Loss of the bloom syndrome helicase increases DNA ligase 4-independent genome rearrangements and tumorigenesis in aging *Drosophila*

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**Abstract**

**Background:** The BLM DNA helicase plays a vital role in maintaining genome stability. Mutations in *BLM* cause Bloom syndrome, a rare disorder associated with cancer predisposition and premature aging. Humans and mice with *blm* mutations have increased frequencies of spontaneous mutagenesis, but the molecular basis of this increase is not well understood. In addition, the effect of aging on spontaneous mutagenesis in *blm* mutants has not been characterized. To address this, we used a *lacZ* reporter system in wild-type and several mutant strains of *Drosophila melanogaster* to analyze mechanisms of mutagenesis throughout their lifespan.

**Results:** Our data show that *Drosophila* lacking BLM have an elevated frequency of spontaneous genome rearrangements that increases with age. Although in normal flies most genome rearrangements occur through DNA ligase 4-dependent classical end joining, most rearrangements that accumulate during aging in *blm* mutants do not require DNA ligase 4, suggesting the influence of an alternative end-joining mechanism. Adult *blm* mutants also display reduced lifespan and ligase 4-independent enhanced tumorigenesis in mitotically active tissues.

**Conclusions:** These results suggest that *Drosophila* BLM suppresses error-prone alternative end-joining repair of DNA double-strand breaks that can result in genome instability and tumor formation during aging. In addition, since loss of BLM significantly affects lifespan and tumorigenesis, the data provide a link between error-prone end joining, genome rearrangements, and tumor formation in a model metazoan.

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**Background**

Bloom syndrome is a rare, autosomal recessive disorder whose most striking characteristic is a predisposition to all types of cancers (reviewed in [1,2]). It is caused by mutations in the *BLM* gene, a member of the RecQ family of DNA helicases [3]. Cells derived from Bloom syndrome patients and *Blm* hypomorphic mice exhibit greatly elevated genome instability, including a dramatically increased frequency of sister chromatid exchanges [4,5].

Cells with defective BLM also have a heightened mutation frequency that is partially independent of the increase in sister chromatid exchanges [6-9]. The cause of this is poorly understood, but it may be related to the increased number of chromosome aberrations and translocations that are observed in *BLM* mutant cells. Because these types of mutations are hypothesized to be driving forces in the development and progression of cancer in Bloom syndrome patients, understanding their origin is important.

The *mus309* gene encodes the *Drosophila melanogaster* BLM ortholog DmBlm [10]. Flies lacking DmBlm phenocopy many of the characteristics of human Bloom syndrome, including reduced fertility and increased...
mitotic sister chromatid exchanges [10-12]. DmBlm is required for accurate homologous recombination (HR) repair of site-specific DNA double-strand breaks (DSBs) and for prevention of mitotic crossovers [13,14]. In its absence, DSB repair frequently proceeds through non-conservative, deletion-prone repair mechanisms. Although initiation of homologous recombination is required for the deletions observed in blm mutants [15], the molecular mechanisms responsible for deletion formation remain unknown.

Here, we report that spontaneous mutagenesis at a lacZ reporter locus is increased in Drosophila blm mutants, similar to what is observed in human Bloom syndrome patients. The increase persists through the adult lifespan and is largely due to elevated genome rearrangements, almost 50% of which involve repetitive genomic regions. Interestingly, although spontaneous rearrangements in wild-type flies depend largely on DNA ligase 4 (Lig4)-dependent classical end joining, rearrangements in blm mutants are mostly Lig4-independent. Flies lacking DmBlm, but not Lig4, also have increased tumorigenesis, suggesting that, in the absence of Blm, alternative end joining may be involved in the formation of genome rearrangements that can drive tumorigenesis.

Results and discussion

Genome rearrangements involving repeated sequences are common in blm mutants

To characterize the consequences of DmBlm deficiency on spontaneous mutagenesis, we crossed a null mus309 mutation [12] into a w^1118 background harboring a lacZ reporter construct that allows us to measure somatic mutation frequencies and mutation spectra [16] (Additional file 1). For these studies, we used a lacZ reporter integrated at cytological position 87E on chromosome 3, which exhibits a spontaneous mutation frequency nearly identical to the average obtained using multiple other insertion sites [17]. The blm mutant stocks containing the reporter construct were isogenized and homozygous mutants were recovered at days 1, 14, and 28 after adult eclosion. Plasmid rescue was performed and mutation frequencies were calculated on a per locus basis as the number of mutant lacZ copies versus the total number of copies rescued from a given amount of DNA (Additional file 2).

