., Verdun, R. E., Haggblom, C. I. & Karlseder, J. Defective telomere lagging strand synthesis in cells lacking WRN helicase activity. *Science* **306**, 1951–1953 (2004).
88. Arnoult, N., Saintome, C., Ourliac-Garnier, I., Riou, J. F. & Londoño-Vallejo, A. Human POT1 is required for efficient telomere C-rich strand replication in the absence of WRN. *Genes Dev.* **23**, 2915–2924 (2009).
89. Stewart, J. A. *et al.* Human CST promotes telomere duplex replication and general replication restart after fork stalling. *EMBO J.* **31**, 3537–3549 (2012).
90. Kasbek, C., Wang, F. & Price, C. M. Human TEN1 maintains telomere integrity and functions in genome-wide replication restart. *J. Biol. Chem.* **288**, 30139–30150 (2013).
91. Ye, J. *et al.* TRF2 and Apollo cooperate with topoisomerase 2 α to protect human telomeres from replicative damage. *Cell* **142**, 230–242 (2010).
92. Pierce, A. J., Hu, P., Han, M., Ellis, N. & Jasin, M. Ku DNA end-binding protein modulates homologous repair of double-strand breaks in mammalian cells. *Genes Dev.* **15**, 3237–3242 (2001).
93. Celli, G. B., Denchi, E. L. & de Lange, T. Ku70 stimulates fusion of dysfunctional telomeres yet protects chromosome ends from homologous recombination. *Nat. Cell Biol.* **8**, 885–890 (2006).
94. Palm, W., Hockemeyer, D., Kibe, T. & de Lange, T. Functional dissection of human and mouse POT1 proteins. *Mol. Cell Biol.* **29**, 471–482 (2009).
95. Arat, N. O. & Griffith, J. D. Human Rap1 interacts directly with telomeric DNA and regulates TRF2 localization at the telomere. *J. Biol. Chem.* **287**, 41583–41594 (2012).
96. Sarek, G., Vannier, J. B., Panier, S., Petrini, J. H. & Boulton, S. J. TRF2 recruits RTEL1 to telomeres in S phase to promote t-loop unwinding. *Mol. Cell* **57**, 622–635 (2015).
97. Vannier, J. B., Pavicic-Kaltenbrunner, V., Petalcorin, M. I., Ding, H. & Boulton, S. J. RTEL1 dismantles T loops and counteracts telomeric G4-DNA to maintain telomere integrity. *Cell* **149**, 795–806 (2012).
98. Wang, R. C., Smogorzewska, A. & de Lange, T. Homologous recombination generates Floop-sized deletions at human telomeres. *Cell* **119**, 355–368 (2004).
99. Compton, S. A., Choi, J. H., Cesare, A. J., Ozgur, S. & Griffith, J. D. Xrcc3 and Nbs1 are required for the production of extrachromosomal telomeric circles in human alternative lengthening of telomere cells. *Cancer Res.* **67**, 1513–1519 (2007).
100. Pickett, H. A., Cesare, A. J., Johnston, R. L., Neumann, A. A. & Reddel, R. R. Control of telomere length by a trimming mechanism that involves generation of t-circles. *EMBO J.* **28**, 799–809 (2009).
101. Pickett, H. A., Henson, J. D., Au, A. Y., Neumann, A. A. & Reddel, R. R. Normal mammalian cells negatively regulate telomere length by telomere trimming. *Hum. Mol. Genet.* **20**, 4684–4692 (2011).
102. Bryan, T. M., Englezou, A., Dalla-Pozza, L., Dunham, M. A. & Reddel, R. R. Evidence for an alternative mechanism for maintaining telomere length in human tumors and tumor-derived cell lines. *Nat. Med.* **3**, 1271–1274 (1997).
This report identifies ALT as a telomerase-independent mechanism that maintains telomeres in a subset of tumours.
103. Dunham, M. A., Neumann, A. A., Fasching, C. L. & Reddel, R. R. Telomere maintenance by recombination in human cells. *Nat. Genet.* **26**, 447–450 (2000).
104. Yeager, T. R. *et al.* Telomerase-negative immortalized human cells contain a novel type of promyelocytic leukemia (PML) body. *Cancer Res.* **59**, 4175–4179 (1999).
