Migrating bubble synthesis promotes mutagenesis through lesions in its template

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ABSTRACT

Break-induced replication (BIR) proceeds via a migrating D-loop for hundreds of kilobases and is highly mutagenic. Previous studies identified long single-stranded (ss) nascent DNA that accumulates during leading strand synthesis to be a target for DNA damage and a primary source of BIR-induced mutagenesis. Here, we describe a new important source of mutagenic ssDNA formed during BIR: the ssDNA template for leading strand BIR synthesis formed during D-loop migration. Specifically, we demonstrate that this D-loop bottom template strand (D-BTS) is susceptible to APOBEC3A (A3A)-induced DNA lesions leading to mutations associated with BIR. Also, we demonstrate that BIR-associated ssDNA promotes an additional type of genetic instability: replication slippage between microhomologies stimulated by inverted DNA repeats. Based on our results we propose that these events are stimulated by both known sources of ssDNA formed during BIR, nascent DNA formed by leading strand synthesis, and the D-BTS that we describe here. Together we report a new source of mutagenesis during BIR that may also be shared by other homologous recombination pathways driven by D-loop repair synthesis.

INTRODUCTION

For much of the cell cycle, DNA remains double-stranded. However, during some processes of DNA metabolism, single-stranded DNA (ssDNA) becomes transiently exposed. Examples of such processes include S-phase replication, homologous recombination, transcription and DNA resection at uncapped telomeres or double strand breaks (DSBs) (reviewed in (1)). While lesions in the context of double-stranded DNA (dsDNA) can often be excised and repaired using the opposite strand as a template (reviewed in (2)), exposed ssDNA is vulnerable to a variety of damages, and is more limited in its capacity for error-free repair of DNA lesions. Moreover, with the exception of direct reversal, repair of lesions in ssDNA often leads to mutations (1).

Hypermutability of ssDNA has been observed in the presence of damaging agents that preferentially target ssDNA. Such ssDNA damages include base-alterations inflicted by alkylating agents (e.g. methyl methanesulfonate (MMS)) (3–5), oxidative damage (6,7), and cytidine deamination induced by cytidine deaminases such as AID/APOBEC enzymes (5,8–15). Damages inflicted in ssDNA often lead to the formation of mutation clusters (groups of closely spaced mutations) that are formed in regions of exposed ssDNA (1,4,5,11,14,16–21). These mutation clusters are also often strand-coordinated (comprised of the same mutant base present in the same DNA strand) (1,4,5,8,17,22). Thus, mutation clusters observed in yeast model systems and in human cancers have served as markers of persistent ssDNA (1,4,5,8–10,12,14,16,18,22–25). In addition, the pattern of strand coordination in mutation clusters can inform on the mechanism of ssDNA formation. For example, ssDNA formed by bidirectional resection of a DSB usually leads to the formation of clusters with switching strand coordination (1,5,17,25), while ssDNA formed in the nascent strand of BIR often leads to non-switching coordinated mutation clusters (4,17,21).

Another important property of persistent ssDNA is its ability to stimulate formation of secondary (non-B) DNA structures. S-phase DNA replication often stalls at secondary DNA structures formed in the template, and this leads to genetic instabilities, including mutations and chromosomal rearrangements (26–30). For example, in humans, G-quadruplexes (G4) that form in ssDNA during transcription were shown to promote recombination in patients with Bloom syndrome, presumably due to stalling...
of replication at G4 structures (31). Extensive analysis of cancer genomes revealed significant association between sequences with the potential to form non-B DNA structures (Hairpin/cruciform, G4, triplex, etc.) and rearrangement breakpoints (26,28,32,33). Additionally, studies in yeast demonstrated that inverted DNA repeats (IR) that can adopt a hairpin (stem-and-loop) structure in ssDNA, stimulate deletions and genomic rearrangements (34–42). Hairpin structures may form in ssDNA exposed during lagging strand synthesis, promoting stalling of the replicative DNA polymerase at the base of the hairpin, and often leading to replication slippage that produces deletions (26,34,36,37). Replication slippage (based on results obtained in various organisms (26,36,37,43,44)) often proceeds between short repeats (microhomologies) that are brought into close proximity by formation of the hairpin by IRs. It was proposed that the frequency of such IR-induced polymerase slippage at positions of microhomology is directly proportional to the frequency of hairpin formation, which is tied to the length and persistence of ssDNA (26,34–37,45).

Aside from S-phase replication, some types of repair DNA synthesis promote accumulation of ssDNA (1,46–50). For example, repair of double-strand DNA breaks (DSBs) through a pathway called break-induced replication (BIR) is particularly susceptible to accumulation of ssDNA that is both longer and more persistent than the relatively short-lived ssDNA exposed during S-phase replication (4,21,51). BIR is initiated when only one DSB end can find homology in the genome for strand invasion, resulting in the formation of a D-loop structure that is typical of all homologous recombination (HR) (reviewed in (52–55)). Studies of BIR using the yeast Saccharomyces cerevisiae demonstrated that BIR, like other types of HR, is preceded by extensive 5′-to-3′ end resection, exposing a long single-stranded 3′ end that is then covered by RPA, and later by Rad51 protein to deoxyuridine (dU) (10,24). The dU lesions produced by HO endonuclease at the MATα locus, where a DSB can be introduced following HO induction by addition of galactose, are isogenic derivatives of AM1003, which contains two copies of Chromosome III (Chr III); one copy (recipient) is truncated and contains a recognition site for HO endonuclease at the MATα locus, where a DSB can be introduced following HO induction by addition of galactose. Another copy of chromosome III (the donor) is full-length and cannot be cut by HO due to MATα-inc mutation. For a complete list of all strains constructed for this study, and those used in this work that were published previously, see Supplementary Data S1. Construction of AM1003 is fully described in (64) and AM1003 has the following genotype: hmlΔ::ADE1/hmrΔ::ADE3 MATα-LEU2 tel/MATα-inc hmrΔ::HPH FS2Δ::NAT/FS2 leu2/leu2-3,112 thr4 ura3-52 ade3::GAL::HO ade1 met13

MATERIALS AND METHODS

Yeast strain construction and growth conditions

The yeast strains used for all experiments in this study are isogenic derivatives of AM1003, which contains two copies of Chromosome III (Chr III); one copy (recipient) is truncated and contains a recognition site for HO endonuclease at the MATα locus, where a DSB can be introduced following HO induction by addition of galactose. Another copy of chromosome III (the donor) is full-length and cannot be cut by HO due to MATα-inc mutation. For a complete list of all strains constructed for this study, and those used in this work that were published previously, see Supplementary Data S1. Construction of AM1003 is fully described in (64) and AM1003 has the following genotype: hmlΔ::ADE1/hmrΔ::ADE3 MATα-LEU2 tel/MATα-inc hmrΔ::HPH FS2Δ::NAT/FS2 leu2/leu2-3,112 thr4 ura3-52 ade3::GAL::HO ade1 met13
To construct the strains containing the lys2-InsH reporter at three different positions in the donor chromosome, we used three derivatives of AM1003 that contained THR4 inserted at the thr4 position of the MATα inc chromosome, and an insertion of LYS2 at the MATa (replacing MATα-inc region starting from the position located 249bp centromere-proximal from the border of the X–Y regions of the MAT locus and finishing at the position 5bp centromere-proximal to the Ye-Z1 border), at 16 kb or at 36 kb positions (see (65) for details).

