

# The Role of Double-Strand Break Repair Pathways at Functional and Dysfunctional Telomeres

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Telomeres have evolved to protect the ends of linear chromosomes from the myriad of threats posed by the cellular DNA damage signaling and repair pathways. Mammalian telomeres have to block nonhomologous end joining (NHEJ), thus preventing chromosome fusions; they need to control homologous recombination (HR), which could change telomere lengths; they have to avoid activating the ATM (ataxia telangiectasia mutated) and ATR (ATM- and RAD3-related) kinase pathways, which could induce cell cycle arrest; and they have to protect chromosome ends from hyperresection. Recent studies of telomeres have provided insights into the mechanisms of NHEJ and HR, how these double-strand break (DSB) repair pathways can be thwarted, and how telomeres have co-opted DNA repair factors to help in the protection of chromosome ends. These aspects of telomere biology are reviewed here with particular emphasis on recombination, the main focus of this collection.

Mammalian telomeres are highly conserved in structure and function. They are built on long tandem arrays of duplex TTAGGG repeats that form the binding sites for the abundant telomere-specific protein complex, called shelterin (Fig. 1A). The telomeric repeat array ends in a 50- to 400-nt 3' protrusion of the G-rich strand. The presence of duplex telomeric repeats, a telomere-specific protein complex, and a 3' protrusion are general themes for all eukaryotic telomeres but the nature of the repeats and proteins vary widely. For instance, although a shelterin-like complex can be recognized in fission yeast and even trypanosomes, budding yeast telomeres function with a different set of proteins (Li et al. 2005; Lewis and Wuttke 2012). Another telomeric theme that is nearly universal is the mode of telomeric DNA

maintenance, which involves the specialized reverse transcriptase, telomerase. Telomerase uses the 3' end of the G-rich repeat strand as a primer and an internal RNA as a template to add telomeric repeats, thereby counteracting sequence loss resulting from semiconservative replication and nucleolytic processing (Blackburn and Collins 2011). Although there are organisms that maintain linear chromosomes without the help of telomerase (e.g., dipteran insects), they are rare. Mammalian cells can also maintain their telomeres in a telomerase-independent manner, using an homologous recombination (HR)-mediated process referred to as alternative lengthening of telomeres (ALT), which will be discussed in detail below.

Shelterin is the main mechanism by which telomeres solve the so-called end-protection

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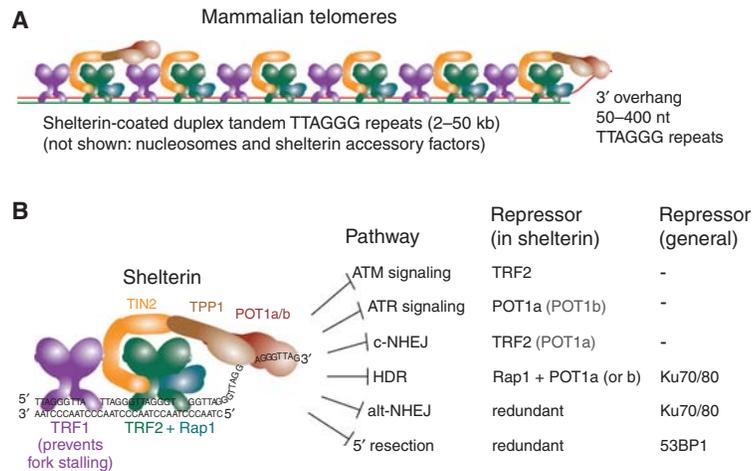
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**Figure 1.** Telomeres, shelterin, and the end-protection problem. (A) The structure of mammalian telomeres, including the telomeric double-stranded DNA, the telomeric 3' overhang, and the shelterin complex. (B) Schematic of the interactions among the six subunits that make up shelterin, their interactions with DNA, and their combined repression of the pathways that threaten telomeres (the end-protection problem).

problem. The end-protection problem refers to the collection of DNA damage response and repair pathways that can be deleterious at telomeres and therefore need to be repressed. Shelterin is composed of six structurally distinct proteins (Fig. 1B) (reviewed in Palm and de Lange 2008). TRF1 and TRF2 are diverged paralogs that have a similar domain structure, and both bind to duplex telomeric repeats as homodimers or -tetramers. Their large dimerization domains, called the TRF homology (TRFH) domains, also provide a binding surface for other shelterin components and shelterin-associated proteins. TRF1 and TRF2 are linked by TIN2, which helps to stabilize TRF1 and TRF2 on the telomeric DNA and, in addition, is crucial for the recruitment of two other shelterin proteins, TPP1 and POT1. POT1 binds to telomeric DNA in single-stranded form through an interaction of two OB-folds in the amino terminus with a 5'-(T)TAGGGTTAG-3' recognition site. Although POT1 binds better to this site at a 3' end, it will also associate with TTAGGG repeats when not located at a DNA end. Human cells have a single POT1 protein, whereas rodents have duplicated the POT1 gene, resulting in two functionally distinct but structurally closely related POT1 proteins at their telomeres, POT1a and POT1b.

The sixth structurally distinct shelterin component is Rap1, which interacts with TRF2 but not with any of the other shelterin proteins.