105. Arora, R. *et al.* rNaseH1 regulates TERRA-telomeric DNA hybrids and telomere maintenance in ALT tumour cells. *Nat. Commun.* **5**, 5220 (2014).
106. Londoño-Vallejo, J. A., Der-Sarkissian, H., Cazes, L., Bacchetti, S. & Reddel, R. R. Alternative lengthening of telomeres is characterized by high rates of telomeric exchange. *Cancer Res.* **64**, 2324–2327 (2004).
107. Bryan, T. M., Englezou, A., Gupta, J., Bacchetti, S. & Reddel, R. R. Telomere elongation in immortal human cells without detectable telomerase activity. *EMBO J.* **14**, 4240–4248 (1995).
108. Conomos, D. *et al.* Variant repeats are interspersed throughout the telomeres and recruit nuclear receptors in ALT cells. *J. Cell Biol.* **199**, 893–906 (2012).
109. Gocha, A. R., Harris, J. & Groden, J. Alternative mechanisms of telomere lengthening: permissive mutations, DNA repair proteins and tumorigenic progression. *Mutat. Res.* **743–744**, 142–150 (2013).
110. Heaphy, C. M. *et al.* Altered telomeres in tumors with ATRX and DAXX mutations. *Science* **333**, 425 (2011).
The authors of this report discover recurrent, inactivating mutations in ATRX and DAXX in telomerase-positive human pancreatic neuroendocrine tumours, which rely on ALT to maintain telomeres.
111. Jiao, Y. *et al.* DAXX/ATRX, MEN1, and mTOR pathway genes are frequently altered in pancreatic neuroendocrine tumors. *Science* **331**, 1199–1203 (2011).
112. Clynes, D. *et al.* Suppression of the alternative lengthening of telomere pathway by the chromatin remodelling factor ATRX. *Nat. Commun.* **6**, 7538 (2015).

113. Napier, C. E. *et al.* ATRX represses alternative lengthening of telomeres. *Oncotarget* **6**, 16543–16558 (2015).
114. Gibbons, R. J., Picketts, D. J., Villard, L. & Higgs, D. R. Mutations in a putative global transcriptional regulator cause X-linked mental retardation with α -thalassaemia (ATR-X syndrome). *Cell* **80**, 837–845 (1995).
115. Garcia-Cao, M., O'Sullivan, R., Peters, A. H., Jenuwein, T. & Blasco, M. A. Epigenetic regulation of telomere length in mammalian cells by the Suv39h1 and Suv39h2 histone methyltransferases. *Nat. Genet.* **36**, 94–99 (2004).
116. Goldberg, A. D. *et al.* Distinct factors control histone variant H3.3 localization at specific genomic regions. *Cell* **140**, 678–691 (2010).
117. Wong, L. H. *et al.* ATRX interacts with H3.3 in maintaining telomere structural integrity in pluripotent embryonic stem cells. *Genome Res.* **20**, 351–360 (2010).
118. Law, M. J. *et al.* ATR-X syndrome protein targets tandem repeats and influences allele-specific expression in a size-dependent manner. *Cell* **143**, 367–378 (2010).
119. Flynn, R. L. *et al.* Alternative lengthening of telomeres renders cancer cells hypersensitive to ATR inhibitors. *Science* **347**, 273–277 (2015).
120. O'Sullivan, R. J. *et al.* Rapid induction of alternative lengthening of telomeres by depletion of the histone chaperone ASF1. *Nat. Struct. Mol. Biol.* **21**, 167–174 (2014).
121. Conomos, D., Reddel, R. R. & Pickett, H. A. NuRD-ZNF827 recruitment to telomeres creates a molecular scaffold for homologous recombination. *Nat. Struct. Mol. Biol.* **21**, 760–770 (2014).
122. Ramamoorthy, M. & Smith, S. Loss of ATRX suppresses resolution of telomere cohesion to control recombination in ALT cancer cells. *Cancer Cell* **28**, 357–369 (2015).
123. Cho, N. W., Dilley, R. L., Lampson, M. A. & Greenberg, R. A. Interchromosomal homology searches drive directional ALT telomere movement and synapsis. *Cell* **159**, 108–121 (2014).
124. Peuscher, M. H. & Jacobs, J. J. DNA-damage response and repair activities at uncapped telomeres depend on RNF8. *Nat. Cell Biol.* **13**, 1159–1145 (2011).