The strains with insertions of LYS2 at MAT and 16 kb positions have been described previously (65), while the strain with LYS2 at 36 kb has been constructed here by transformation of THR4 AM1003 derivative ((64), see Supplementary Data S1) with a DNA fragment obtained by amplifying the LYS2 gene from the pL12 plasmid (66) using the following primers (5′ to 3′) where lower-case letters indicate homology to the wild-type LYS2 gene sequence and capital letters indicate homology to the respective position on Chr III:

Forward primer (FP):
ATCGTAAATACATAAGGTCGTCGGGATATGTACCTAG
AATAGTGTGCAGCAACTGTGACAGATACATTAC
AATTGCTCATTAAaatataaataatatcggag
CTGTTGGCATCATATC
ATTTGATAAAGAATGTTTTGCTTATCAT
reverse primer (RP):
AATCGGAAATCTTTCCTTGCTTGGCCCTATCAT
TTTTCTTTCCGGATTATGCTTATTATCT
AACTGGAAATGCTTTCCCTTTTGCCCTATCATTA
ATTGATAAAGAATGTTTTGCTTATCATAT

To insert the lys2-InsH reporter at three positions in the donor chromosome, a pCORE construct (containing KanMX and URA3 cassettes (67,68)) was inserted into the LYS2 gene, and then replaced by lys2-InsH sequence described in (36). Construction of strains containing the lys2-Δ4 reporter cassette is described in (65). Reporter strains were confirmed by PCR and phenotype at each step of construction. “No DSB” strains were created by plating on YEP-Gal media and selection of colonies with an alpha- marker amplification contained tail sequences homologous to the first and last 80 bp of the open reading frame of each gene to be disrupted (69). Successful disruptions were confirmed by both PCR and by the phenotypes of transformants (where possible). Mutants harboring the pol3-ΔY708A and pol3-Δt mutations were constructed by restriction digest of plasmids p170 (71) and p171 (72) respectively with HpaI prior to transformation. The pol3-Δt mutant was created by co-transformation of the CRISPR-Cas9 expression vector bRA89 (73) modified to express sgRNAs targeting the sequences (5′-3′): TCTTTGATATCGAGTGTC and the following repair template (5′-3′): CAGCCTCATTGCTATCATGTCCTTTGGCTATGC
GTTGCTGGTAAAGGATCTGCGCTTCCGGAACTGGAATCGATCCC.

Synthetic dropout (Sc), rich growth media (YPED – yeast extract, peptone, and dextrose), YEP-lactate (YPE-Lac) and YEP-galactose (YPE-Gal) media were prepared similar to (74). Expression of APOBEC3A was achieved by transformation with an A3A-expression vector (pSR355) or EV control (pSR419) plasmids containing an HPH marker (22). Selection for correct transformants was done similar to (21). Yeast cultures were grown at 30°C for most experiments. Yeast strains harboring the temperature sensitive pol3-t mutation were grown at 18°C prior to DSB induction.

Determining the rate of BIR-associated mutagenesis

Yeast strains from single colonies were grown with agitation in Sc media lacking leucine for approximately 20 h, diluted 20× with YEP-Lac and grown to logarithmic phase (for ~16 h). DSBs were induced by addition of galactose to a final concentration of 2% (w/v). Due to residual BIR, which is capable of affecting Lys+ frequency even prior to galactose addition (74), the level of Lys+ during S-phase replication was determined in no-DSB control strains where HO recognition sites have been removed (64,65).

“No DSB” control strains were grown under the same conditions. In experiments including A3A and empty vector expression, hygromycin (1% w/v) was added to the YEP-Lac medium. After DSB induction, cultures were incubated at 30°C (or 20°C for low-temperature experiments as specified in Supplementary Figure S3) for 7h with agitation.

Appropriate amounts of culture were plated at 0h (before galactose addition) and 7h (after galactose addition) time points on YEPD and Sc-lysine dropout media (Sc-Lys) (and on Sc-adenine dropout (Sc-Ade) and Sc-adenine/lysine dropout (Sc-Ade/Lys) media for POL3 mutant experiments) for strains harboring the lys2-InsH reporter cassette, or Sc-Ade dropout and Sc-adenine/uracil dropout media (Sc-Ade/Ura) for strains harboring the ura3-29 reporter cassette. In experiments using lys2-InsH reporter strains, DSBs were initiated, DSB repair outcomes were identified, BIR efficiencies were calculated, and frequencies and rates of Lys+ reversions were determined as described in (65,75). For ura3-29 reporter strains containing A3A or empty vectors, experiments were performed, BIR repair outcomes were identified, BIR efficiencies were calculated, and Ura+ reversion rates were determined similar to (51). Rates and frequencies are reported as median values and 95% confidence intervals (or ranges for experiments with fewer than 6 biological replicates). Mann–Whitney U test was used to draw statistical comparisons of mutation rates between different strains. Fisher’s exact test was used to make statistical comparisons between fractions of different categories of repair outcomes.

To analyze mutation spectra, Lys+ or Ura+ colonies were selected randomly from 7h plates from experiments where cells underwent BIR (all colonies were considered to be independent BIR events due to no additional cell...
divisions before plating and big increase of 7 h over 0 h mutation frequency). For “No DSB” controls, only one colony (Lys+) was selected from each independent culture after plating (similar to (65)). Lys+ mutation spectra were determined by PCR amplification of LYS2 alleles using the following primers in Lys+ outcomes (5’ to 3’): CCATCCACCTTCATCTGAAAGACC, and AAATGTCACTGCAAATTAGTGGCGGAAGAC. The PCR products were then Sanger sequenced using the following primer (5’ to 3’): GTTCGTACCCCTCTCGAGAATA. Lys+ outcomes were confirmed to be heterozygous after completion of BIR using the following primer pair, where the forward primer anneals to the spacer region between the repeats of the InsH quasi-palindrome and is indicative of the presence of an unaltered lys2-InsH allele (Lys−), while the reverse primer anneals to the LYS2 sequence outside of the quasi-palindrome (5’ to 3’): ATCCTGGAAAACGGGAAAGG and AAATGTCACTGCAAATTAGTGGCGGAAGAC respectively. Outcomes that did not produce a product using these two primers were considered to be homozygous and excluded from spectra results. Likewise, Ura+ mutation spectra were determined by PCR amplification of Ura+ outcomes using the following primers (5’ to 3’): CTGGAGTTAGTTGAAGCATTAGG. Homozygous and excluded from spectra results. Likewise, Ura+ variants with allelic frequencies lower than 0.85 were called heterozygous. Identical mutations that occurred at the same chromosomesal positions in different biological replicates were classified as existing prior to BIR induction and were thus discarded. C to N variants along the BIR track were appropriated as markers of A3A-induced lesions in the ssDNA template for lagging strand synthesis. Likewise, G to N variants along the BIR track were appropriated as markers of A3A-induced cytidine deamination in ssDNA of the template for the leading strand synthesis (D-loop bottom template strand (D-BTS)). Only C to N and G to N mutations on the right arm of Chr III were counted as representing BIR-related A3A-induced mutations.

Simulations of randomly distributed G to N mutations were performed by identifying the cumulative number of G to N positions across all sequenced samples on all chromosomes (total of 98 mutations) and redistributing them randomly across a synthetic genome to generate 100 000 unique samplings. From each of these samplings, the number of G to N mutations that occurred on the right arm of Chr III was recorded and used to create a frequency histogram and kernel density estimation. All custom code used for the variant filtering, simulations and graphing is available through GitHub (https://github.com/malkovalab/WGS-A3A-Tools).