Shelterin is ubiquitously expressed and sufficiently abundant to cover all telomeric repeats, even at the exceedingly long (20- to 50-kb) telomeres of *Mus musculus*. There is also sufficient TPP1/POT1 to engage the single-stranded telomeric DNA, although TPP1/POT1 are at least 10-fold less abundant than TRF1, TRF2, TIN2, and Rap1 (Takai et al. 2010). What regulates the abundance of shelterin is not yet known.

The repression of DNA damage signaling by shelterin is primarily the role of TRF2 and POT1 (POT1a in the mouse) (Fig. 1B). TRF2 is required to prevent the activation of the ATM kinase at chromosome ends, and the absence of POT1 leads to robust signaling by the ATR kinase. These aspects of the end-protection problem have been reviewed elsewhere (Palm and de Lange 2008; de Lange 2009). Shelterin is also involved in the regulation of telomerase-dependent maintenance of telomeres, mediating both the recruitment of telomerase and controlling the homeostasis of telomere length (Smogorzewska and de Lange 2004; Nandakumar and Cech 2013). Below, we will briefly discuss how shelterin controls classical- and alter-

native nonhomologous end joining (NHEJ). The major part of this article will focus on how telomeres engage the cellular HR-related pathways—including 5' end resection—to achieve their optimal protected state, how telomeres prevent deleterious HR reactions, and what is known about the HR-dependent maintenance of telomeres in ALT cells.

## PHYSIOLOGICAL AND PATHOLOGICAL RESECTION AT TELOMERES

### Generating the Telomeric Overhang: Shelterin-Controlled 5' End Processing

The first step in the initiation of homologous recombination and all forms of homology-directed repair is the resection of the 5' ended strand at a DSB to generate a 3' overhang. 5' end resection also occurs at functional telomeres, but, interestingly, this process is very different from the processing of DSBs (for additional material on DSB end resection, see Symington [2014]). All mammalian telomeres have an overhang, implying that the telomeres formed by leading- and lagging-strand DNA synthesis (referred to as leading-end and lagging-end telomeres) acquire a 3' protrusion after DNA replication (Fig. 2) (Makarov et al. 1997; McElligott and Wellinger 1997). This overhang is longer at lagging-end telomeres, at least in telomerase-negative human cells (Chai et al. 2006; Chow et al. 2012). Because leading-strand DNA synthesis will not generate a 3' overhang and telomerase makes the overhangs longer but is not required for their presence (Hemann and Greider 1999; Chai et al. 2006; Wu et al. 2012), 5' end resection is the most likely explanation for the acquisition of the 3' overhangs at leading-end telomeres. In fact, it is now clear that 5' end resection takes place at both leading-end telomeres and lagging-end telomeres. The generation of the 3' overhangs has emerged as a multistep process that is closely controlled by shelterin.

The first indication that shelterin controls the formation of the 3' overhang came from partial (short hairpin RNA [shRNA]) knockdown studies of human POT1, which revealed a reduction in the telomeric overhang signal and a change in the sequence at the 5' end of telomeres.

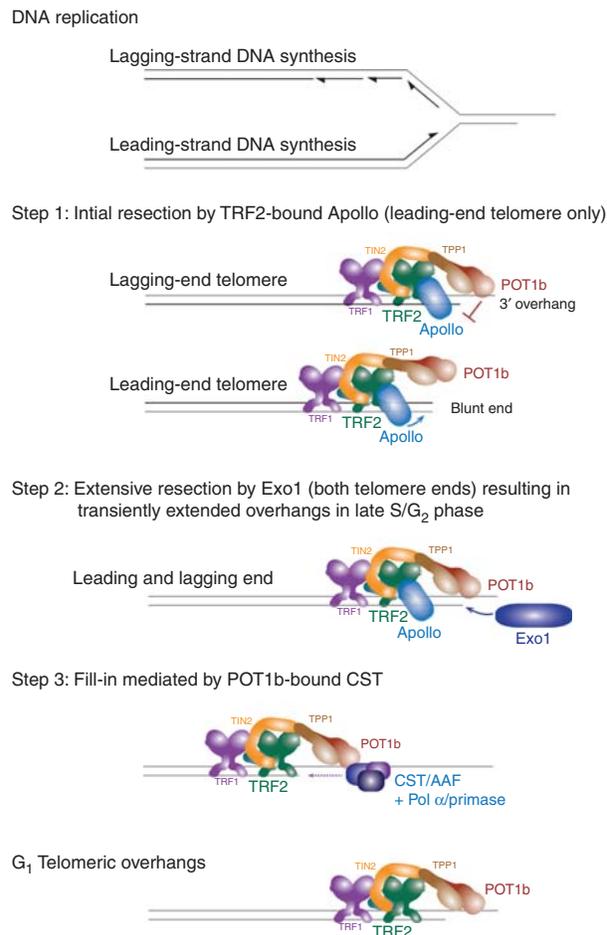
Remarkably, all human telomeres end on the sequence 3'-CCAATC-5' (Sfeir et al. 2005) but when POT1 is depleted, this terminal specificity is lost (Hockemeyer et al. 2005). However, the partial removal of POT1 from human telomeres also activates a DNA damage response (Hockemeyer et al. 2005), confounding the interpretation of the data. In the mouse, however, the repression of ATR kinase signaling is primarily the task of POT1a, whereas POT1b is dedicated to the control of the telomeric overhang, clarifying the interpretation of phenotypes obtained with deletion of POT1b (Hockemeyer et al. 2006).

