Genetic Control of Genomic Alterations Induced in Yeast by Interstitial Telomeric Sequences

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ABSTRACT In many organisms, telomeric sequences can be located internally on the chromosome in addition to their usual positions at the ends of the chromosome. In humans, such interstitial telomeric sequences (ITSs) are nonrandomly associated with translocation breakpoints in tumor cells and with chromosome fragile sites (regions of the chromosome that break in response to perturbed DNA replication). We previously showed that ITSs in yeast generated several different types of instability, including terminal inversions (recombination between the ITS and the “true” chromosome telomere) and point mutations in DNA sequences adjacent to the ITS. In the current study, we examine the genetic control of these events. We show that the terminal inversions occur by the single-strand annealing pathway of DNA repair following the formation of a double-stranded DNA break within the ITS. The point mutations induced by the ITS require the error-prone DNA polymerase ζ. Unlike the terminal inversions, these events are not initiated by a double-stranded DNA break, but likely result from the error-prone repair of a single-stranded DNA gap or recruitment of DNA polymerase ζ in the absence of DNA damage.

KEYWORDS recombination; mutagenesis; interstitial telomeric repeats; DNA replication; inversions

T HE ends (telomeres) of the chromosomes have several properties that distinguish them from most genomic sequences. In most organisms, telomeres are replicated by the telomerase ribonucleoprotein complex (Greider and Blackburn 1985). In addition, telomeres have a structure (the cap) that distinguishes the natural end of the chromosome from ends produced by DNA damage, preventing telomere–telomere fusions (Wellinger and Zakian 2012). One last unusual property of the telomeres in some organisms (including Saccharomyces cerevisiae) is their ability to transcriptionally silence genes located nearby (Gottschling et al. 1990).

In yeast, as in most organisms, the telomeric sequences are simple repeats (Wellinger and Zakian 2012). Unlike most organisms, in S. cerevisiae, the repeat is imperfect of the form C1–3A/G1–3T. Wild-type strains have a double-stranded region of this sequence of ~300 bp, and a single-stranded 3′ G1–3T “tail” of ~15 bases (Wellinger and Zakian 2012). There are a large number of proteins that bind directly to telomeric DNA or indirectly to the telomere by binding to other proteins including Rap1p, Cdc13p, Stn1p, Ten1p, Rif1p, Rif2p, Sir2p, Sir3p, Sir4p, Yku70p, Yku80p, Rfa1p, Rfa2p, Rfa3p, Est1p, Est2p, and Est3p, as well as the telomerase RNA TLC1 (Kupiec 2014). In addition to the proteins that usually reside at the telomere, there are proteins that have a more transient interaction [such as Tel1p/Mec1p and the Mre11p/Rad50p/Xrs2p (MRX) complex], present at certain times in the cell cycle or when the telomeres shorten below a critical length.

In many organisms, telomeric sequences can be found embedded within the chromosome. These interstitial telomeric sequences (ITSs) were first detected by in situ hybridization (Meyne et al. 1990; Azzalin et al. 1997) and, subsequently, by DNA sequencing. The human genome contains > 100 ITSs that have a minimum repeat length of four repeats (Azzalin et al. 2001). Two lines of evidence...
suggest that ITSs may represent breakage-prone sites in the mammalian genome (Lin and Yan 2008). First, ITSs are significantly overrepresented in translocation breakpoints observed in tumor cells. Second, fragile sites (regions of the genome prone to breakage in conditions of DNA replication stress) are enriched for ITSs.

We have recently begun to investigate the effect of ITSs on genome stability in the yeast S. cerevisiae (Aksenova et al. 2013, 2015). Using a system that will be described in detail in the Results section, we showed that ITSs stimulated two types of genomic alterations: point mutations near the site of the ITS insertion and chromosome rearrangements. The most common rearrangement was a terminal inversion between the ITS and the true chromosome telomere. In our 2013 study, we restricted our analysis to a wild-type strain. In the current study, we examine the rates of ITS-induced genomic alterations in strains with mutations affecting DNA repair/recombination and telomere metabolism. We show that ITS-induced point mutations require the error-prone DNA polymerase ζ and are not initiated by a double-strand DNA break (DSB). In contrast, the terminal inversions are likely initiated by a DSB and are repaired by a single-strand annealing (SSA) pathway.