We observed a significantly higher mutation frequency for blm mutants relative to wild-type at all ages (Figure 1a; P < 0.001 for both sexes, one-way ANOVA). As found previously, mutation frequencies were slightly higher in females than in males [17]. This difference is largely due to an increased frequency of point mutations in females (see below). Since point mutations often result from replication errors, females, which have a larger body size, may have more opportunities to accumulate point mutations.

The increased mutation frequency in blm mutants was not specific to a particular region of the fly, but was observed in head, thorax, and abdominal tissues (Figure 1b). Although the differences in tissue-specific mutation frequencies between blm mutant and wild-type males
were less significant compared to those observed using whole flies, this is likely a consequence of a higher mutation frequency in the wild-type males in this particular experiment (data not shown).

To examine the types of mutations that occurred in blm mutants, we performed restriction analysis on a subset of plasmids. Plasmids that show no size change after digestion with the Aval restriction enzyme generally harbor point mutations, while plasmids showing a size change after digestion involve rearrangements with one breakpoint in the lacZ gene and the other elsewhere in the fly genome. We have previously shown that genome rearrangements, including deletions and both intra- and inter-chromosomal translocations, are the predominant type of mutation in flies [17]. Interestingly, the increased mutation accumulation in blm mutants appeared to be almost entirely due to an increase in genome rearrangements (Figure 1c). This increase was evident in both males and females ($P < 0.001$, Fisher’s exact test).

To further characterize the genome rearrangements recovered from the blm mutants, we sequenced the lacZ gene in a representative number of mutants and identified the breakpoints. Using BLAST alignment of the recovered sequence against the Drosophila genome, we definitively mapped the breakpoints for 11 out of 21 rearrangements (Additional file 3). Three rearrangements were deletions in the lacZ transgene, while eight were larger deletions or inversions with the breakpoints outside the reporter gene in the same chromosome. Surprisingly, we were unable to map the breakpoints for 48% of the rearrangements isolated from blm mutants because one side of the breakpoint was located in highly repetitive sequences. Six of these sequences corresponded to various transposons, including accord/818, stalker, and copia, while four matched repetitive sequences that are highly represented in both euchromatic and heterochromatic regions of the fly genome. Previous studies using the same lacZ reporter found a much lower proportion of genome rearrangements involving repetitive sequences (6 out of 55, or 11%, $P = 0.0003$ compared to blm mutants, two-tailed Fisher’s exact test) [17,18]. Therefore, the increased somatic mutation frequency in blm mutants is due, at least in part, to an increased frequency of genome rearrangements involving highly repetitive sequences.

Differential requirements for rearrangement formation in wild type and blm mutants

The simplest model to explain the observed genome rearrangements involves inaccurate repair of two simultaneous DSBs. It is possible that recombination between two breaks could result in deletions and translocations and that DmBlm could suppress these types of rearrangements by preventing homeologous recombination between closely related sequences [19]. However, careful inspection of the junction sequences at the sites of the genome rearrangements isolated from wild type [18] and blm mutants did not reveal extensive regions of homology. To determine if classical non-homologous end joining (C-NHEJ) might be required for the formation of these rearrangements, we repeated our analysis in flies lacking DNA ligase 4, which is required for C-NHEJ [20,21]. Surprisingly, we saw no age-dependent increase in mutation frequency in lig4 mutants (Figure 1a). In addition, the percentage of genome rearrangements in lig4 mutants was significantly less compared to wild-type flies (Figure 1c; $P \leq 0.01$ for both sexes at all ages, two-tailed Fisher’s exact test), indicating that Lig4-dependent C-NHEJ is responsible for most of the genome rearrangements that accumulate during normal development and aging.