125. Porro, A., Feuerhahn, S. & Lingner, J. TERRA-reinforced association of LSD1 with MRE11 promotes processing of uncapped telomeres. *Cell Rep.* **6**, 765–776 (2014).
126. Bunting, S. F. *et al.* 53BP1 inhibits homologous recombination in *Brca1*-deficient cells by blocking resection of DNA breaks. *Cell* **141**, 243–254 (2010). **In this report, the authors demonstrate that loss of 53BP1 rescues phenotypes associated with BRCA1 deficiency by allowing cells to overcome defects in homologous recombination.**
127. Bouwman, P. *et al.* 53BP1 loss rescues BRCA1 deficiency and is associated with triple-negative and BRCA-mutated breast cancers. *Nat. Struct. Mol. Biol.* **17**, 688–695 (2010).
128. Dimitrova, N., Chen, Y. C., Spector, D. L. & de Lange, T. 53BP1 promotes non-homologous end joining of telomeres by increasing chromatin mobility. *Nature* **456**, 524–528 (2008).
129. Zimmermann, M., Lottersberger, F., Buonomo, S. B., Sfeir, A. & de Lange, T. 53BP1 regulates DSB repair using Rif1 to control 5' end resection. *Science* **339**, 700–704 (2013).
130. Callen, E. *et al.* 53BP1 mediates productive and mutagenic DNA repair through distinct phosphoprotein interactions. *Cell* **153**, 1266–1280 (2013).
131. Escribano-Diaz, C. *et al.* A cell cycle-dependent regulatory circuit composed of 53BP1-RIF1 and BRCA1-CtIP controls DNA repair pathway choice. *Mol. Cell* **49**, 872–883 (2013). **This publication defines the molecular mechanisms that allow c-NHEJ to dominate in the G1 phase of the cell cycle and enables homologous recombination to be favoured in S/G2. The authors identify RIF1 as a key effector of 53BP1 that inhibits resection at DSBs.**
132. Chapman, J. R. *et al.* RIF1 is essential for 53BP1-dependent nonhomologous end joining and suppression of DNA double-strand break resection. *Mol. Cell* **49**, 858–871 (2013).
133. Di Virgilio, M. *et al.* Rif1 prevents resection of DNA breaks and promotes immunoglobulin class switching. *Science* **339**, 711–715 (2013).
134. Boersma, V. *et al.* MAD2L2 controls DNA repair at telomeres and DNA breaks by inhibiting 5' end resection. *Nature* **521**, 537–540 (2015).
135. Mateos-Gomez, P. A. *et al.* Mammalian polymerase θ promotes alternative NHEJ and suppresses recombination. *Nature* **518**, 254–257 (2015).
136. Simsek, D. *et al.* DNA ligase III promotes alternative nonhomologous end-joining during chromosomal translocation formation. *PLoS Genet.* **7**, e1002080 (2011). **This study identifies Pol θ as a key factor that promotes alt-NHEJ. Inhibition of Pol θ increases the frequency of homologous recombination and compromised the survival of homologous recombination-defective cells.**
137. Seki, M., Marini, F. & Wood, R. D. POLQ (Pol θ), a DNA polymerase and DNA-dependent ATPase in human cells. *Nucleic Acids Res.* **31**, 6117–6126 (2003).
138. Kent, T., Chandramouly, G., McDevitt, S. M., Ozdemir, A. Y. & Pomerantz, R. T. Mechanism of microhomology-mediated end-joining promoted by human DNA polymerase θ . *Nat. Struct. Mol. Biol.* **22**, 230–237 (2015).
139. Ceccaldi, R. *et al.* Homologous-recombination-deficient tumours are dependent on Pol θ -mediated repair. *Nature* **518**, 258–262 (2015).
140. Yousefzadeh, M. J. *et al.* Mechanism of suppression of chromosomal instability by DNA polymerase POLQ. *PLoS Genet.* **10**, e1004654 (2014).
141. Chan, S. H., Yu, A. M. & McVey, M. Dual roles for DNA polymerase θ in alternative end-joining repair of double-strand breaks in *Drosophila*. *PLoS Genet.* **6**, e1001005 (2010).
142. Koole, W. *et al.* A polymerase theta-dependent repair pathway suppresses extensive genomic instability at endogenous G4 DNA sites. *Nat. Commun.* **5**, 3216 (2014).