Materials and Methods

Yeast strains and plasmids

The genotypes and details of construction for the strains used in this study are presented in Supplemental Material, Table S1, and the primers used in strain construction/strain analysis are in Tables S2 and S3. All strains are derivatives of the previously described haploid SMY749 (Aksenova et al. 2013). This strain is isogenic with the commonly-used lab strain S288c and has the genotype: MATα leu2-Δ1 trp1-Δ63 ura3-52 his3-200 ade2Δ::kanMX4 III(75594–75641)::URA3-Int-(TGTGTGGG)15-TRP1; the reporter gene with the ITS is located on chromosome III (Figure 1A). As described in Table S1, we isolated a derivative of SMY749 (PG329) in which we replaced ura3-52 with a drug resistance marker (natMX4). Mutant derivatives of PG329 lacking various genes involved in DNA repair, recombination, or telomere elongation were also generated (Table S1). The deletions were constructed by standard procedures using PCR fragments containing drug resistance markers; these fragments were obtained by amplifying plasmids containing either hphMX4 or natMX6 as described in Table S1. All insertions and deletions were confirmed by PCR using primers described in Table S3.

Yeast media

In most experiments, standard media were used (Guthrie and Fink 1991). Medium with 5-fluoro-orotate (5-FOA) contained 1 g of 5-FOA and 40 mg of uracil per liter in synthetic dextrose (SD) complete (SD-complete) medium. Medium containing 5-FOA and canavanine had the same amount of uracil and 5-FOA, but contained 120 mg of canavanine per liter; in addition, the medium lacked arginine. The medium used to check for silencing of TRP1 expression was SD-tryptophan medium containing 70 mg of nicotinamide per liter. Sporulation medium was standard (Guthrie and Fink 1991).

Measurements of rates of genetic alterations in strains with URA3-Int-(TGTGTGGG)15-TRP1 on chromosome III

We measured several types of genetic instability in our analysis. In most of the studies, we first determined the rate of 5-FOAR (5-FOA-resistant) derivatives in strains that contained the URA3-Int-(TGTGTGGG)15-TRP1 reporter gene integrated on chromosome III. For these experiments, the strains were streaked on rich growth medium (YPD) and allowed to form colonies at 30°C. Individual colonies were then grown as small patches on YPD for 1 day. Individual patches were resuspended in water and various dilutions were plated to SD-complete or SD medium containing 5-FOA. In each experiment, we examined 15–20 independent colonies and we performed two separate experiments for each strain. Based on the total number of cells in the patch and the number of 5-FOAR derivatives, we calculated a rate of formation of 5-FOA resistance for each strain using the method of the median (Lea and Coulson 1949).

We found three classes of 5-FOAR derivatives: class 1 (point mutations within the URA3 reporter), class 2 (terminal inversions involving the ITS), and class 3 (uncharacterized chromosome rearrangements). For each strain, we examined at least 20 independent 5-FOA derivatives using a series of tests. First, we performed PCR with all the isolates using the primers UIRL1/UIRL2 and UIRL2/CHR413R (Figure 1A). UIRL1/UIRL2 flank the ITS and produce a PCR product if there is a point mutation (class 1). The UIRL2 and CHR413R primers produce a PCR product if there is a terminal inversion (class 2). Failure to detect a PCR product with either set of primers indicates class 3. Second, we tested whether the 5-FOAR derivatives failed to grow in medium lacking tryptophan but could grow in medium lacking tryptophan that contained nicotinamide. Because terminal inversions result in a larger insertion of telomeric sequences near the TRP1 gene, the TRP1 gene is silenced epigenetically, and this silencing is reversible by nicotinamide (Aksenova et al. 2013). In contrast, most class 3 strains were Trp−, but the tryptophan requirement could not be reversed with nicotinamide. Using the criteria described above, we found that >90% of the strains were classes 1 and 2, with class 3 representing a minor fraction. We calculated the rates of formation of classes 1 and 2 by multiplying the proportion of these events among the 5-FOAR derivatives by the rate of 5-FOA.