Deep sequencing

Strains containing the Ori1 or Ori2 lys2-InsH reporter at the 16 kb position (Supplementary Data S1) were used to perform BIR-induction experiments as described in (74). For DNA purification, 5 ml of cell cultures containing ∼3 × 10⁶ cells/ml were collected before (0 h) galactose addition and 12 h after galactose addition, and genomic DNA was extracted using the glass bead protocol as described in (21). The region corresponding to the insertion of insH in LYS2 (∼700 bp-long) was amplified by PCR from these samples by using the following primers:

Forward primer 1 (5’ to 3’): AAATGTCACTGCAAATTAGTGGCGGAAGAC
Reverse primer 1 (5’ to 3’): TGATAGTTGAATAAACCTTTGGATCCCTCGAC

The PCR products were separated by gel electrophoresis (using 1% agarose). Following electrophoresis, PCR fragments of ∼400–700 bp were excised from the gel and used for gel extraction using QIAquick Gel Extraction Kit (QIAGEN #28704). The gel extracted products were subsequently used for a second PCR amplification of a ∼500 bp region using the following primers:

Forward primer 2 (5’ to 3’): GTTCGTACCCCTCTCAGAAT
Reverse primer 2 (5’ to 3’): CCATCCACCTTCATCTGAAAGACC

The PCR products were purified by using QIAquick PCR Purification Kit (QIAGEN, #28104). The resulting DNA samples (∼20 ng/µl) were submitted to GENEWIZ
for deep sequencing through the Amplicon-EZ sequencing pipeline (without Illumina® partial adapters). To analyze the deep sequencing data, reads were first trimmed (Trimmomatic-0.39). Next, reads with a Q30-dust scores lower than 0.07 were removed (PRINSEQ+++, version 1.2). Only reads containing an unchanged sequence of either primer used for the final amplicon (Forward and Reverse primers 2) were next selected and trimmed to their 5' ends. All reads shorter than 220 bp and reads supporting no-deletion events were discarded. Remaining reads were sorted and grouped by common sequences. Junction positions of most common deletion events were manually verified and used for alignment (tolerance of two errors) of remaining reads. The custom code used for the analysis is available through GitHub (https://github.com/malkovalab/DeepSeqTools).

## Determining the frequency of IR-mediated deletions by droplet digital PCR (ddPCR)

Yeast cells containing the Ori1 or Ori2 lys2-InhH reporters at the 16 kb position were used to perform BIR-induction experiments as described in (74,76). 1.5 ml of yeast cultures with 5 x 10⁷ cells/ml were collected by centrifugation 12 h post-BIR induction.

The cells were resuspended in 1 ml Spheroplasting buffer (0.4 M sorbitol, 0.4 M KCl, 40 mM sodium phosphate buffer pH 7.2, 0.5 mM MgCl₂), then digested by addition of 5 µl Zymolase buffer (0.1 g/ml 20 T zymolase (MP Biomedicals, #08320921) dissolved in 2% glucose, 50 mM Tris–HCl, pH 7.5), and incubated at 37°C for 15 min. After that, cells were collected by centrifugation at 3000 rpm for 2 min with all liquid removed. Cells were then resuspended by addition of 500 µl 1× Cut smart buffer (50 mM potassium acetate, 20 mM Tris-acetate, 10 mM magnesium acetate, 100 µg/ml BSA, pH 7.9 at 25°C). Next, 5 µl 10% SDS buffer and glass beads (about 300 µl volume) were added to the resuspended cells and then suspensions were vortexed for 1 min to break all cells. To remove RNA, 20 µl 10 mg/ml RNase buffer was added to the mixture, which was then incubated at 37°C for 30 min. After incubation, 500 µl phenol:chloroform:isoamyl alcohol (25:24:1) (Fisher, #15593-049) were added to the mixture followed by brief vortexing and centrifugation at 13 000rpm for 15 min. The upper layer was transferred to a new tube by careful pipetting. Another round of wash by phenol:chloroform:isoamyl alcohol (25:24:1) followed by centrifugation was repeated. The upper layer after centrifugation was transferred to a new tube and mixed with 50 µl 3M sodium acetate buffer (PH 5.2) and 500 µl isopropanol. After brief vortexing, the mixture was kept at room temperature for 20 min and then centrifugated at 13 000 rpm for 20 min. All liquids were removed without touching the pellet of DNA. Then 500 µl 80% ethanol were added to the tube followed by centrifugation at 13 000 rpm for 5 min. All liquid was removed, and pellets were left to air dry for at least 10 min. 50 µl water was added to dissolve the dried DNA, which was then quantified by Qubit and stored at −20°C if not immediately used.

To detect Type I events, 2 µl of undiluted DNA was mixed with 10 µl 2x ddPCR supermix for probe (no dUTP Bio-rad 1863023), 7 µl water, and 1 µl of 20× primer sets that specifically recognize the junction formed in Type I events (see sequences below). To determine the amount of yeast genomic DNA, the original DNA solutions were diluted by serial dilution to ~0.2 ng/µl, and then 2 µl of the diluted DNA were mixed with 10µl 2× ddPCR supermix for probe (no dUTP), 7µl water, and 1 µl of 20× ACT1 primer set (sequences are listed below). The PCR mixture and Droplet generation oil (Biorad, #1863005) were used to generate droplets using QX200 Droplet Digital PCR (ddPCR) System. PCR reactions for both Type I events and for the ACT1 locus were conducted with the following program: step 1: 95°C for 10 min; step 2: 94°C for 30 s (2°C change per seconds), 60°C for 2min 30 s (2°C change per second); step 3, repeat step 2 for 39 cycles; step 4: 98°C for 10 min; step 5: 12°C for 30 min. After PCR, the reactions were analyzed by QX200 Droplet Digital PCR (ddPCR) System. The Type I events were analyzed by channel 1 to detect the FAM signal, while the ACT1 locus was analyzed by channel 2 to detect the HEX signal. The sequences for the primers sets (ordered from IDT) and used for the ddPCR analysis are as follows: Type1 (ratio of primers and probe was 4:1 and the probe was labeled by FAM): Forward (5’ to 3’): CGTGGTGTTCGCAATCTCGCA Reverse (5’ to 3’): TCTTTACATCAGTGCGTACAG Probe (5’ to 3’): AGGCAGGTATCTATTATAGGTCGTAAGTTGGA. ACT1 (ratio of primers and probe was 4:1 and the probe was labeled by HEX): Forward (5’ to 3’): GCCTTCCTAGTTTCCTATCCA Reverse (5’ to 3’): GGGTAGAGAAGACCAGGGTAA Probe (5’ to 3’): TCGGGGGATGGTGTACTCTAGTGCT