Deletion of POT1b results in a 2- to 4-fold increase in the telomeric overhang signal (Hockemeyer et al. 2006). This effect accelerates telomere shortening when there is insufficient telomerase activity in the cells, consistent with the increase in single-stranded DNA being caused by exonucleolytic attack (Hockemeyer et al. 2008; He et al. 2009). The exact nature of the defect caused by loss of POT1b became clear in recent work with compound genetic mouse cells lacking two critical nucleases, Apollo and Exonuclease 1.

Apollo is an SMN1B/PSO2-type nuclease with a role in interstrand cross-link repair that binds to the TRFH domain of TRF2 (Freibaum and Counter 2006; Lenain et al. 2006; van Overbeek and de Lange 2006; Chen et al. 2008). At telomeres, the TRF2-bound Apollo is required for the generation of the 3' overhang at the leading ends (Lam et al. 2010; Wu et al. 2010, 2012). When Apollo is absent or cannot interact with TRF2, leading-end telomeres (separated on CsCl gradient [Wu et al. 2012]) show a diminished overhang signal, activate the ATM kinase, and fuse to other leading-end telomeres.

The resection of newly replicated telomeres by Apollo is controlled by POT1b (Wu et al. 2012). When POT1b is absent, hyperresection by Apollo occurs at both leading- and lagging-end telomeres. Presumably, POT1b will normally inhibit Apollo at lagging-end telomeres immediately after DNA replication, because they will have a short overhang to which POT1b can bind. At leading-end telomeres, POT1b is proposed to inhibit Apollo after the nuclease has generated a POT1b binding site (Fig. 2).

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**Figure 2.** Generation of the telomeric 3' overhang. Schematic of the three steps involved in the regeneration of the 3' overhang at the telomere replicated by leading- and lagging-strand DNA synthesis. See text for details. (Figure based on data from Wu et al. 2012.)

The second nuclease, EXO1, acts at both leading- and lagging-end telomeres, presumably after Apollo. EXO1 generates highly extended overhangs but their presence is transient (Wu et al. 2012). This transient extension of the overhangs in S phase has been observed in mouse and human cells (Dai et al. 2010; Wu et al. 2010, 2012). How EXO1 is recruited to telomeres in absence of a DNA damage response is not clear. At DSBs, EXO1 is dependent on RPA, which presumably is not available at functional/protected telomeres (Nimonkar et al. 2011; Cannavo et al. 2013; Chen et al. 2013). Perhaps TPP1/POT1a can provide this function.

Because the extended overhangs generated by EXO1 are returned to a shorter length by the time cells are in G<sub>1</sub>, it appears that a fill-in reaction is taking place. The candidate for the critical factor mediating this fill-in reaction, CST, emerged from work on budding yeast telomeres and polymerase  $\alpha$  (Pol  $\alpha$ )/primase accessory factor (reviewed in Price et al. 2010). Budding yeast telomeres are protected by a complex of Cdc13, Stn1, and Ten1 (CST), which prevents the activation of the ATR kinase homolog, Mec1. The RPA-like CST complex is not related to shelterin, although it uses OB-folds to bind to single-stranded DNA, as does POT1 (Gao et al.



2007; Lewis and Wuttke 2012). The mammalian ortholog of this complex was recognized by two groups as a complex of Ctc1/Stn1/Ten1 and also identified as the Pol  $\alpha$ /primase accessory factor AAF (Casteel et al. 2009; Miyake et al. 2009; Surovtseva et al. 2009; Wan et al. 2009). In human cells, CST interacts with TPP1 (Wan et al. 2009), and knockdown of Stn1 results in an increase in the telomeric overhang (Miyake et al. 2009; Surovtseva et al. 2009), although the cell cycle effects associated with Stn1 inhibition might be a confounder. In mouse cells, CST interacts with POT1b and mutations in the POT1b residues required for this interaction also lead to extended overhangs (Wu et al. 2012). These extended overhangs arise in late S/G<sub>2</sub> and fail to be shortened to their normal G<sub>1</sub> length. These data suggest that the POT1b-mediated recruitment of CST is needed for a fill-in step that returns the telomeric overhangs back to their normal lengths after EXO1 processing. Essentially, the same conclusion was reached on the necessity for CST-mediated fill-in at human telomeres (Huang et al. 2012; Wang et al. 2012; Kasbek et al. 2013). Given the ability of CST/AAF to promote Pol  $\alpha$ /primase activity on single-stranded DNA (Nakaoka et al. 2012), it is reasonable to assume that this is the way CST acts at telomeres (Fig. 2).

This highly regulated processing of the newly replicated telomere ends appears to be designed to ensure that both sister telomeres acquire an overhang. In addition, the tight regulation of telomere-end processing provides a way to control the rate of sequence loss at telomeres. Resection-dependent sequence loss ultimately determines the rate of telomere shortening in cells lacking telomerase. The gradual and progressive shortening of telomeres in the human soma is thought to represent a tumor suppressor mechanism that curbs the proliferative potential of incipient tumor cells. It may therefore be important for human somatic cells to control the rate of shortening by modulating the resection and fill-in steps.