Determination of 95% confidence limits on rate measurements

For the calculation of 95% confidence limits (CL) on the median rates of 5-FOAR derivatives, we used the ranking method described previously (Wierdl et al. 1996). The procedure for
ITS, interstitial telomeric sequence.

a consequence of epigenetic silencing of when the cells are analyzed on plates lacking tryptophan but containing the Sir2p-inhibitor nicotinamide. 5-FOA, 5-FOA-resistant; DSB, double-strand break; (TGTGTGGG). Strains with this reporter are Ura+ Trp+, and we selected 5-FOA− derivatives to detect genomic alterations. (B) Point mutations induced by the ITSs are inserted within an intron embedded with URA3; the length of the ITS is 120 bp, representing 15 copies of an 8-bp repeat (TGTGTGGG). Strains with this reporter are Ura+ Trp+, and we selected 5-FOA− derivatives to detect genomic alterations. (B) Point mutations induced by the ITSs. These derivatives are Ura+ Trp+ and have a PCR fragment of the expected size when the purple primers flanking the ITS are used (URL1 and URL2 in Table S2). Sequencing of PCR fragments demonstrates that these events are associated with a point mutation in the URA3 coding sequences flanking the ITS. (C) Terminal inversions induced by the ITSs. In this class, no PCR fragment is found with the purple primers, but a PCR fragment is observed using the orange primer (Chr3_413R in Table S2) and the centromere-proximal purple primer (UIRL2). In addition to the 5-FOA− phenotype, most of this class are Trp−, as a consequence of epigenetic silencing of URA3 caused by the longer region of telomeric repeats resulting from the inversion; the Trp− phenotype is reversed when the cells are analyzed on plates lacking tryptophan but containing the Sir2p-inhibitor nicotinamide. 5-FOA+, 5-FOA-resistant; DSB, double-strand break; ITS, interstitial telomeric sequence.

determining the 95% CL of the rates of class 1 and class 2 isolates is more complex because both the CLs on the rates of 5-FOA8 and on the proportions of each class must be considered. The 95% limits on the proportions of each class were calculated using the Vassarstat website (http://vassarstats.net). If the rate of class 1 or class 2 events is Q, the rate of 5-FOA8 is A, and the proportion of class 1 or class 2 events is B, we calculate the error (confidence interval) for Q (δQ) by the equation:

\[
\delta Q = Q \times \sqrt{\left(\frac{\delta A}{A}\right)^2 + \left(\frac{\delta B}{AB}\right)^2}
\]

(http://ipl.physics.harvard.edu/wp-uploads/2013/03/PS3_Error_Propagation_sp13.pdf).