## RESULTS

BIR promotes deletions between microhomologies flanking inverted DNA repeats

When IRs are included into single-stranded DNA (ssDNA) they form secondary DNA structures (hairpins) promoting replication stalling and polymerase slippage at microhomologies leading to deletions between them. It is believed that the frequency of replication slippage-induced deletions is determined by the frequency of hairpin formation, and therefore by the persistence of ssDNA (26,34–36). Because BIR is a source of long, persistent ssDNA (4,21,51), we asked whether this ssDNA would promote hairpin-induced deletions at microhomologies flanking IRs placed on the synthesis track of BIR. To accomplish this, we used our BIR experimental system in Saccharomyces cerevisiae dizomic for chromosome III (64). In this system, a galactose-inducible HO-endonuclease initiates a DSB at the MATα locus on a Chromosome III (Chr III) that is truncated by the insertion of LEU2 and telomere sequence centromere distal to MAT. BIR is the dominant repair pathway in this system and is initiated by 5’ to 3’ resection that can proceed for long distances (up to the centromere) followed by invasion of the broken chromosome into the homologous region of the full copy of Chr III (the donor) that contains a MATα-inc allele that cannot be cut by HO-endonuclease (Figure 1A). Because the length of resection is variable, the exact site of
Figure 1. BIR promotes deletions between microhomologies flanking inverted DNA repeats. (A) The lys2-InsH reporter before (0 h) and after (7 h) DSB. Top. BIR is initiated by induction of a DSB by HO endonuclease at MATα located in a truncated (recipient) chromosome (R) of yeast disomic for Chr. III. Invasion of one broken end of the recipient into the homologous full-length donor (D) chromosome containing MATα-inc establishes BIR synthesis that can progress approximately 100kb to the end of the chromosome. The lys2-InsH (Lys−) reporter consists of 69bp inverted repeats (yellow) separated by a 9bp spacer (purple) and is flanked by 9bp direct repeats (red). The construct was inserted at three different positions along the track of BIR synthesis in the donor chromosome: “MAT” (MATα-inc), “16kb” centromere distal from MATα-inc, and “36 kb” centromere distal from MATα-inc. Bottom. Replication slippage during BIR proceeding through the lys2-InsH reporter can result in deletion of InsH producing a functional allele (LYS2*) conferring a Lys+ phenotype. Positions of primers (FP: forward primer, RP: reverse primer) for confirming deletions by PCR are shown for both LYS2* and lys2-InsH alleles. (B) Lys+ reversion rates for the strains containing lys2-InsH reporter cassette at the three positions specified in (A) prior to DSB induction (DSB 0h), following BIR (DSB 7h), as well as in isogenic strains lacking the HO cut-site (No DSB). Asterisks indicate statistically significant difference (P < 0.005) from Lys+ reversion rates at 0 h (DSB 0h). Numbers above bars indicate median rates (see Supplementary Data S2 for precise P-values and 95% CIs). (C) Representative gel image showing PCR-amplified products of the lys2-InsH allele and the LYS2* allele (404 bp) present in a single BIR outcome (lane 1). PCR products of WT LYS2 (374 bp) (lane 3) and the lys2-InsH construct before BIR (530 bp) (lane 4) are also shown for comparison, along with a 100-bp DNA ladder (L) in lane 2. Primers used for all PCR products are those shown in (A, bottom) (FP and RP). Analysis of lys2-InsH deletion products (like LYS2*) was performed by Sanger sequencing.
The polarity of insH deletions induced by BIR

Previous studies reported the precise excision (deletion) of insH that occurred during S-phase DNA replication and was mediated by replication slippage at the flanking 9 bp direct repeats of the LYS2 sequence (34,36). Because we observed that PCR products from 7 h Lys+ revertants often did not match the wild-type LYS2 PCR product by size, we asked whether deletions of insH formed during BIR synthesis are imprecise. We Sanger sequenced 7 hr Lys+ BIR outcomes from the 16 kb lys2-InsH reporter strain and Lys+ isolates from isogenic “no DSB” strains for comparison. We observed that the majority of insH deletions during BIR were imprecise and asymmetrical such that the deletion products retained a part of the insH quasi-palindrome; the remaining sequence was either from the left inverted repeat (Type I deletion) or from the right inverted repeat (Type II deletion) (Figure 2A–C). The Type II deletion also resulted in a loss of a portion of the LYS2 gene sequence, yet still produced a Lys+ phenotype, despite slower growth of colonies on synthetic media lacking Lysine (Figure 2A, B, C). All three of the deletion types observed (Precise, Type I and Type II) produced an in-frame LYS2 gene sequence (Figure 2A, B) and contained microhomologies on their breakpoints (9-bp-long for precise deletions and 6-bp-long for imprecise deletions) (Figure 2C). Of the Lys+ isolates from the “No DSB” strains (reflecting deletions resulting from S-phase DNA replication), there were significantly more precise deletions of insH as compared to BIR (Figure 2B), though the majority were still imprecise Type I deletions. Importantly, all three deletion types (precise, Type I and Type II) were stimulated by BIR, while Type I occurred most frequently. When we inserted lys2-InsH at the same position (16 kb) in inverted orientation (Ori2) with respect to that of the original strain (Ori1) (see schematic in Figure 2B), the frequency of BIR-associated Lys+ reversions was increased 3.3 times as compared to what we observed in Ori1 (Figure 2D), and all of the Lys+ outcomes (20/20) analyzed by Sanger sequencing) were Type I (Figure 2B). By direct comparison, Type I frequencies were 5.1 × more frequent during BIR with the Ori2 reporter than with the Ori1 reporter, even though in both orientations Type I deletions were greatly stimulated by BIR (Figure 2D). We believe that the lack of Type II events among Ori2 outcomes results from a significant decrease in the fraction of Type II events (from 33% among Ori1 outcomes to ~2% among Ori2 outcomes). This calculation is based on the 5.1 × increase in Type I events (from leading to lagging strand (Figure 2D)) and therefore on 5.1 × decrease of Type II events (from lagging to leading strand). We also performed Sanger sequencing of Lys+ BIR outcomes from Ori1 MAT and 36 kb reporter positions and found that they produced only Type I imprecise deletions (Supplementary Figure 1A). To assess whether the ability for the reporter to produce a Lys+ phenotype may differ between reporter positions (and could result, for example from differences in the levels of LYS2 expression between different reporter locations), we transformed strains with reporters in different positions with both Type I and Type II imprecise deletion fragments obtained from the 16 kb Ori1 strain. Indeed, we found that only Type I deletion fragments supported a Lys+ phenotype at MAT and 36 kb positions (even though with varying efficiency), while both Type I and Type II fragments did so at 16 kb in the Ori1 strain (Supplementary Figure 1B). Due to the possibility that the selection for Lys+ reversions could be affected by the level of LYS2 expression for two orientations of the reporter as well, and also because Lys+ selection precludes identification of any deletions that inherently do not produce Lys+ outcomes (e.g. out of frame deletions), we next analyzed non-selected post-BIR cells by deep sequencing. Specifically, yeast cultures were collected 12 h following DSB induction, at the point when BIR was expected to complete in the majority of the cells. The DNA purified from these cells was subjected to
Figure 2. (A) Sequences of full lys2-InsH reporter and its derivatives following precise (restoring the wild type LYS2 allele) and imprecise (Type I and Type II) deletions of InsH. Colors correspond to those depicted in Figure 1A from the full-length lys2-InsH allele in the donor chromosome. Gray highlighted and underlined bases are indicative of breakpoint microhomologies. (B) Schematic of lys2-InsH deletion polarities for observed deletion types. Gray arrows indicate the implied direction of synthesis. The colors of microhomologies are similar to (E).
PCR amplification and deep sequencing of the lys2-InsH region followed by identification of reads containing deletions of InsH. This method allowed us to reveal the main types of deletions missed in reporter experiments using Lys⁺ selection and to identify the most frequent deletion types among them.

By deep sequencing, we detected Type I (which was the most frequent following BIR in both Ori1 and Ori2 reporters), Type II, and precise deletion types, and we detected several new types not seen in reporter experiments (Figure 2E, Supplementary Data S3, S4). One of the more frequent new types that we detected, which we called J1 (type 7 in Figure 2E; Supplementary Data S3, S4), resulted from deletion of 125 nucleotides and contained 5bp microhomologies on its boundaries (Figure 2E; Supplementary Data S3, S4). Because this deletion was not in-frame it could not produce a Lys⁺ outcome but appeared to be a major type due to its relative frequency among the other deletion classes.

One limitation of the deep sequencing method is that it cannot yield accurate absolute frequencies of BIR-induced deletion events due to the requirement for standard PCR amplification, which can introduce bias among deletion outcomes and is biased against outcomes without deletions that contain the hairpin-forming IR and thus do not amplify faithfully to calculate their frequency in the cell population. To address this limitation, we next used a highly sensitive digital droplet PCR (ddPCR) method to detect the absolute frequencies of the most abundant deletion types during BIR. Although the frequencies of Type II and J1 deletions were below the threshold for ddPCR detection, the Type I deletion was detectable, and we determined that the frequencies of Type I during BIR were \( \sim 4 \times 10^{-5} \) in Ori1 and \( \sim 1 \times 10^{-3} \) in Ori2 (Figure 2F, Supplementary Figure 2). Importantly, the frequencies of Type I for No-DSB controls for both Ori1 and Ori2 were below ddPCR detection threshold (Supplementary Figure 2), thus confirming that the Type I frequencies calculated by ddPCR following BIR were indeed BIR-specific.

