When transcription meets recombination: a lesson from the human RECQ protein complexes

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Abstract

Since the cloning of the first human RECQ gene, RECQ1, more than 15 years ago, RECQ helicases have been a major focus in cancer research. Recent studies of human RECQ protein complexes are providing insight into their roles in various DNA metabolic pathways that protect the integrity of our genome.

Introduction and context

The highly conserved RecQ helicase was first identified in Escherichia coli about 25 years ago and was subsequently found to be required for suppressing illegitimate recombination in both E. coli and yeast [1-3]. Ten years later, discovery of the link between the human homolog, BLM, and Bloom syndrome demonstrated the importance of the RecQ family helicases in human health [4]. Bloom syndrome is a devastating genetic disorder that causes cancer predisposition. Cells derived from Bloom syndrome patients show high levels of sister chromatid exchange resulting from DNA recombination and genome rearrangements reminiscent of those found in lower organisms [5].

Importantly, BLM is not the only RECQ homolog to be identified in humans. During evolution, the RecQ gene has been amplified and has diverged from a single copy in bacteria to five RECQ homologs in humans, namely, RECQ1, BLM, WRN, RECQ4, and RECQ5. Importantly, although there are multiple REC helicases in humans, they are not redundant, as mutations in different RECQ helicases are responsible for distinct clinical diseases [6]. Bloom syndrome, Werner syndrome, and Rothmund-Thomson syndrome have been linked to defects in BLM, WRN, and RECQ4, respectively. In the past decade, much research has been dedicated to uncovering molecular functions of RECQ family helicases linked with different clinical disorders.

Major recent advances

Among the five human RECQ helicases, BLM has been the most extensively studied. BLM is found in a complex with topoisomerase IIIα (TOPOIIIα) and the RecQ-mediated genome instability (RMI) proteins RMI1 and RMI2. Cumulative data indicate that BLM makes use of four possible mechanisms to repair DNA breaks by both promoting and suppressing homologous recombination (Figure 1) [7]. First, BLM, together with TOPOIIIα and RMI1/2, dissolves double Holliday junction intermediates into non-crossover products [8]. Second, BLM may contribute to the synthesis-dependent single-stranded annealing pathway, leading exclusively to products without sister chromatid exchange [9]. Most recently, BLM has been implicated in two opposite roles: disassembling the RAD51 filament from the break site to prevent homologous recombination; and facilitating end resection to promote homologous recombination [10-12].

While it is clear that BLM is an essential player in safeguarding the progression and influencing the outcome of homologous recombination events, how these pro- and anti-recombinational roles of BLM are coordinated during
homologous recombination is still a puzzle. Further complicating the story, it had been thought that elevated levels of sister chromatid exchange accompanied by chromosome rearrangement are a hallmark of Bloom syndrome cells and a key feature in the clinical diagnosis of Bloom syndrome, however, recent studies have shown that mouse embryonic stem cells defective in RECQ5 exhibit a phenotype similar to that of Bloom syndrome cells [13]. Furthermore, even though RECQ5 mutations are not linked to Bloom syndrome or any other clinical disease in humans, RECQ5 deletion in mice results in a significant increase in the development of solid tumors in various organs such as the lung, liver, and breast [14]. Notably, the recq5-/- blm-/- double mutant exhibited an even greater frequency of sister chromatid exchange than either of the single mutants, suggesting that these two proteins contribute to regulation of sister chromatid exchange via different mechanisms [13].