Previous studies from our lab and others have demonstrated that repair of site-specific DSBs in blm mutants is frequently accompanied by large deletions [14,15], a feature characteristic of Lig4-independent alternative end joining [22,23]. We therefore wished to determine if the genome rearrangements that occur in blm mutants rely on classical or alternative end joining. We constructed lig4 blm double mutants containing the lacZ reporter construct and measured the mutation frequency and spectra in 28-day-old flies. Spontaneous mutagenesis frequencies for the lig4 blm flies were similar to blm single mutants, but were significantly different from lig4 mutants (Figure 2a; $P = 0.029$ for males, $P < 0.001$ for females, one-way ANOVA). Similarly, the majority of mutations in lig4 blm flies were genome rearrangements, paralleling the blm mutant phenotype (Figure 2b). In contrast to wild-type flies, therefore, most of the genome rearrangements that occur in flies lacking the DmBlm helicase do not require DNA ligase 4 and are instead generated through an alternative end-joining mechanism.

Increased tumor formation and reduced lifespan in blm mutant flies

One hallmark of Bloom syndrome is cancer predisposition at an early age [24]. Recently, several groups have reported that adult flies develop dysplasia and tumors in mitotically active tissues, including intestinal and germline tissues [25,26]. These tumors arise more frequently in older flies [27]. We reasoned that the increased frequency of genome rearrangements observed in the absence of DmBlm might promote tumorigenesis. To test this, we performed histopathological staining on sectioned tissues of wild-type and blm mutant flies at 35, 50, and 60 days of age. Representative histological
sections from 35-day-old control and age-matched blm mutant flies are shown in Figure 3. Normal fly tissues (Figure 3a, c, e) are characterized by a high degree of cytological regularity and orderliness of tissue architecture. In contrast, tumors (Figure 3b, d, f) are composed of densely crowded masses of irregularly arrayed tumor cells. Although no metastases to distant tissues were observed, the gut tumors were frequently observed to partially or fully occlude the intestinal lumen. Germline tumors consisted of masses of proliferating tumor cells that were morphologically similar to stem cells. Like many mammalian tumor cells, individual fly tumor cells were variable in size and shape and showed high nucleus to cytoplasm ratios.

Similar to Bloom syndrome patients, flies lacking DmBlm had an increased frequency of tumor development at early ages. We observed a significant difference in overall tumor formation between 35-day-old wild-

Figure 2 Increased frequency of genome rearrangements in flies lacking DmBlm is independent of classical non-homologous end joining (a) Somatic mutation frequencies in 28-day-old flies, grouped by sex. Each bar represents four to six independent measurements. Error bars represent standard deviations. *P < 0.05, ***P < 0.001 compared to wild type (one-way ANOVA). (b) Somatic mutation frequencies that involved either a size change or no size change, grouped by sex.

Figure 3 blm mutant adults are predisposed to early tumor development (a) Transverse section through midgut of wild-type fly. Normal, uniform-appearing villi (V) protrude into the gut lumen (L). A cluster of bacteria and yeasts (normal gut flora) is present within the lumen. (b) Tumor within the midgut of a blm mutant fly. A dense cluster of small tumor cells (T) arises from within the mucosa and forms an irregular mass that protrudes into the gut lumen. (c) Normal testis from a wild-type fly. Normal spermatocytes (S) in various stages of development are present. (d) Tumor within the testis of a blm mutant fly. Large numbers of small tumor cells (T) resembling early germ line precursors have replaced normal spermatocytes within the testis. (e) Normal maturing ovarian follicles from a wild-type fly. The largest (arrowhead) contains normal cyst cells surrounded by a single layer of follicular cells. (f) Ovarian tumor from a blm mutant fly. Large numbers of tumor cells (T) resembling immature germ line precursor cells replace normal cystocytes. All sections were prepared from 35-day-old flies and were stained with hematoxylin and eosin. Each scale bar = 50 microns. (g) Histogram showing tumor frequency in 35- to 60-day-old adult flies. *P < 0.05 compared to wild type (WT; two-tailed Fisher’s exact test), **P < 0.01.
type and blm mutants (Figure 3g; $P = 0.05$ and $P = 0.0087$ for males and females, respectively). Older males also had increased tumorigenesis relative to wild type ($P = 0.0047$ at 50 days), but tumor frequency was similar in older wild-type and blm mutant females. Germline tumors, which are rare in wild-type flies, comprised a significant fraction of tumors in blm mutants, particularly in males. This could indicate that certain cell populations within the testes of blm males may be more prone to neoplastic transformation. Interestingly, there was no significant difference in tumor frequency between wild type and lig4 mutants (Additional file 4), suggesting that increased tumorigenesis is not a result of loss of DSB repair capacity in general. Overall, these results agree with previous findings that tumors accumulate with age in Drosophila [27] and indicate that tumorigenesis in flies can be influenced by the status of tumor suppressor genes like BLM.