Together, we conclude that BIR promotes deletions at microhomologies flanking IRs. The majority of these deletions have a polarity (with one microhomology located inside the inverted repeat and another outside of the repeat (Figure 2E, G)). This type of polarity was previously observed for similar events mediated by Polδ during S-phase lagging strand synthesis (36). There it was proposed that following hairpin formation, Polδ can copy via displacement synthesis inside a hairpin but undergoes frequent template switches from inside to the outside the hairpin (36). It was also proposed that the polarity of the deletion events reflects the direction of synthesis (from the microhomology inside the hairpin towards one outside). Using the same logic for BIR, we propose that Type I events result from slippage during lagging strand synthesis in Ori2 and during leading strand synthesis in Ori1 (Figure 2G, see also Discussion and the Figure 6B for details). Even though the frequency of the latter is lower than the frequency of the former, our data suggest that ssDNA in the template for leading strand synthesis forms and persists, which was not previously appreciated.

The effect of polymerase mutations on InsH deletions

Polδ was recently confirmed as the main replicative polymerase driving both leading and lagging strand synthesis during BIR (77), which makes it likely that it also mediates deletions of InsH. Yet, stalling of Polδ during BIR was proposed to recruit translesion polymerase ζ (Polζ) to mediate template switching at microhomology (75). Because the deletions of InsH that we observed during BIR were mediated by microhomology, we sought to determine whether a similar switch from Polδ to Polζ might also take place in the process of InsH deletion. We observed that deletion of REV3, encoding the catalytic subunit of Polζ, had no significant effect on the rate of Lys⁺ reversions of the lys2-InsH reporter at the 16 kb position during BIR (Figure 3A, Supplementary Data S2). To assess possible redundancy with Polymerase η (Polη), another translesion polymerase that has the capacity to substitute for Polδ at lesions (reviewed in (78)), we also created rad50Δ (deletion of the gene encoding the catalytic subunit of Polη) and rev3Δ rad50Δ mutants. Neither of these mutations had any significant effect on the rate of Lys⁺ reversion of the lys2-InsH reporter after BIR as compared to the wild-type (Figure 3A, Supplementary Data S2). In addition, we did not observe any effect following the deletion of POL4 (Supplementary Data S2).

Because we did not find any evidence supporting participation of translesion polymerases in the deletions of InsH, we next tested the effects of various mutations affecting Polδ, the main polymerase driving both leading and lagging strand BIR synthesis (77). Three mutations in POL3, pol3-t, polβ-Y708A and pol3-01 were selected for this analysis. In particular, pol3-t (Polδ active site mutation (36,37,79)), was previously shown to greatly increase deletions at microhomologies promoted by inverted repeats in S-phase (36,37). A second mutation, pol3-Y708A (Polδ nucleotide binding pocket mutation (71)), was characterized as having a mutator phenotype dependent on Polζ (71,80,81). This mutation was shown to decrease processivity during BIR, leading to increased half-crossover outcomes with reduced BIR efficiency (82), but the distance that Polδ carrying the pol3-Y708A mutation could synthesize during BIR remained unknown. A third mutation, pol3-01, leads to proofreading deficiency of Polδ (83) and was shown to eliminate frame shifts and complex events involving template switches during gene conversion (84). Based on these data and on preceding investigations (85), the pol3-01 mutant was hypothesized to create a more processive Polδ during BIR. Therefore, we hypothesized that if this is true, then a more processive Polδ might displace secondary structures more easily, and therefore affect the frequency of deletions and their spectrum in our system.

For pol3-Y708A (and for pol3-Y708A rev3Δ) mutants, we observed that BIR synthesis never reaches the InsH position (16kb) in our system. This was concluded based on a very high frequency of CL and HC (defective DSB repair outcomes) (Figure 3C), which was consistent with previous observations (82), as well as on a very low level of Lys⁺ reversions which did not increase following DSB induction (Figure 3B, Supplementary Data S5). Based on these observations, we concluded that even Ade⁺ Leu⁻ colonies that were observed in this mutant were not completed BIR events, but
Figure 3. The effect of polymerase mutations on InsH deletions. (A) Eliminating Pol η (rad30Δ) or Pol ζ (rev3Δ) does not affect the frequency of Lys⁺ resulting from deletions in lys2-InsH at 16kb during BIR. “NO GAL” = rates prior to the addition of galactose. “GAL” = Lys⁺ rates following BIR. Median values are listed above the bars. N.S. = not significantly different from wild-type (RAD30 REV3) (P ≥ 0.05 determined by Mann-Whitney U test, see Supplementary Data S5 for P-values and 95% CI). (B) The effects of mutations in POL3, including: pol3-01, pol3-Y708A, and pol3-t on Lys⁺ reversion rate after BIR induction in strains harboring the lys2-InsH reporter at the 16kb position. Sc-Lys: Lys⁺ were selected for lysine prototrophy only. Sc-Ade/Lys: Lys⁺ were selected for lysine and adenine prototrophy to eliminate chromosome loss and half-crossover outcomes which exhibit adenine auxotrophy. Asterisks indicate values that were significantly different (P < 0.01) from wild-type (POL3) and N.S. indicates no significant difference (P ≥ 0.05) from wild-type (POL3). “<1” indicates that rates were not calculable (see Supplementary Data S5 for frequencies used in rate calculation). Other details similar to (A). (C) BIR efficiency is reduced (and chromosome loss and half-crossovers increased) in pol3-Y708A and pol3-t mutants as compared to POL3. BIR efficiency is not affected in pol3-01. Asterisks indicate significant differences in DSB repair outcome fractions from POL3 (P < 0.05) measured by Fisher’s exact test. Different DSB repair outcomes are illustrated in the schematic (left). (D) lys2-InsH deletion spectra in POL3, pol3-t, and pol3-01 strains. P-values are listed to indicate statistically significant differences (P < 0.05) in the fractions of individual deletion types compared to POL3 measured by Fisher’s exact test. N.S. = no significant difference (P ≥ 0.05).
rather aberrant repair outcomes (similar to those described in (82)), where BIR synthesis was interrupted before reaching the 16kb position.

Next, Lys⁻ release rate in the pol3-01 mutant was not significantly different from the wild-type POL3 strain (Figure 3B), nor did this mutation have any effect on BIR efficiency (Figure 3C). Additionally, the pol3-01 mutant showed no difference in InsH deletion spectrum as compared to the spectrum of the wild-type POL3 strain (Figure 3D). Combined with Lys⁻ release frequency data for this mutant, our results do not show any evidence of increased displacement activity of Polδ in the pol3-01 mutant in our BIR system.

Finally, we observed that in pol3-t mutants, Lys⁺ release rate during BIR was modestly (1.9×), but significantly increased as compared to the wild-type POL3 strain, indicating a higher rate of InsH deletion (Figure 3B, left (Sc-Lys)). As expected, BIR efficiency was also reduced in pol3-t mutants and the frequency of chromosome loss (CL) and half-crossover (HC) outcomes was increased, (Figure 3C), consistent with previous observations (82). When experiments were carried out in the absence of adenine to eliminate CL and HC outcomes (see Figure 3C, schematic), pol3-t Lys⁺ release rate was more dramatically increased (3.4×) as compared to wild-type POL3 strains (Figure 3B, right (Sc-Ade/Lys)). In addition, we observed that pol3-t mutation altered the distribution of deletions, such that precise deletions were favored, which was different from the spectrum observed in wild-type POL3 where Type I imprecise deletions were predominant (Figure 3D). One possible explanation for this change was that progression of synthesis by pol3-t is kinetically slower than POL3 and this allows the formation of longer ssDNA regions allowing the full InsH hairpin to form more often. However, when BIR was induced in POL3 (wt) strains at 20°C (which could also slow down BIR progression), Lys⁺ release rate and the spectrum of InsH deletions were similar to what we observed at 30°C and did not tend towards the rate and spectrum observed in the pol3-t mutant (Supplementary Figure 3, Supplementary Data S2).