143. Lemea, F. *et al.* DNA polymerase θ up-regulation is associated with poor survival in breast cancer, perturbs DNA replication, and promotes genetic instability. *Proc. Natl Acad. Sci. USA* **107**, 13390–13395 (2010).
144. Higgins, G. S. *et al.* Overexpression of POLQ confers a poor prognosis in early breast cancer patients. *Oncotarget* **1**, 175–184 (2010).
145. Heiss, N. S. X-linked dyskeratosis congenita is caused by mutations in a highly conserved gene with putative nuclear functions. *Nat. Genet.* **19**, 32–38 (1998).
146. Vulliamy, T. J. *et al.* Mutations in the reverse transcriptase component of telomerase (TERT) in patients with bone marrow failure. *Blood Cells Mol. Dis.* **34**, 257–263 (2005).
147. Vulliamy, T. *et al.* The RNA component of telomerase is mutated in autosomal dominant dyskeratosis congenita. *Nature* **413**, 432–435 (2001).
148. Walne, A. J. *et al.* Genetic heterogeneity in autosomal recessive dyskeratosis congenita with one subtype due to mutations in the telomerase-associated protein NOP10. *Hum. Mol. Genet.* **16**, 1619–1629 (2007).
149. Walne, A. J., Vulliamy, T., Beswick, R., Kirwan, M. & Dokal, I. TINF2 mutations result in very short telomeres: analysis of a large cohort of patients with dyskeratosis congenita and related bone marrow failure syndromes. *Blood* **112**, 3594–3600 (2008).
150. Savage, S. A. *et al.* TINF2, a component of the shelterin telomere protection complex, is mutated in dyskeratosis congenita. *Am. J. Hum. Genet.* **82**, 501–509 (2008).
151. Zhong, F. *et al.* Disruption of telomerase trafficking by *TCAB1* mutation causes dyskeratosis congenita. *Genes Dev.* **25**, 11–16 (2011).
152. Lee, J. W. Telomere shortening by mutations in the *RTEL1* helicase cause severe form of dyskeratosis congenita, Hoyerall-Hreidarsson syndrome. *Clin. Genet.* **84**, 210 (2013).
153. Walne, A. J., Vulliamy, T., Kirwan, M., Plagnol, V. & Dokal, I. Constitutional mutations in *RTEL1* cause severe dyskeratosis congenita. *Am. J. Hum. Genet.* **92**, 448–453 (2013).
154. Ballaw, B. J. *et al.* Germline mutations of regulator of telomere elongation helicase 1, *RTEL1*, in dyskeratosis congenita. *Hum. Genet.* **132**, 473–480 (2013).
155. Savage, S. A. Human telomeres and telomere biology disorders. *Progress Mol. Biol. Translat. Sci.* **125**, 41–66 (2014).
156. Greenberg, R. A. *et al.* Short dysfunctional telomeres impair tumorigenesis in the INK4a^{2/3} cancer-prone mouse. *Cell* **97**, 515–525 (1999).
157. Perera, S. A. *et al.* Telomere dysfunction promotes genome instability and metastatic potential in a K-ras p53 mouse model of lung cancer. *Carcinogenesis* **29**, 747–753 (2008).
158. Feldser, D. M. & Greider, C. W. Short telomeres limit tumor progression *in vivo* by inducing senescence. *Cancer Cell* **11**, 461–469 (2007).
159. Kaul, Z., Cesare, A. J., Huschtscha, L. I., Neumann, A. A. & Reddel, R. R. Five dysfunctional telomeres predict onset of senescence in human cells. *EMBO Rep.* **13**, 52–59 (2012).
160. Cesare, A. J., Hayashi, M. T., Crabbe, L. & Karlseder, J. The telomere deprotection response is functionally distinct from the genomic DNA damage response. *Mol. Cell* **51**, 141–155 (2013).
161. Capper, R. *et al.* The nature of telomere fusion and a definition of the critical telomere length in human cells. *Genes Dev.* **21**, 2495–2508 (2007).