The 3' overhang itself is important for several aspects of telomere function. It functions as the priming site of telomerase, which is incapable of acting on a blunt end, and the 3' overhang

is thought to be crucial for the protection of chromosome ends.

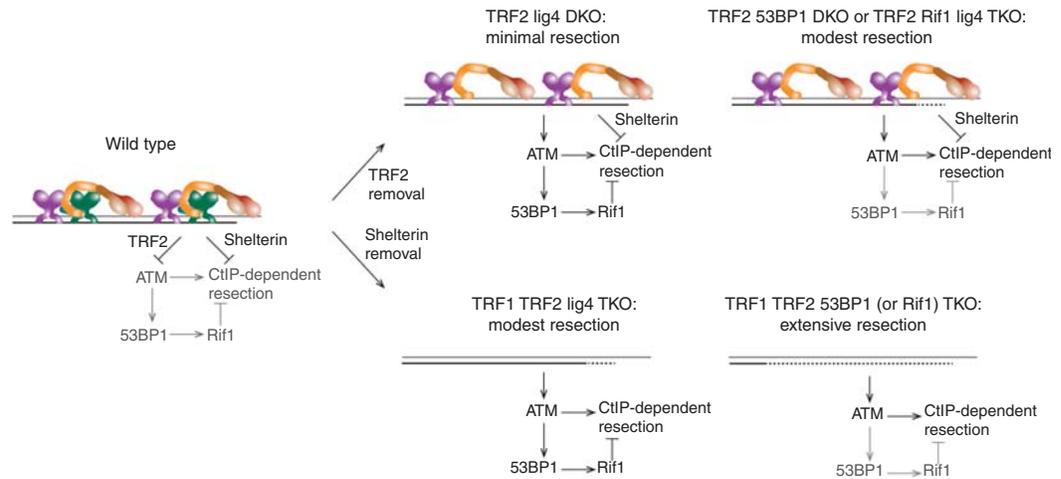
The best-understood role for the 3' overhang is mediating the formation of the T-loop structure, by strand invasion into the double-stranded telomeric repeats (see Fig. 4) (Griffith et al. 1999). T-loops have been proposed to provide an architectural solution to the end-protection problem by sequestering the end of the chromosome from end-initiated DNA damage response pathways. T-loop formation itself could be a mechanism to block hyperresection at telomeres similarly to what happens after strand invasion in homologous recombination (Sugawara et al. 2003).

### Hyperresection at Dysfunctional Telomeres

When POT1b is removed from telomeres, hyperresection of the newly replicated telomeres occurs, but this resection is distinct from resection at DSBs in that it is not directed by DNA damage signaling. ATM-dependent resection can take place at dysfunctional telomeres but only in cells that lack 53BP1, which has recently emerged as a general repressor of DSB resection in mammalian cells (Fig. 1B) (Bothmer et al. 2010; Bunting et al. 2010; Noon and Goodarzi 2011). When TRF2 is removed from telomeres in 53BP1<sup>-/-</sup> cells, telomere ends are processed in S phase by ATM- and CtIP-dependent resection (Fig. 3) (Lottersberger et al. 2013). Curiously, this resection only takes place at leading-end telomeres; the reason for this preference is not clear. The leading-end telomeres are supposedly blunt after replication and in TRF2-deficient cells they will not undergo their normal Apollo-dependent resection. Perhaps at these leading ends, which lack binding sites for POT1a and POT1b, CtIP-dependent resection can take place, whereas at the lagging-end telomeres POT1a and POT1b are protective.

TRF2 is not the only component within shelterin required to repress hyperresection at telomeres. Modest resection also occurs after TRF1 deletion from 53BP1<sup>-/-</sup> cells (Sfeir and de Lange 2012). However, when all shelterin proteins (and all shelterin-associated factors) are removed from telomeres, the hyperresection

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**Figure 3.** Repression of hyperresection at telomeres. ATM/CtIP-dependent resection at telomeres is repressed by several independent pathways. On the one hand, the 53BP1 binding partner RIF1 inhibits resection at telomeres that have become dysfunctional such that the ATM kinase pathway has been activated. On the other hand, resection is repressed by TRF2 in shelterin preventing the activation of the ATM kinase and protecting telomeres from moderate resection when 53BP1 is absent (*top left and right*). In addition, other components in shelterin repress resection so that extensive hyperresection only occurs in the absence of the complete shelterin complex (*bottom right*).

is dramatically increased, indicating that different shelterin subunits act redundantly in blocking nucleolytic attack on chromosome ends (Fig. 3) (Sfeir and de Lange 2012). RNAi knock-down experiments have implicated CtIP, EXO1, and the BLM RECQ helicase in this resection but other factors are not excluded. Thus, the hyperresection taking place at dysfunctional telomeres that activate the ATM kinase signaling pathway bears a strong resemblance to the resection of DSBs (see Symington 2014).