In Drosophila, DSB repair relies more heavily on HR as flies age [28]. Because DmBlm is important for successful completion of HR and plays an important role in spontaneous genome rearrangements, we hypothesized that it might also act to promote longevity. To test this, we performed lifespan analysis on isogenic flies age [28]. Because DmBlm is important for successful completion of HR and plays an important role in spontaneous genome rearrangements, we hypothesized that it might also act to promote longevity. To test this, we performed lifespan analysis on isogenic wild-type, lig4, and blm mutant males and females. Loss of DmBlm significantly shortened mean and maximum lifespan in both sexes ($P < 0.0001$, Wilcoxon rank-sum test; Figure 4). The survivorship plots of the blm mutants diverged from the wild-type plots at approximately 20 days, and the mean lifespan of blm mutants was reduced more than 35% relative to wild type. In contrast, loss of Lig4 had no significant effect on the mean or maximum lifespan of female flies ($P = 0.83$). The survivorship plot of the lig4 mutant males closely paralleled that of wild-type flies until approximately 45 days, at which point it diverged, intersecting with the blm curve near the maximum lifespan of both mutants.

We also quantified tumor formation and lifespan in the double mutants. For both males and females, tumor frequency in lig4 blm mutants was similar to levels observed in blm single mutants (Additional file 4), matching the spontaneous mutagenesis results (Figure 2). Interestingly, the mean and maximum lifespan of both males and females was further decreased in lig4 blm flies relative to blm single mutants (Additional file 5). Thus, C-NHEJ repair of DNA DSBs becomes important for organism lifespan when HR repair is compromised.

Conclusions

Taken together, our results indicate that genome instability in Drosophila, specifically in the form of chromosome rearrangements, is affected to a greater extent by impairment of HR repair than by loss of C-NHEJ. Unlike genome rearrangements that occur during aging in wild-type flies, most rearrangements in blm mutants are Lig4-independent. Because blm mutants are proficient in the initial resection and strand invasion steps of HR [15], we hypothesize that DSB repair in the absence of DmBlm is more likely to proceed by alternative end-joining mechanisms that involve extensive resection at DNA ends. This leads to an increase in genome rearrangements and may predispose to tumorigenesis and reduced longevity. Interestingly, recent findings demonstrate that alternative end joining also plays a dominant role in translocation formation in mammals [29,30]. In combination with these observations, our study raises the interesting possibility that inaccurate end-joining repair may be a primary cause of pathology in Bloom syndrome and may also be a critical factor in aging.

Materials and methods

Drosophila stocks and genetics

Flies were raised on standard cornmeal agar medium. All stocks contained a lacZ reporter construct integrated
at chromosome 3 position 87E (line 11) [17]. The lig4\textsuperscript{169a} mutation deletes the majority of the LIG4 locus, including the promoter [31]. The mus309\textsuperscript{N1} mutation removes the region encoding the amino terminus of DmBlm, including part of the conserved helicase domain [12]. Females from the lacZ wild-type and mutant stocks were subjected to at least four back-crosses with \textsuperscript{w}\textsuperscript{118} males to create isogenic stocks. Heterozygous lacZ\textsuperscript{11} mus309\textsuperscript{N1}/TM3, lig4\textsuperscript{169a}/FM7w; lacZ\textsuperscript{11}/TM3, and lig4\textsuperscript{169a}/FM7w; lacZ\textsuperscript{11} mus309\textsuperscript{N1}/TM3 stocks were created by standard genetic crosses and mutations were verified by PCR.