162. Maciejowski, J., Li, Y., Bosco, N., Campbell, P. J. & de Lange, T. Chromothripsis and kataegis induced by telomere crisis. *Cell* **163**, 1641–1654 (2015). **This publication shows that dicentric chromosomes can persist through mitosis and are resolved 3–20 hours after anaphase by the cytoplasmic nuclease TREX1. This is thought to promote chromothripsis and kataegis.**
163. Jones, R. E. *et al.* Escape from telomere-driven crisis is DNA ligase III dependent. *Cell Rep.* **8**, 1063–1076 (2014).
164. Artandi, S. E. *et al.* Telomere dysfunction promotes non-reciprocal translocations and epithelial cancers in mice. *Nature* **406**, 641–645 (2000). **Using telomerase-deficient mice, the authors provide *in vivo* evidence that telomere erosion in a p53-null background promotes the development of epithelial tumours that display complex chromosomal rearrangements, including non-reciprocal translocations.**
165. Rudolph, K. L., Millard, M., Bosenberg, M. W. & DePinho, R. A. Telomere dysfunction and evolution of intestinal carcinoma in mice and humans. *Nat. Genet.* **28**, 155–159 (2001).
166. Ding, Z. *et al.* Telomerase reactivation following telomere dysfunction yields murine prostate tumors with bone metastases. *Cell* **148**, 896–907 (2012). **In this report, the authors show that reactivation of telomerase in cells with dysfunctional telomeres augments the metastatic potential of prostate tumours.**
167. Chin, K. *et al.* *In situ* analyses of genome instability in breast cancer. *Nat. Genet.* **36**, 984–988 (2004).
168. Jeon, H. S. *et al.* Telomere length of tumor tissues and survival in patients with early stage non-small cell lung cancer. *Mol. Carcinog.* **53**, 272–279 (2014).
169. Strefford, J. C. *et al.* Telomere length predicts progression and overall survival in chronic lymphocytic leukemia: data from the UK LRF CLL4 trial. *Leukemia* **29**, 2411–2414 (2015).
170. Kim, N. W. *et al.* Specific association of human telomerase activity with immortal cells and cancer. *Science* **266**, 2011–2015 (1994).
171. Hahn, W. C. *et al.* Inhibition of telomerase limits the growth of human cancer cells. *Nat. Med.* **5**, 1164–1170 (1999).
172. Horn, S. *et al.* TERT promoter mutations in familial and sporadic melanoma. *Science* **339**, 959–961 (2013). **In this report, the authors identify a germline mutation in the promoter of TERT, which creates a binding motif for ETS and TCF transcription factors.**
173. Huang, F. W. *et al.* Highly recurrent TERT promoter mutations in human melanoma. *Science* **339**, 957–959 (2013).
174. Heidenreich, B., Rachakonda, P. S., Hemminki, K. & Kumar, R. TERT promoter mutations in cancer development. *Curr. Opin. Genet. Dev.* **24**, 30–37 (2014).
175. Killela, P. J. *et al.* TERT promoter mutations occur frequently in gliomas and a subset of tumors derived from cells with low rates of self-renewal. *Proc. Natl Acad. Sci. USA* **110**, 6021–6026 (2013).
176. Kinde, I. *et al.* TERT promoter mutations occur early in urothelial neoplasia and are biomarkers of early disease and disease recurrence in urine. *Cancer Res.* **73**, 7162–7167 (2013).
177. Remke, M. *et al.* TERT promoter mutations are highly recurrent in SHH subgroup medulloblastoma. *Acta Neuropathol.* **126**, 917–929 (2013).
178. Quaas, A. *et al.* Frequency of TERT promoter mutations in primary tumors of the liver. *Virchows Arch.* **465**, 673–677 (2014).
179. Atala, A. Re: TERT promoter mutations and telomerase reactivation in urothelial cancer. *J. Urol.* **194**, 848–849 (2015).

180. Peifer, M. *et al.* Telomerase activation by genomic rearrangements in high-risk neuroblastoma. *Nature* **526**, 700–704 (2015).
181. Borah, S. *et al.* TERT promoter mutations and telomerase reactivation in urothelial cancer. *Science* **347**, 1006–1010 (2015).
182. Bell, R. J. *et al.* The transcription factor GABP selectively binds and activates the mutant TERT promoter in cancer. *Science* **348**, 1036–1039 (2015).
183. Chiba, K. *et al.* Cancer-associated TERT promoter mutations abrogate telomerase silencing. *eLife* **4**, e07918 (2015).