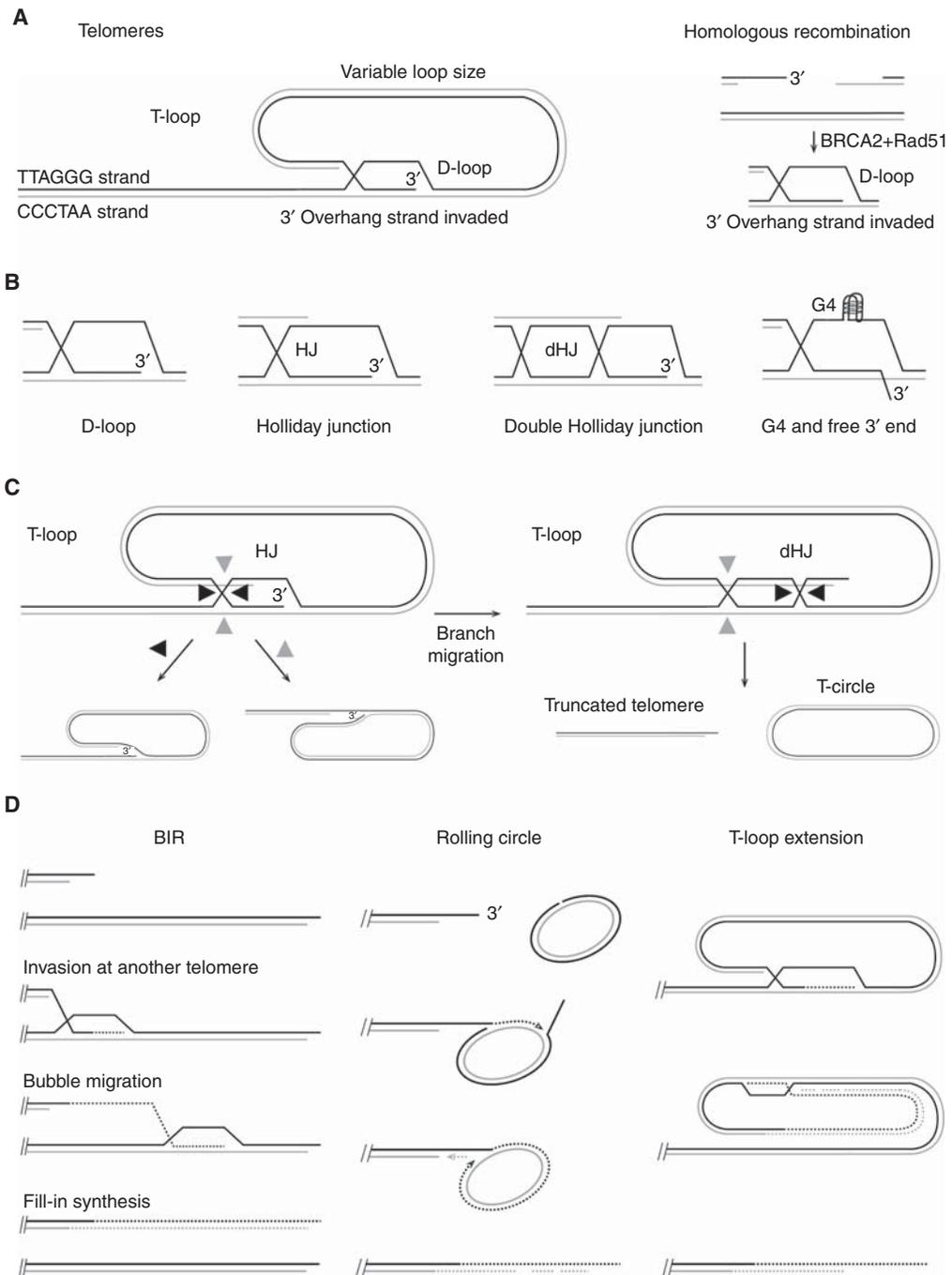
Recent data has revealed how 53BP1 represses resection at telomeres and DSBs (Chapman et al. 2013; Di Virgilio et al. 2013; Escibano-Diaz et al. 2013; Feng et al. 2013; Zimmermann et al. 2013). The main factor in this process is RIF1, a protein originally identified as a part of the telomeric protein complex in budding yeast (Hardy et al. 1992). In mammals, RIF1 is not found at telomeres but localizes to DSBs in an ATM- and 53BP1-dependent manner (Silverman et al. 2004). It was found that the amino-terminal S/TQ ATM target sites of 53BP1 are required for most of 53BP1's functions, including blocking resection in various contexts

(Bothmer et al. 2011; Lottersberger et al. 2013), and for the recruitment of RIF1 to sites of DNA damage (Chapman et al. 2013; Di Virgilio et al. 2013; Escibano-Diaz et al. 2013; Feng et al. 2013; Zimmermann et al. 2013). RIF1's role in repressing resection at telomeres was shown directly in the context of dysfunctional telomeres, thus identifying RIF1 as a factor downstream from 53BP1 in suppressing resection (Fig. 3) (Zimmermann et al. 2013). The same pathway was found to act at DSBs, further reinforcing the similarities between DSBs and dysfunctional telomeres (Chapman et al. 2013; Di Virgilio et al. 2013; Escibano-Diaz et al. 2013; Feng et al. 2013; Zimmermann et al. 2013).