A recombination event is initiated by a DNA break and leads to crossover or non-crossover products. First, the DNA ends are resected to produce single-stranded DNA, which in a reaction catalyzed by RAD51 recombinase can then invade the homologous duplex DNA to form a recombination intermediate, or D-loop structure. After homologous recombination is initiated by RAD51 at one end of the broken DNA, the second DNA end may be captured by the unpaired strand of the homologous DNA. The resulting recombination intermediate can proceed to form double Holliday junctions (HJs), and any remaining gaps may be filled by DNA synthesis. HJs can be cleaved by HJ resolvase (center right) or dissociated by the BLM-dependent pathway (center left). In mitotic cells, it is argued that the majority of homologous recombination is completed without the formation of HJs. Instead, most homologous recombination events are completed by synthesis-dependent strand annealing (SDSA) to generate non-crossover products that result from dissociation of the D-loop after DNA synthesis (left). The newly synthesized DNA is available to re-anneal with the second end of the same DNA molecule at the break site. DNA synthesis continues to fill the gap and restore the integrity of the DNA without crossover. Alternatively, D-loop structure can be cleaved to generate sister chromatid exchange (SCE) products (right). Possible points of RECQ5 action discussed in the text are indicated by (1), (2), and (3).
While RECQ5 does not dissolve double Holliday junctions [15], it is possible that it may function to promote synthesis-dependent single-strand annealing (Figure 1, mechanism 1). Alternatively, it has been shown that RECQ5 has an antagonistic role in preventing RAD51 filament formation (Figure 1, mechanism 2) [14]. However, both of these RAD51-dependent mechanisms are redundant to BLM function and are not likely to represent the primary role of RECQ5 in suppressing sister chromatid exchanges. Indeed, a recent report demonstrated an additive sensitivity to camptothecin treatment in recq5-rad51 double mutants [16], arguing for a role of RECQ5 in a RAD51-independent pathway.

A recent surprise in the field came with the purification from human cells of a multiprotein complex containing the RECQ5 helicase. Unlike BLM, which forms a stable complex with TOPOIIIα and RMI1/2 to function as a ‘dissolvasome’ in homologous recombination [7], RECQ5 primarily interacts with RNA polymerase II, the enzyme responsible for messenger RNA synthesis in eukaryotes (Figure 2) [17,18]. Importantly, this interaction is specific to RECQ5 but not to other RECQ helicases [17]. Further biochemical analysis revealed that RECQ5 efficiently suppresses RNA polymerase II-dependent transcription via direct protein-protein interaction [19]. Together, these data suggest a unique role of RECQ5 in transcription.

**Future directions**

The biochemical activities of the BLM-TOPOIIIα complex are clearly consistent with a role in homologous recombination pathways leading exclusively to formation of non-sister chromatid exchange products (Figure 1). On the other hand, many questions remain regarding the physiological connection between the RECQ5-RNA polymerase II interaction and sister chromatid exchange. Nonetheless, the potential link between transcription and recombination may not be unexpected. While transcription is essential for cells to function and grow, the process itself can generate undesirable DNA lesions and result in genome instability [20]. Indeed, mutations in RNA processing factors, such as ASF/SF2 and THO/TREX, result in double-stranded break formation and transcription-associated recombination [21,22]. It is very reasonable, therefore, to expect that RECQ5 could have a unique function in preventing DNA break formation during transcription (Figure 1, mechanism 3). According to this hypothesis, transcription-dependent double-stranded breaks would be generated in the absence of RECQ5 and subsequently repaired by homologous recombination, leading to an increase in sister chromatid exchange. It is worth noting that transcription-associated recombination was also observed in yeast [20], yet Sgs1, the only RecQ homolog in yeast, has not been implicated in this process. Further, Sgs1 is not known to interact with RNA polymerase II. Thus, it is possible that a role for RECQ-like helicases in transcription-associated recombination has been acquired during the course of evolution and is unique to higher eukaryotes.

Given that the key to solving the puzzle regarding RECQ5 and BLM may lie within the distinct protein complexes they form in cells, the identification of other RECQ helicase complexes may provide important clues to the unique functions of the remaining RECQ helicases in DNA metabolism. Such discoveries would allow us to start to understand how the human RECQ helicases associate with different clinical syndromes that have specific cancer spectra.

**Abbreviations**

RMI, RecQ-mediated genome instability; TOPOIIIα, topoisomerase IIIα.

**Competing interests**

The authors declare that they have no competing interests.

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