ssDNA in the template for BIR leading strand synthesis is susceptible to APOBEC3A deamination

The high frequency of Type I deletions in Ori1 orientation of insH during BIR, allowed us to hypothesize that ample ssDNA is exposed in the template for BIR within the D-loop structure. Therefore, we asked whether this ssDNA can be detected by its susceptibility to APOBEC-induced damage. Previously, we expressed APOBEC3A (A3A), a cytosine deaminase that converts cytidine in the context of ssDNA into deoxyuridine (dU), in yeast cells undergoing BIR (21). This led to the formation of long mutation clusters on the track of BIR and allowed us to conclude that long stretches of ssDNA accumulate behind the BIR migrating bubble formed by leading strand BIR synthesis and by resection of the DSB end. However, in those studies we never asked whether mutagenic ssDNA can also accumulate within the D-loop bottom template strand (D-BTS). To investigate this, we placed a ura3-29 base substitution reporter cassette (86) at the 16kb position centromere-distal from MATα-inc in the donor chromosome of our disomic galactose-inducible DSB system (Figure 4A). The ura3-29 reporter contains a T to C transition at position 257 in the MATα-inc reporter as compared to BIR-associated mutagenesis in the EV-harboring strains (Figure 4B, Supplementary Data S6). Because the reporter cytosine in Ori1 is most likely included into the ssDNA during BIR when it is formed in the D-BTS region (see Figure 4A, schematic), the observed increase implies that a significant amount of ssDNA is persisting during BIR in the D-BTS region. In strains harboring the ura3-29 reporter in Ori2, we observed a 30-fold increase in Ura⁺ reversion rate during BIR in the presence of A3A as compared to EV (Figure 4B, Supplementary Data S6), consistent with the high mutagenicity of the long persistent NS ssDNA that we previously reported (21). To ensure that the A3A-induced increase seen in Ori1 strains after BIR was not unique to the 16 kb position of the reporter, we repeated the experiment with the Ori1 ura3-29 reporter cassette at a position 90 kb centromere-distal from MAT (Figure 4A). We observed a similar 4.7-fold increase of Ura⁺ during BIR in the presence of A3A as compared to EV (Figure 4B, Supplementary Data S6).

dU lesions in the D-BTS are poor substrates for error-free repair

Previously, we demonstrated that dU generated by A3A in the nascent ssDNA of BIR are frequently converted to AP sites by uracil-DNA glycosylase Ung1, and this conversion leads to reduction of mutagenesis via an error-free repair pathway (21). Here, we asked whether dU lesions generated in the D-BTS (see Figure 4A, schematic) are repaired through error-free pathways equally often. To this end, we tested the effect of deleting UNG1 on A3A mutagenesis during BIR in strains harboring ura3-29 reporter at 16-kb and 90-kb positions. Like what was previously observed for 90-kb position (21), the rate of Ura⁻ reversions increased dramatically (115-fold) in unglΔ Ori2 strains as compared to wild-type (UNG1 Ori2) for 16-kb position (Supplementary Data S6). However, such a dramatic increase was not observed in unglΔ Ori1 strains (Supplementary Data S6). Rather, the 90-kb reporter position produced a rate of Ura⁻ that was only 4.2× higher in unglΔ as compared to UNG1,
which suggests that ∼24% of dUs introduced into the D-BTS led to mutations (as compared to only 5% of dUs that led to mutations when they were introduced into the ss-DNA of the newly synthesized leading strand at the same location (Ori2 90 kb, see in [21])). The data obtained at the 16 kb position support this idea. Specifically, we did not observe an increase of Ura⁺ frequency following BIR in Ori1 ung1Δ strains as compared to the pre-BIR level, even though an increase was expected based on the observed increase in ung1Δ strains containing ura3-29 in Ori2 (Supplementary Data S6). Also, when we analyzed the mutation spectra of BIR/A3A Ura⁺ revertants in UNG1 strains with the reporter at this (16 kb) position, we observed significantly fewer C to G transversions (reflective of translesion
synthesis across from abasic (AP) sites than in Ori2 (Figure 4C). Further, our calculations (based on combining results shown in Figure 4B and C) demonstrated that, while the frequencies of both C to T and C to G substitutions were drastically increased following BIR in the presence of A3A (as compared to EV) in Ori2, only C to T substitutions were increased in Ori1 (BIR/A3A versus BIR/EV; Figure 4D). The lack of C to G increase in Ori1 suggests that dU lesions formed in the D-BTS region during BIR might be converted into AP sites more rarely by Ung1 as compared to lesions introduced into a nascent leading BIR strand (21).

We next asked whether A3A-induced damage in the D-BTS could be a significant contributor to mutagenesis along the entire track of BIR. To address this question, we performed whole genome sequencing (WGS) analysis for the outcomes of BIR exposed to A3A in both UNG1 and unglΔ backgrounds. For these experiments, we combined previously analyzed BIR outcomes from (21) with outcomes from newly performed experiments (see Materials and Methods). We identified all base substitutions on Chr III and called C to N and G to N (with respect to the Watson strand) substitutions separately (see Supplementary Data S7). Because the exact site of invasion during BIR could occur anywhere along the length of resection (which can be up to the centromere (4,60,21)), we considered mutations called on the right arm of Chr. III to be part of the BIR repair track, while mutations called on the left arm were assumed to result from another source, such as S-phase replication (Figure 5A). On Chr III, C to N substitutions on the BIR track indicate mutations incorporated due to A3A damage to the nascent ssDNA that serves as a template for lagging strand synthesis. Meanwhile, G to N substitutions likely indicate mutations incorporated due to A3A damage to the D-BTS ssDNA. In total, based on sequencing of 62 UNG1 BIR outcomes, we observed that the outcomes accumulated a total of 41 G to N mutations on the right arm of chromosome III (Figure 5B). This is comparatively fewer than the 345 C to N mutations observed in the same chromosome region in the same 62 outcomes (Figure 5A, B), consistent with the higher frequency of A3A-induced Ura+ reversions in the Ori2 reporter than in the Ori1 reporter of our UNG1 strains (Figure 4B). In the unglΔ strains, C to N mutations accumulated massively on the BIR track with 2596 mutations across all 45 outcomes (Figure 5B). This accumulation of C to N mutations is in accordance with the idea that the majority of A3A-inflicted lesions (dU) are introduced in the nascent strand (lagging strand template) and that most of these lesions are channeled into an error-free pathway of repair by conversion into AP sites mediated by Ung1, as we previously proposed in (21). By contrast, we observed only 41 G to N mutations on the right arm of chromosome III in the unglΔ background among all 45 outcomes analyzed (Figure 5B). We next compared the number of G to A mutations on the BIR track per outcome between UNG1 (mean of 0.21 mutations per outcome) and unglΔ (mean of 0.82 mutations per outcome) backgrounds and found that loss of UNG1 promoted a significant increase (Figure 5C). In addition, when we compared the number of G to A mutations (mean of 0.21 per outcome) and G to C mutations (mean of 0.35 per outcome) among UNG1 outcomes, there was no significant difference between the two, supporting that dU were efficiently converted into AP sites at least at some of the chromosomal positions (Figure 5D).