## HOMOLOGY-DIRECTED REPAIR REACTIONS AT TELOMERES

### Formation of T-Loops and Their Function

In its simplest form, the T-loop structure resembles an early step in homologous recombination because the 3' overhang has strand-invaded homologous sequences, resulting in the formation of a D-loop (Fig. 4A). In homologous recom-



**Figure 4.** Homologous recombination (HR)-related structures invoked at telomeres. (A) The structure of the T-loop compared to the strand invasion step in HR (telomeric sequences written from 5' to 3'). (B) Possible structures at the base of the T-loop compatible with T-loop detection after psoralen cross-linking. (C) Products predicted from the indicated Holliday junction (HJ) resolution (cleavage and ligation) of the two T-loop structures shown. The products of the double Holliday junction (dHJ) schematic have been detected in cells overexpressing a mutant form of TRF2 lacking the amino-terminal basic domain (Wang et al. 2004). (D) Proposed mechanisms of telomere DNA synthesis in ALT cells. See text for discussion.

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bination, the strand invasion is mediated by BRCA2-loaded RAD51, and the invading strand would be initially coated by RAD51. Whether BRCA2 and RAD51 are also involved in T-loop formation has not yet been established. Recent experiments have indicated that TRF2 is the main component of shelterin required for the formation/maintenance of T-loops (Doksani et al. 2013), and recombinant TRF2 has the ability to promote a loop-forming reaction on telomere model substrates in vitro (Griffith et al. 1999; Stansel et al. 2001).

How might TRF2 promote looping in vitro? One possibility is based on the inherent ability of TRF2 to change the topology of the DNA it is bound to, which results in untwisting of the surrounding sequences and therefore can stimulate strand invasion (Amiard et al. 2007; Poulet et al. 2009). In addition, TRF2 has an amino-terminal domain, referred to as the basic domain, that binds Holliday junctions (HJs) in vitro and may therefore stabilize the strand-invasion event (Fouche et al. 2006; Poulet et al. 2009). These features, together with the propensity of TRF2 to initiate binding near the end of the telomeric repeat array in vitro (Stansel et al. 2001), may explain how TRF2 can form T-loop-like structures in vitro; however, whether these features are relevant to the in vivo situation is not yet known.

It is not unlikely that TRF2 acts in conjunction with other factors to establish the T-loop structure and to protect it (see also below). It is important to note that the exact nature of the structure at the base of the T-loop is not known (Fig. 4B gives four options but others are not excluded). Given that the basic domain of TRF2 can promote the formation of HJs and stabilize them (Fouche et al. 2006; Poulet et al. 2009), it is tempting to speculate that this is the most likely configuration (Fig. 4B, second option).

T-loops have been proposed to represent an architectural solution to the end-protection problem posed by linear chromosomes (Griffith et al. 1999). It appears likely that the telomere end, when in the T-loop configuration, would be impervious to end-loading factors that initiate DNA damage response pathways. Two prominent examples are the MRN (MRE11/

RAD50/NBS1)-initiated ATM kinase signaling pathway and KU70/80-dependent classical NHEJ. Neither MRN nor KU70/80 may be able to recognize the telomere for what it is, a DNA end, when the terminus is sequestered in the T-loop. Indeed, the shelterin component implicated in T-loop formation/maintenance, TRF2, also is critical for the repression of ATM signaling and classical NHEJ (de Lange 2009; Doksani et al. 2013).

It is generally assumed that T-loops are not a substrate for telomerase because the 3' end would be base-paired. However, given that the exact structure at the base of the T-loop is not known, it is not excluded that part of the 3' overhang is extruded and allows telomerase access (see Fig. 4B). Of course, it is also possible that telomerase gains access to the 3' terminus when T-loops are resolved during DNA replication.

### Protecting of T-Loops from Resolution and Other Steps in HR

Although the T-loop offers an architectural solution to many aspects of the end-protection problem, T-loops also create several challenges because of their structural resemblance to HR intermediates. The D-loop and possible HJ structures at the base of the loop are good substrates for nucleases and HJ resolution activities, such as MUS81/EME1, SLX4/SLX1, or GEN1 (Gaillard et al. 2003; Osman et al. 2003; Wyatt et al. 2013; Wyatt and West 2014). If a dHJ is generated by branch migration, its resolution can cleave off the loop part, generating a shortened telomere and a circular product often referred to as a T-circle (Fig. 4C, right). These products have been detected in cells induced to overexpress a TRF2 mutant lacking the amino-terminal basic domain (Wang et al. 2004). The interpretation of these findings is that the basic domain normally occupies the junction at the base of the T-loop and hence prevents HJ resolvases from gaining access and/or blocks branch migration. However, even without extensive branch migration, the D-loop itself or the single HJ intermediate will probably need protection because its resolution could give rise



to unwanted products when processed by nucleases and/or resolvases (Fig. 4C, left).

A second problem inherent to the T-loop structure is that the 3' end could be extended by polymerases, as it would be during HR (Mehta and Haber 2014). How this is prevented is not clear. It is also not clear what prevents the T-loop from forming on the sister telomere (in *trans*), or on chromosome-internal telomeric sequences (in *trans* or in *cis*) (e.g., Zhu et al. 2003). Finally, it has been argued that T-loops need to be resolved before the replication fork can progress through the telomeres and that the RTEL1 helicase is required for this resolution (Vannier et al. 2012). However, because the MCM replicative helicase is a 3' to 5' translocase (Fu et al. 2011), it should by itself be able to resolve the strand invasion that locks down the T-loop.