To illustrate the method used to calculate the 95% CL of class 1 and class 2 events, we will use the data from the wild-type strain PG329 as an example. The median rate of 5-FOA8 derivatives was 18.6 \times 10^{-7} (95% CL of 14–24 \times 10^{-7}); the median rate and the 95% CL on the rate were calculated as described by Wierdl et al. (1996). Of 107 5-FOA8 mutants examined, 52 had point mutations; this proportion is 0.486 (95% CL of 0.39–0.58). The 95% limits on the proportions were calculated using the “The Confidence Interval of a Proportion” section of the Vassarstat website. The rate of point mutations was 18.6 \times 10^{-7} \times 0.486 = 9.0 \times 10^{-7}. To obtain the upper CL limit on this rate, we used the following protocol. First, (step 1) subtract the median rate of 5-FOA8 from the rate representing the upper 95% CL (24 \times 10^{-7}–18.6 \times 10^{-7} = 5.4 \times 10^{-7}); (step 2) divide the value of step 1 by the median rate of 5-FOA5 [(5.4 \times 10^{-7})/(18.6 \times 10^{-7}) = 0.30]; and (step 3) square the value obtained in step 2; [(0.30)^2 = 0.09]. For steps 4–6, perform the comparable steps with the data for the proportion: (step 4) 0.58–0.486 = 0.094; (step 5) (0.094)/(0.486) = 0.19; and (step 6) (0.19)^2 = 0.036. For step 7, add the values calculated for steps 3 and 6 (0.09 + 0.036 = 0.126). For step 8, take the square root of the value calculated for step 7 [(0.126)^{1/2} = 0.35]. For step 9, multiply the value obtained in step 8 by the rate of point mutations (0.35 \times 9.0 \times 10^{-7} = 3.15 \times 10^{-7}). The upper 95% CL is equal to the median rate of the point mutations plus the value calculated in step 9 (9.0 \times 10^{-7} + 3.15 \times 10^{-7} or 12.2 \times 10^{-7}). Performing comparable steps for the lower 95% CL, we calculate a value of 6.1 \times 10^{-7}. In summary, for point mutations, the rate is 9.0 \times 10^{-7} (95% CL of 6.1–12.2 \times 10^{-7}). Similar calculations for the terminal inversions yield a rate of 9 \times 10^{-7} (CL of 5.9–12 \times 10^{-7}). These values and those for the mutant strains are in Table 1.

Data availability

All data and strains used in this study are available on request. Supplemental material available at Figshare: https://doi.org/10.25386/genetics.6013667.

Results

Experimental system

The system used by Aksenova et al. (2013) and in the present study is shown in Figure 1A. The reporter URA3 gene has an intron with 15 copies of the yeast telomeric repeat (TGTGTGGG). This gene (URA3::Int (TGTGTGGG)15) was inserted on chromosome III near ARS306 \sim70 kb from the left end. We determined the mutation rate of this gene by measuring the frequency of 5-FOA5, a phenotype associated with ura3
mutant strains (Boeke et al. 1984). The purple and orange arrows in Figure 1 show the location of primers that were used to diagnose the genomic alterations (as described in the section below).

The rate of 5-FOAR was elevated by the ITS ~100-fold compared to the rate for an intron-containing URA3 in the same location without the ITS (Aksenova et al. 2013). About half of these Ura− derivatives had point mutations in ura3 (Figure 1B) and the remainder had chromosome alterations of several different types. A common chromosome alteration was a terminal inversion of the sequences located between the ITS and the true telomere; we assume that this rearrangement was initiated by a DSB within the ITS (Figure 1C). We previously observed chromosome rearrangements in addition to the terminal inversion; some of these rearrangements involved an interaction between Ty elements located near the ITS on chromosome III and the Ty insertion in the ura3 gene (ura3-52) on chromosome V (Aksenova et al. 2013). To avoid getting this class of rearrangement, in the current study, we deleted the ura3-52 allele. In this strain (PG329), as described below, almost all (> 90%) of the 5-FOAR derivatives had point mutations (class 1) or terminal inversions (class 2).

### ITS-induced genomic alterations in the wild-type strain PG329

The rate of 5-FOAR in the wild-type strain was 1.9 × 10⁻⁶/division (95% CL of 1.4 × 10⁻⁶) (Table 1), 52-fold higher than the rate in an isogenic strain (AM21) with the same URA3 reporter gene lacking an ITS [3.5 × 10⁻⁸ (1.7–23 × 10⁻⁸)]. To confirm that ITS-induced mutagenesis in PG329 occurred locally rather than on a genome-wide basis, we examined mutation rates at the CAN1 locus in PG329 and in a strain lacking the ITS insertion (SMY803). The mutation rate to canavanine resistance was very similar in the two strains: 2.1 × 10⁻⁷/division (95% CL of 1.1–3.6 × 10⁻⁷) in PG329 and 1.4 × 10⁻⁷/division (95% CL of 0.6–2.6 × 10⁻⁷) in SMY803.