184. Quesada, V. *et al.* Exome sequencing identifies recurrent mutations of the splicing factor *SF3B1* gene in chronic lymphocytic leukemia. *Nat. Genet.* **44**, 47–52 (2012).
185. Ramsay, A. J. *et al.* *POT1* mutations cause telomere dysfunction in chronic lymphocytic leukemia. *Nat. Genet.* **45**, 526–530 (2013).
This sequence analysis of > 300 patients with chronic lymphocytic leukaemia identifies recurrent somatic mutations in *POT1*. The mutations mostly cluster in the DNA-binding domains of *POT1* and affect its telomere-protective function.
186. Robles-Espinoza, C. D. *et al.* *POT1* loss-of-function variants predispose to familial melanoma. *Nat. Genet.* **46**, 478–481 (2014).
187. Shi, J. *et al.* Rare missense variants in *POT1* predispose to familial cutaneous malignant melanoma. *Nat. Genet.* **46**, 482–486 (2014).
188. Bainbridge, M. N. *et al.* Germline mutations in shelterin complex genes are associated with familial glioma. *J. Natl Cancer Inst.* **107**, 384 (2015).
189. Kataoka, K. *et al.* Integrated molecular analysis of adult T cell leukemia/lymphoma. *Nat. Genet.* **47**, 1304–1315 (2015).
190. Landau, D. A. *et al.* Evolution and impact of subclonal mutations in chronic lymphocytic leukemia. *Cell* **152**, 714–726 (2013).
191. Calvete, O. *et al.* A mutation in the *POT1* gene is responsible for cardiac angiosarcoma in TP53-negative Li-Fraumeni-like families. *Nat. Commun.* **6**, 8383 (2015).
192. Aoude, L. G. *et al.* Nonsense mutations in the shelterin complex genes *ACD* and *TERF2IP* in familial melanoma. *J. Natl Cancer Inst.* **107**, dju408 (2015).
193. Hartmann, K. *et al.* Gene dosage reductions of *Trf1* and/or *Tin2* induce telomere DNA damage and lymphoma formation in aging mice. *Leukemia* **30**, 749–753 (2015).
194. Sfeir, A. & Symington, L. S. Microhomology-mediated end joining: a back-up survival mechanism or dedicated pathway? *Trends Biochem. Sci.* **40**, 701–714 (2015).
195. Deriano, L. & Roth, D. B. Modernizing the nonhomologous end-joining repertoire: alternative and classical NHEJ share the stage. *Annu. Rev. Genet.* **47**, 433–455 (2013).
196. Attwooll, C. L., Akpınar, M. & Petrini, J. H. The Mre11 complex and the response to dysfunctional telomeres. *Mol. Cell. Biol.* **29**, 5540–5551 (2009).
197. Dimitrova, N. & de Lange, T. Cell cycle-dependent role of MRN at dysfunctional telomeres: ATM signaling-dependent induction of nonhomologous end joining (NHEJ) in G1 and resection-mediated inhibition of NHEJ in G2. *Mol. Cell. Biol.* **29**, 5552–5563 (2009).
198. Deng, Y., Guo, X., Ferguson, D. O. & Chang, S. Multiple roles for MRE11 at uncapped telomeres. *Nature* **460**, 914–918 (2009).
199. Dimitrova, N. & de Lange, T. MDC1 accelerates nonhomologous end-joining of dysfunctional telomeres. *Genes Dev.* **20**, 3238–3243 (2006).
200. Botuyan, M. V. *et al.* Structural basis for the methylation state-specific recognition of histone H4-K20 by 53BP1 and Crb2 in DNA repair. *Cell* **127**, 1361–1373 (2006).

Acknowledgements

The authors apologize to the colleagues whose work they could not cite owing to space constraints. They thank A. Penev, S. K. Deng, Lynda Grocock, F. Lotterberger and D. Conomos for comments on the manuscript. Research in the authors' laboratories is supported by grants from the US National Institutes of Health (DP2CA195767 and DK102562 to A.S., and AG038677 to E.L.D.) and the American Cancer Society (RSG-14-186) to E.L.D. A.S. is a Pew-Stewart Scholar, a Damon Runyon-Rachleff grant recipient and a fellow of the David and Lucille Packard foundation.

Competing interests statement

The authors declare no competing interests.