### Repression of HR between Sister Telomeres

Sequence exchanges between sister telomeres, referred to as telomere sister chromatid exchanges or T-SCEs, are harmless as long as they are equal. If an unequal exchange occurs, however, one daughter cell will inherit a shortened telomere, which will dictate a shorter replicative life span if the cell lacks a telomere maintenance system. Because each nucleus contains a large number of telomeres, sequence exchanges between sister telomeres have to be stringently repressed to provide the population as a whole with its proper proliferative capacity. Telomeres, representing a DNA end, are inherently recombinogenic. How is their recombination avoided? One obvious way is by forming a T-loop, which will sequester the telomere terminus; but genetic analysis shows that there are additional mechanisms at work. The KU70/80 heterodimer is an important repressor of HR at telomeres (Celli et al. 2006; Wang et al. 2009), analogous to its ability to repress HR at DSBs (Pierce et al. 2001). In addition, members of the shelterin complex repress the exchanges between sister telomeres. Notably, T-SCEs are only observed at substantial frequencies (>8% of telomeres) in two settings: in KU70/80-deficient mouse cells lacking either Rap1 or both

POT1a and POT1b (Fig. 1B) (Palm et al. 2009; Sfeir et al. 2010). Thus, the shelterin-dependent repression of HR involves the combined action of Rap1 and one of the POT1 proteins. How these proteins block HR is not clear. In the Rap1/KU70 double-knockout cells, it was notable that there is no DNA damage signaling at telomeres, yet they undergo homologous recombination (Sfeir et al. 2010). Perhaps the lack of requirement for DNA damage signaling in this setting is because of the presence of an overhang at the telomeres, circumventing the need for ATM kinase-dependent resection.

### Alternative Lengthening of the Telomeres

Telomere maintenance in the absence of telomerase was first described in budding yeast and shown to involve recombination (Lundblad and Blackburn 1993). Telomerase-independent telomere maintenance in human cells was first observed in mostly virally transformed cell populations that survived telomere crisis and became immortal (Bryan et al. 1995). Such immortal clones most often express telomerase, but a substantial number were found to maintain telomeres in the absence of detectable telomerase activity. Although first observed in cultured cells, it is now clear that the ALT pathway can sustain telomeres in a considerable subset of human cancers (reviewed in Henson and Reddel 2010).

ALT cells appear to use HR to maintain telomeric sequences. Their telomeres show an elevated rate of T-SCEs and a neo cassette embedded within one telomere was shown to spread to other telomeres (Dunham et al. 2000; Bechter et al. 2004; Londono-Vallejo et al. 2004). Furthermore, extrachromosomal telomeric DNA, including single-stranded and duplex circular telomeric DNA, is observed in ALT cells as if T-loops are more vulnerable to resolution by HJ resolvases (Tokutake et al. 1998; Cesare and Griffith 2004; Wang et al. 2004; Henson et al. 2009). ALT telomeres are also often heterogeneous in length (Bryan et al. 1995), consistent with increased recombination, and can show spreading of variant telomeric repeats through the telomeres (Varley et al. 2002; Conomos et al.

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2012). Finally, the telomeric DNA in ALT cells is often associated with PML bodies and a host of recombination and DNA damage response proteins (Yeager et al. 1999).

Despite these indications that HR is unleashed at ALT telomeres, the actual mechanism by which the telomeres are maintained and how ALT is activated is unknown. The single common denominator of all ALT cells is the presence of extrachromosomal telomeric DNA that is often circular and has been proposed to function as a template for telomere elongation (Fig. 4D) (Henson et al. 2002). Telomere elongation could also occur in *cis*, as has been directly shown by the local amplification of a cassette inserted in a telomere (Muntoni et al. 2009). T-loop extension is a likely mechanism of telomere elongation (see Fig. 4D). The idea that the telomere maintenance in these cells is dependent on HR is consistent with the finding that diminished activity of BRCA2, MUS81, FEN1, FANCD2, FANCA, SMC5/6-dependent sumoylation, RAD50, and other recombination factors affects telomere maintenance in ALT cells (reviewed in Gocha et al. 2013).

Telomere replication problems could play an important role in inducing recombination in ALT cells. First, ALT telomeres appear to have internal nicks or gaps that would cause frequent replication fork collapse at telomeres (Nabetani and Ishikawa 2009). Second, some of the recombination proteins required for telomere maintenance in ALT cells, including MUS81, FEN1, FANCD2, and FANCA, are known to play an important role in replication-coupled repair. Accumulation of broken replication forks at telomeres would be highly recombinogenic considering the repetitive nature of telomeric DNA and thus the abundance of homology for strand invasion. A broken end could invade the sister chromatid at different positions and generate equal or unequal T-SCEs. Strand invasion could also involve a telomere of another chromosome, and, in both cases, the invading end could be involved in break-induced replication (BIR) (Fig. 4D) (Donnianni and Symington 2013; Saini et al. 2013; Wilson et al. 2013; Mehta and Haber 2014).