Three classes of genomic alterations were found among 107 independent 5-FOAR derivatives that were examined. Class 1 strains (52 of 107) were Trp+ on standard omission plates (Table 1). 5-FOAR derivatives in this class were divided into two subgroups based on their resistance to 5-FOAR: 27 (26%) were resistant to 5-FOAR (class 1a) and 25 (24%) were resistant to 5-FOAR but had a lower resistance than the former group (class 1b). Class 1 derivatives included at least 13 different types of mutations: point mutations, insertions, deletions, and inversions. The frequency of these mutations was highest in class 1b, suggesting that these derivatives arose from different types of rearrangements than those in class 1a.

### Table 1 Rates of ura3 point mutations and terminal inversions in strains of different genotypes

| Genotype* | Rate of ura3 point mutations (class 1) (μ × 10⁻⁷) | Rate of class 1 normalized to wild-type | Rate of terminal inversions (class 2) (μ × 10⁻⁷) | Rate of class 2 normalized to wild-type |
|-----------|-----------------------------------------------|----------------------------------------|-----------------------------------------------|----------------------------------------|
| Wild-type | 9.0 (6.1–12) | 1 | 9.0 (5.9–12) | 1 |
| etg1Δ     | 11 (5.5–19) | 1.2 | 5.2 (2.1–10) | 0.6 |
| exo1Δ     | 18 (11–25) | 1.9* | 15 (9.1–22) | 1.2 |
| lig4Δ     | 9.2 (5.4–13) | 1 | 9.2 (5.4–13) | 1 |
| mms2Δ     | 6.4 (3.4–11) | 0.71 | 9.7 (5.4–13) | 1.1 |
| mre11Δ    | 1.1 (0.6–3.6)** | 0.12 | 1.3 (0.7–4.2)** | 0.14 |
| mre11-H125N | 7.9 (4.4–4.9)** | 0.32 | 7.5 (4.5–9.8) | 0.83 |
| mshΔ      | 9.0 (6.2–13 [0.98] | 0.98 | 0.6 (2.1–1.4)** | 0.07 |
| msh6Δ     | 8.7 (5.5–14) | 0.97 | 7.8 (4.8–13) | 0.9 |
| mus81Δ    | 14 (8.2–22)* | 1.5 | 6.9 (3.8–11) | 0.77 |
| rad1Δ     | 14 (8.2–22)* | 1.5 | 1.0 (4.4–3.0)** | 0.15 |
| rad18Δ    | 5.6 (2.2–9.1)* | 0.62 | 8.9 (3.5–12) | 1 |
| rad50Δ    | 3.1 (2.0–4.1)** | 0.34 | 0.9 (0.4–1.8)** | 0.1 |
| rad51Δ    | 19 (12–26)* | 2.1 | 13 (7.4–20)** | 1.5 |
| rad52Δ    | 26 (50–51)** | 4 | 1 (0.07–6)* | 0.1 |
| rad59Δ    | 13 (8.9–19)* | 1.4 | 2.2 (0.9–4.7)** | 0.25 |
| rev3Δ     | 0.9 (0.4–1.9)** | 0.1 | 4.4 (3.1–7.2)* | 0.49 |
| rflΔ      | 11 (8.3–22) | 1.2 | 12 (4.7–16) | 1.3 |
| rrm3Δ     | 11 (7.6–15) | 1.2 | 2.1 (0.8–4.4)** | 0.24 |
| sae2Δ     | 3.4 (1.6–6.6)* | 0.37 | 12 (7.1–18) | 1.3 |
| sgs1Δ     | 14 (6.3–23)* | 1.6 | 1.3 (0.1–3.8)** | 0.2 |
| sir2Δ     | 3.2 (1–8) [0.4]* | 0.4 | 27 (20–34)** | 3 |
| sir3Δ     | 0.55 (0.1–1.9)** | 0.06 | 8.3 (6.3–11) | 0.92 |
| sml1Δ     | 12 (6.8–19) | 1.3 | 11 (6.3–18) | 1.3 |
| srs2Δ     | 7.1 (4.6–9.4) | 0.77 | 2.0 (1–3.3)** | 0.22 |
| tel1Δ     | 17 (11–26)* | 1.9 | 23 (11–33)** | 2.6 |
| tof1Δ     | 3.4 (1.7–6.8)* | 0.38 | 2.1 (1.7–6.8)* | 0.38 |
| tof1Δ mre11Δ | 5.7 (4.1–7.1)* | 0.3 | 0 (0–0.8)* | 0 |
| top1Δ     | 6.1 (3.8–8.7) | 0.67 | 7.2 (4.6–10) | 0.8 |