Genomic DNA isolation
For mutation frequency studies, flies were snap frozen at various ages and stored at -80°C. Each sample consisted of 50 pooled male or female flies. For the tissue-specific experiments, each sample consisted of 100 heads, thoraces, or abdomens that were isolated by cutting snap-frozen flies with a razor blade. Flies were homogenized in 600 μl of lysis buffer (10 mM Tris-HCl, pH 8.0; 10 mM EDTA; 150 mM NaCl) in 2 ml eppendorf tubes using a battery-operated pestle. To the homogenate, 12 μl of Proteinase K (25 mg/ml), 60 μl 10% SDS and 10 μl RNase A (20 μg/ml) were added and samples were incubated at 65°C while rotating during 30 minutes. Genomic DNA was subsequently extracted from these samples using phenol/chloroform as described [16]. DNA yield and quality was subsequently extracted from these samples using phenol/chloroform as described [16]. DNA yield and quality was estimated by electrophoresis of a small amount of the purified DNA on a 1.0% agarose gel and by spectrophotometric measurement.

Mutation analysis
The mutation frequency was determined as described in detail elsewhere [16]. Briefly, isolated DNA from 50 flies or 100 body parts was digested for 1 hour at 37°C with HindIII (40 U) in the presence of magnetic beads coated with lacI-lacZ fusion protein. The lacZ plasmid was then eluted from the beads by incubation with isopropyl β-D-1-thiogalactopyranoside (IPTG), circularized by ligation with T4 DNA ligase (New England Biolabs, Ipswich, MA, USA), precipitated with ethanol and used to electrot transfect Escherichia coli (ΔlacZ, ΔgalE\textsuperscript{-}). Each mutant frequency determination point was based on at least three replicates from the same sample, that is, three 50-fly groups or 100 body parts from the same population, with a minimum of 100, 000 colonies for each rescue. LacZ plasmids from mutant colonies were further characterized as described in detail elsewhere [17]. Sequencing reactions of purified mutant plasmids were conducted at the Albert Einstein College of Medicine Genomics core facility. The returned chromatograms were analyzed with Sequencher (Gene Codes, Ann Arbor, MI, USA). Analysis of large rearrangements consisting of non-lacZ sequences was carried out using the fly genome database [32]. After alignment with the D. melanogaster sequence, the chromosomal origins of the flanking sequences were determined and the orientation and the type of chromosomal rearrangements deduced as described [33].

Histology
Cohorts of 20 flies were immersed in Telly’s fixative (20 parts 70% ethanol, 2 parts 37% formalin, 1 part glacial acetic acid) for a minimum of 48 hours at 4°C prior to processing with a Leica ASP 300 automatic tissue processor using standard techniques. The processed flies were embedded in paraffin blocks, sectioned into ribbons 6 to 7 microns thick, placed onto glass slides, and baked at 65°C for 12 hours to increase tissue adherence prior to staining with hematoxylin and eosin. Digital microphotography was done with an Olympus VANOX-T photomicroscope and a Q Imaging digital camera. All samples were coded and scored without prior knowledge of genotype.

Lifespan analysis
Homzygous flies collected from isogenized stocks were aged in groups of < 20 flies in vials kept at 25°C and constant humidity on a 12-hour light:12-hour dark cycle. Flies were transferred without anesthesia to new food every 2 to 3 days. For each lifespan experiment, at least 150 flies were monitored and deaths were scored every 1 to 2 days. Lifespan experiments were repeated three times for wild-type and blm flies and two times for lig4 flies and lig4 blm flies.

Statistical analysis
Somatic mutation frequencies were compared with one-way ANOVA tests followed by Tukey-Kramer multiple comparisons tests for three or more different genotypes and Mann-Whitney tests for comparisons between two genotypes. Gaussian distributions were confirmed using the Kolmogorov and Smirnov method. For tumor frequencies, two-tailed Fisher’s exact tests were used to compare between genotypes. Survivorship curves were compared using Wilcoxon matched pairs signed-rank tests. All statistical analysis was conducted using GraphPad Prism software.

Additional material

**Additional file 1**: Methods used to calculate spontaneous mutation frequency and analyze mutations
**Additional file 2**: Information about the numbers of colonies obtained in the mutation frequency experiments
**Additional file 3**: Information about the genome rearrangements isolated in this study
**Additional file 4**: Set of tables showing data related to tumor frequency in wild-type and mutant backgrounds and statistical analysis for tumor frequency in different genetic backgrounds
Additional file 5: Lifespan analysis for lig4 blm double mutant flies.

Abbreviations
C-NHEJ: classical non homologous end joining; DmBlm: Drosophila melanogaster Blm; DS8: double-strand break; HR: homologous recombination; Lig4: DNA ligase 4.

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Competing interests
The authors declare that they have no competing interests.

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