Notably, G to N and C to N mutations were nearly absent on the left arm of Chr. III in the UNG1 background, but both were present in the unglΔ background (Figure 5A), indicating that spontaneous dUs responsible for their formation did not frequently lead to mutations in the presence of uracil glycosylase. As a secondary control, we also identified C to N and G to N mutations across the rest of the genome (where BIR synthesis did not take place) in our sequenced outcomes in both UNG1 and unglΔ backgrounds. Particularly in the UNG1 background, these mutations did not occur at the same frequency as those found on the BIR track, which represents only about 1% of the yeast genome (41 mutations observed on the BIR track vs 98 G to N mutations observed across the entirety of the genome (Supplementary Figure S4A; Supplementary Data S7)). Further, we performed simulations to test whether the number of G to N mutations that occur cumulatively across all UNG1 background samples subjected to WGS were likely to occur in the observed frequency at which they exist on the BIR track region of the genome if redistributed randomly across the entire genome (Supplementary Figure 4B). From 100,000 simulations, we saw that most instances had 1–2 G to N mutations on the BIR track and the probability approached 0 between 7 and 8 G to N mutations. This was far fewer than the 41 G to N mutations that we observed in this region among all sequenced UNG1 samples (Supplementary Figure 4B), thus strongly suggesting that these 41 G to N mutations that we report here indeed result from BIR. In addition, we confirmed that the majority of mutations detected in the UNG1 and unglΔ outcomes were part of a TCW motif, a recognition motif for A3A (Supplementary Figure 4C), as expected. From these results, we conclude that the ssDNA of the BIR D-loop is susceptible to A3A-induced damage with each lesion resulting in mutation more frequently than similar lesion introduced into ssDNA of the newly synthesized leading strand (Figure 6, see Discussion).

DISCUSSION

Our results demonstrate that ssDNA formed in the D-BTS region is sufficient to form secondary structures and to incur DNA damage from agents that specifically target persistent ssDNA. This conclusion is based on our observations suggesting that the template for leading strand BIR synthesis is highly vulnerable to A3A damage and also highly susceptible to microhomology-mediated deletions promoted by inverted DNA repeats.

The template for leading strand BIR synthesis is a novel source of mutagenic ssDNA

APOBEC-induced mutagenesis has been an important tool for identifying sources of persistent ssDNA in vivo. Previously, vulnerability to APOBEC-induced mutagenesis helped to identify several important sources of mutagenic ssDNA, including the lagging strand of S-phase DNA synthesis, actively transcribed regions (i.e. tRNA transcription), as well as uncapped telomeres (11,13–15,20,22,24,25,63). Our previous work on BIR (4,21)
Figure 5. (A) A3A-induced mutations at cytosines (C to N: black tick-marks) and guanines (G to A: red, G to C: blue, and G to T: yellow tick-marks) on Chromosome III (Watson or 5' to 3' strand) identified by WGS of BIR outcomes from UNG1 and ung1Δ backgrounds exposed to A3A. Outcomes numbered in blue or green are from independent experiments (see Supplementary Data S7 for details). (B) Summary of total number of G to N and C to N mutations on the right arm (BIR track) in BIR outcomes from UNG1 and ung1Δ backgrounds shown in (A). (C) Number of G to A mutations per outcome on the track of BIR (right arm) in UNG1 and ung1Δ backgrounds. Horizontal lines indicate mean values that are also listed above each plot. P-value obtained by unpaired t-test is listed above the plot to indicate statistical significance (asterisk) or no statistical significance (N.S.) of difference in the mutation number. (D) Number of G to N mutations per outcome (G to A or G to C) on the track of BIR (right arm) in the UNG1 background. Mean values and statistical significance are as in (C).
Figure 6. (A) Cytidine (C*) in the ssDNA of D-BTS is susceptible to A3A-induced deamination producing dU lesions (U*). The fate of dU lesions follows one of two possible paths. In the first (left), the Ung1 enzyme is unable to excise the dU base, leaving it in the template where it is encountered by leading strand synthesis. An A base is placed across from the dU lesion, incorporating it into the nascent strand where it is paired with a T base during lagging strand synthesis. Ung1 may later repair the template strand dU lesion after BIR has proceeded beyond it, but the mutation will stay in the newly synthesized strand. In the second path (right), Ung1 is able to access the dU lesion created by A3A and excise it, leaving an AP-site. This AP-site does not (or rarely) triggers error-free bypass pathways. Instead, a base placed across from the AP site during leading-strand synthesis often results in incorporation of the wrong base into the nascent strand, resulting in a mutation (M). (B) Schematics of Type I InsH deletion following hairpin formation during leading or lagging strand synthesis of BIR. Directions of leading and lagging strand synthesis are defined by the known direction of BIR synthesis for BIR in our system. Microhomologies at InsH deletion breakpoints are indicated in red.

demonstrated that large amounts of stable nascent ssDNA accumulate during BIR leading strand DNA synthesis and following DSB resection preceding BIR. We observed that this ssDNA is susceptible to A3A-inflicted lesions and to alkylating damage, both leading to formation of mutations along the track of BIR that are similar to those termed kataegis that were described in cancer cells (4,5,16). While the majority of mutagenic ssDNA associated with BIR accumulates as the template for lagging strand synthesis, our new data presented here provide the evidence for mutagenic ssDNA formed in the template for the leading strand as well. This mutagenic DNA is formed in shorter stretches (creating isolated mutations rather than mutation clusters) that correspond to the template regions that become single-stranded within a D-loop. This conclusion is based on our analysis of mutation frequencies using the *ura3-29* reporter inserted at 16kb and 90kb positions on the BIR track. It is also supported by the results of our WGS analysis that identified mutations that likely resulted from dUs introduced in the D-BTS through the entire track of BIR.

Our conclusion regarding ssDNA in D-BTS is consistent with the previously published *in vitro* reconstitution
of Polô-driven repair DNA synthesis (87), suggesting that binding of RPA occurs within the D-loop to the template ssDNA and stimulates Polô-driven repair DNA synthesis. The length of such ssDNA-binding RPA was proposed to be at least 30bp (required for binding one RPA molecule). The results of our study suggest that the length of this ssDNA may be much longer (at least 130-150 bp), because deletions of InsH, which provide an estimate of the size of the hairpin structure formed, were usually longer than 120bp. In addition, the results of our in vivo study allowed us to conclude that the ssDNA region within a D-loop not only exists but is likely persistent enough to become a significant source of mutagenesis. It is also possible that ssDNA in the D-BTS arises most readily when BIR proceeds slowly or interrupts. Nonetheless, our observation that D-BTS-associated mutations occur through the entire track of BIR suggests that instances of mutagenic ssDNA in the D-BTS arise often, regardless of whether perturbed BIR synthesis is prerequisite.

Previous studies demonstrated that correction of dUs introduced during replication or transcription usually (in ~95% of the cases) proceeds via error-free repair (20,63). That was also the case for dU lesions introduced by APOBEC in the nascent strand of BIR (21). Here we demonstrate that dUs formed in the D-BTS are less frequently repaired in an error-free fashion and therefore lead to mutations more often. This could be caused by lower uracil glycosylase efficiency in excising of dUs or by lower efficiency of error-free repair pathways (template switching, homologous recombination, etc.) recruited for the repair of AP lesions. Variations in the efficiency of dU correction by Ung1 between different sources of ssDNA have been previously reported. For example, dUs introduced by APOBEC into yeast uncapped telomeres or into the nascent strand of BIR were almost all converted into AP sites by Ung1 as evidenced by the equal numbers of C to T and C to G base substitutions among repair outcomes and by increase in mutation frequency following elimination of Ung1 (11,21,63). Conversely, excision of dUs formed during S-phase replication (primarily in the lagging strand ssDNA) by Ung1 was ~9% less efficient as compared to dUs excised during repair synthesis at uncapped telomeres (63). Our data reported here suggest that decreased efficiency of dU excision contributes to the increased level of mutations resulting from dUs in the D-BTS at the 16kb ura3-29 position. However, because G to A and G to C base substitutions were observed in equal numbers through the track of BIR, it appears that Ung1 is capable of excising dUs at many BIR track locations. More striking, however, is the lack (or only modest increase) of G to N mutation frequencies in ung1Δ as compared to Ung1 through the entire track of BIR. This data suggests that even when dUs in the D-BTS are converted into AP sites, error-free repair of these AP sites is not as efficient as was observed for the nascent BIR strand (Figure 6A versus (21)) or for S-phase lagging strand synthesis (63). Together, we propose that the transient state of ssDNA in the D-BTS region does not provide enough time for conversion of dUs into AP sites or for the channeling of AP sites into error-free repair pathways. An alternative possibility is that it might be difficult for Ung1 or for proteins participating in error-free repair to access their substrates inside of the D-loop, and likewise in the D-BTS region.