The only consistent genetic defect so far identified in ALT is the loss of ATRX (Heaphy et al. 2011a,b; Lovejoy et al. 2012). ATRX is a SWI/SNF-related chromatin remodeling enzyme that was shown to deposit the histone variant H3.3 into telomeric chromatin in mouse embryonic stem cells (Goldberg et al. 2010; Wong et al. 2010). How the loss of ATRX or H3.3 could lead to a higher level of telomere recombination is entirely unclear at this stage (e.g., Clynes and Gibbons 2013; Conomos et al. 2013). It is likely that in addition to ATRX deletion, a second genetic or epigenetic change is required to unleash ALT. So far, no changes in shelterin have been found in ALT cells and the mechanism of ALT activation remains elusive.

## REPRESSION OF NHEJ

### Classical NHEJ

Telomeres are threatened by classical KU70/80- and DNA ligase IV-dependent NHEJ (Smogorzewska et al. 2002; Celli and de Lange 2005; Celli et al. 2006). This type of DSB processing of dysfunctional telomeres takes place primarily in G<sub>1</sub>, resulting in chromosome-type fusions (the telomeres of both chromatids are fused to other telomeres) in metaphase (Konishi and de Lange 2008). However, postreplicative DNA ligase IV-dependent telomere fusions, which typically give rise to chromatid-type fusions (one chromatid fused to another) or sister telomere fusions, have also been observed (Smogorzewska et al. 2002; Dimitrova and de Lange 2009; Hsiao and Smith 2009). TRF2 is the main component of shelterin involved in blocking classical NHEJ. The mechanism by which TRF2 acts most likely involves the formation of the T-loop configuration. However, other aspects of TRF2 may be important as well because TRF2 has been reported to repress DNA repair when tethered to a nontelomeric site next to an inducible DSB, a context in which T-loop formation is impossible (Fumagalli et al. 2012). One way TRF2 may be acting in this context is through the action of its recently discovered IDDR domain (Okamoto et al. 2013). This



small region in the TRF2 linker between the TRFH domain and the Myb/SANT DNA binding domain interacts with both the MRN complex and the BRCC36 deubiquitylating enzyme, and through these two modules, it inhibits the loading of 53BP1 at sites of DNA damage. In the absence of 53BP1, classical NHEJ could be thwarted both at telomeres and at other sites through induction of excessive resection.

In addition to TRF2, the engagement of the POT1 proteins on the single-stranded DNA may be acting as a deterrent to classical NHEJ. Deletion of POT1a results in a low but significant level of fusions between sister telomeres that is exacerbated when POT1b is deleted as well (Hockemeyer et al. 2006). However, it has not been established whether these are classical NHEJ events. The idea that the 3' overhang (with POT1 proteins engaged) could block the loading of KU70/80 was also invoked to explain the paradoxical finding on TRF2 removal from telomeres in ATM-deficient cells. It was found that leading-end telomeres that had lost TRF2 were protected from NHEJ by ATM-mediated resection (Attwooll et al. 2009; Dimitrova and de Lange 2009; Lotterberger et al. 2013). Whether the 3' overhang can really protect telomeres from NHEJ and, if so, how long the overhang needs to be and whether it has to be coated with POT1 proteins to block NHEJ remain to be determined.

### Alternative NHEJ

The final DSB repair reaction that threatens telomeres is fusion through alternative NHEJ (alt-NHEJ). Hints that such alt-NHEJ might promote telomere fusions were seen in experiments in which multiple components of shelterin were targeted in KU-deficient mouse cells (Rai et al. 2010). Further insight into the type of alt-NHEJ taking place at dysfunctional telomeres recently came from experiments in which telomeres were rendered completely free of all shelterin proteins (Sfeir and de Lange 2012). When shelterin is removed from telomeres in KU-deficient cells, the telomere fusions are extremely prominent, involving up to 65% of the chromosome ends. These fusions are medi-

ated by DNA ligase III and PARP1 and appear to rely on the microhomology provided by the TTAGGG repeats in the 3' overhangs, which can form two base pairs per repeat. Similar to what happens at DSBs, alt-NHEJ at telomeres is repressed by KU70/80 (Wang et al. 2006). Within shelterin, multiple proteins contribute to the repression of alt-NHEJ, because the individual deletion of shelterin components in a KU-deficient background does not unleash the same dramatic alt-NHEJ phenotype as the complete removal of shelterin.

### CONCLUDING REMARKS

The connection between telomere biology and DSB repair pathways is akin to the link between immunology and infectious diseases in which the study of one aspect of biology informs the other and vice versa. It is now clear that mammalian telomeres are threatened by each of the processing and repair pathways that can act on DSBs. In each case, shelterin plays a critical role in repressing these pathways, but much remains to be learned about how this complex acts. Future insights into the biochemical and structural aspects of shelterin will be helpful as will be additional information on how exactly these dangerous pathways are initiated on a substrate like an unprotected telomere. Another question that merits attention is how telomeres use DSB repair factors for protective purposes. An example is how shelterin manages to use EXO1 and Apollo to generate the 3' overhang while avoiding hyperresection. Further insights into these types of questions will not only increase the understanding of telomeres but also further illuminate how DSB repair works.

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## The Role of Double-Strand Break Repair Pathways at Functional and Dysfunctional Telomeres

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