Although mutations resulting from D-BTS lesions rarely formed even small clusters, varying densities of G to A mutations along the entire track of BIR might be indicative of fluctuating lengths and persistence of ssDNA formed inside the D-BTS during BIR. Additional work, particularly in situations where D-loop migration is challenged by obstacles that lead to stalling, may elucidate factors capable of altering the amount of ssDNA present or amount of time that it persists within the D-loop, thereby increasing its propensity to accumulate damage, mutations, and mutation clusters. Another important observation from our WGS results was the presence of template strand A3A lesions upstream (centromere-proximal) to the HO cut site. Previously, A3A-induced mutations identified in the region between the centromere and the HO site were ascribed to long resection of DSB ends preceding strand invasion and BIR initiation (4,21). However, all mutations resulting from resection should occur at cytosines, not guanines. Yet, we identified several G to N mutations bearing A3A’s preferential sequence motif on the proposed resection track. We interpret this as an indication of synthesis in this region that leads to formation of these mutations caused by ssDNA lesions inside the D-BTS. This implies that damage to ssDNA exposed by long resection might not always be converted into mutations because the entire ssDNA region is removed if strand invasion occurs centromere-proximal to them. In this case, A3A-induced mutations likely result from synthesis initiated distant from the DSB following extensive resection.

The ssDNA in the D-BTS promotes deletions at microhomologies stimulated by inverted DNA repeats

Here we report that BIR stimulates deletions that are promoted by inverted repeats and occur between microhomologies showing polarity (with one short repeat located inside of the IR and the other outside (see Figures 2E, G, 6B). Similar polarity was previously reported for IR-mediated deletions observed in LYS2 genes during S-phase replication (36). To explain their mechanism, authors proposed that deletions are initiated by hairpin formation involving IRs included into ssDNA regions formed during lagging strand DNA synthesis. Further, it was proposed that displacement DNA synthesis driven by Polb enters the duplex of the hairpin stem, but is unstable, which leads to frequent template switches from microhomology located inside of the hairpin duplex towards microhomology located outside of the hairpin (36). Based on this model, the polarity of deletion breakpoints allowed authors to postulate the direction of replication that led to deletions. Using the same logic, we use the polarity of insH deletion breakpoints that we observed during BIR (Figures 2B, E, G) to deduce the direction of synthesis producing the deletions. The result of our analysis indicates that both leading and lagging strand BIR synthesis promote IR-mediated deletions. This conclusion can be made based on Type I deletion frequencies following BIR in strains containing InsH in Ori1 and Ori2 orientations. Type I deletions were ~5-fold more frequent in Ori2 (where they likely occur during lagging strand synthesis).
synthesis) as compared to Ori1 (leading strand synthesis) (Figures 2D, 6B). This difference is expected as ssDNA in the template for lagging strand BIR synthesis is expected to be longer and more stable as compared to the template for leading strand. Yet, deletions during leading strand synthesis were hundreds of times more frequent as compared to frequencies observed during S-phase DNA synthesis (Figure 1B, 2D, no DSB versus 7 h DSB). Similarly, the contribution of both leading and lagging strand BIR synthesis to the formation of IR-promoted deletions follows from the spectra of deletions observed by deep sequencing following BIR in Ori1 and Ori2 strains. In particular, after BIR, deletions of both polarities were observed (Type I-like (“left” microhomology inside the IR and the “right” microhomology outside—shown in red in Figures 2B, E, G, and 6B) and Type II-like (“right” microhomology inside the IR the “left” microhomology outside—shown in blue). The presence of both polarities following BIR in each strain is consistent with contributions of both leading and lagging strands to the induction of IR-promoted deletions.

Because Pol δ mediates both leading and lagging strand synthesis during BIR (77), and imprecise deletions were the predominant outcomes from both leading and lagging BIR synthesis, it is likely that both cases result from template switching occurring during displacement synthesis by Pol δ that partially opens hairpin stems and then undergoes slippage and switches to microhomology outside of the hairpin. We did not observe any evidence of translesion polymerase participation in the process, which also supports this model where slippage events are entirely mediated by Pol δ. The effect of the pol3-t mutation in this context (including the increase in the overall frequency of InsH deletions, and in the fraction of precise deletions among them) might result from reduced displacement ability of Pol δ in the pol3-t mutant leading to more frequent stalling of Pol δ at the base of hairpins. An additional explanation for the high frequency of imprecise deletions observed during BIR is that they result from the formation of incomplete hairpins due to a limited amount of ssDNA exposed during BIR. This could explain the effects of pol3-t on events attributed to leading strand synthesis as this mutant might increase the length of ssDNA in the D-BTS. However, this cannot explain why BIR lagging strand synthesis does not solely produce precise deletions when the amount of ssDNA accumulated after leading strand synthesis should be sufficient for full hairpin formation, even in POL3 (wt) strains. In addition, our failure to recapitulate the effect of pol3-t in POL3 (wild-type) cells by executing BIR at a lower temperature to slow down BIR synthesis also makes the second explanation less likely, even though it is possible that the decreased temperature slows down not only DNA synthesis but also DNA unwinding. Overall, the first explanation (Pol δ-mediated displacement synthesis as an explanation for imprecise deletions of InsH) appears more likely, even though the latter explanation could represent another contributing factor. It is also possible that ssDNA accumulated in the D-BTS region is less accessible for RPA binding, which could represent an additional factor provoking more efficient hairpin formation. Finally, questions remain about whether the genetic requirements for deletions in pol3-t are different from those in POL3. For example, we cannot exclude that deletions of InsH in pol3-t could be mediated by another polymerase (for example by Polζ).

**BIR-associated mutagenic ssDNA: future questions**

The observations made in this study prompted us to formulate several additional questions.

First, our findings led us to wonder whether D-BTS regions form during other homologous recombination pathways (e.g., during gene conversion (GC)) and are also mutagenic. Because the D-BTS is likely a common feature of GC and BIR, the mutagenic properties of the D-BTS could be shared between these two pathways. It would be especially relevant to investigate this with respect to meiotic recombination events because many D-loops are formed during every meiosis and because mutagenesis associated with meiosis could have severe consequences for progeny, potentially leading to birth defects in humans.

The results obtained in this work might also help in the interpreting of the mechanisms responsible for the formation of APOBEC-induced mutation clusters that are detected in various cancers. For example, it was believed that only non-switching C-coordinated or G-coordinated clusters can be ascribed to BIR (1,17). It is clear from this study that mutagenesis from both leading and lagging BIR strands is expected to produce more complex patterns, such as clusters where most of the mutations are G- or C-coordinated with rare cases of mutations in the opposing base (e.g. rare mutation in G in otherwise C-coordinated clusters and vis versa), similar to what was observed in (17).

**DATA AVAILABILITY**

Raw sequencing reads can be accessed from the NCBI Sequence Read Archive database under bioproject accession number PRJNA821991. In addition, the raw reads that were re-analyzed in this work and were originally from (21) can be accessed from NCBI Sequence Read Archive database under accession number PRJNA517571. All custom code used for the analysis is available through GitHub https://github.com/malkovalab/WGS-A3A-Tools; (https://github.com/malkovalab/DeepSeqTools).

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

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