Rates were calculated as described in the text. 95% confidence limits are shown in parentheses and the rates normalized to the wild-type rate are shown in squared brackets. Single asterisks indicate that the rate is outside of the 95% confidence limits of the wild-type strain, and double asterisks show that the rate is higher or lower than the comparable wild-type rates, respectively.

*Strain names for the various genotypes listed from top to bottom are: PG329, MD651, MD653, MD687, AM17, AM13, MD740, AM24, MD659, MD658, MD689, AM26, MD691, MD655, MD649, MD686, AM12, MD647, AM14, MD708, MD688, MD657, MD736, MD646, AM15, MD644, MD674, MD734, and MD661. Genotypes shown in bold are the ones used most extensively in developing our models for the genetic regulation of interstitial telomeric sequence-induced instability.
UIRL2 flanking the ITS (shown in Figure 1B). The ura3 genes from 29 class 1 strains were sequenced and all had mutations in the coding sequence flanking the ITS insertion (Figure S1). Genomic DNA samples from class 2 strains (51 of 107) had a PCR fragment when analyzed with the primers CHR413R and UIRL2 (Figure 1C), indicating that these strains had a terminal inversion. These strains were Trp− on standard omission medium, but Trp+ on medium lacking tryptophan but containing nicotinamide; nicotinamide is a drug that reverses silencing induced by telomeric repeats (Bitterman et al. 2002). These phenotypes are also diagnostic of a terminal inversion because such inversions resulted in a larger insertion of telomeric sequences near the TRPI gene, producing an epigenetic silencing of TRPI (Aksenova et al. 2013). Class 3 strains are Trp+ on standard omission medium and fail to produce a PCR product with either set of primers used to diagnose class 1 and 2 strains. Since class 3 strains represent < 10% of the events (Table S4), we did not further characterize them. Based on our previous study (Aksenova et al. 2013), most of the class 3 strains likely reflect recombination between Ty elements located near URA3::Int (TGTGTGGG)15 and other Ty elements in the genome. In summary, the ITS-induced genomic alterations in PG329 were approximately equal numbers of point mutations and terminal inversions, with a much lower frequency of other types of events.

Genetic regulation of ITS-induced genomic changes: general considerations

We constructed a series of strains isogenic with PG329 except for mutations affecting various pathways of DNA repair, DNA replication, and telomere metabolism. For each strain, we measured 5-FOAR in two independent experiments, each involving at least 15 cultures. At least 20 independent 5-FOAR isolates were classified as class 1, 2, or 3 by the tests described above. Based on these values, we calculated the rates of formation of class 1 and 2 events for each mutant strain (Table 1).

It is likely that both class 1 and 2 events are initiated by a DNA lesion (DSB or single-stranded nick) or a stall of the DNA replication fork at the ITS, since the ITS does not result in a mutator phenotype at other genomic loci; the nature of the initiating event may be different or the same for class 1 and 2 events. Following the initiating event, the subsequent mechanisms that produce class 1 and 2 events are likely quite different, although some of the same enzymes may be involved. The frequencies of the two classes of events can be regulated either at the level of initiation or at subsequent steps.

The genes that we examined for their effects on ITS-mediated instability and the functions regulated by the proteins encoded by these genes are in Table 2. We analyzed proteins involved in regulating the stability of the replication fork (Rrm3p, Sgs1p, Srs2p, and Tof1p), since previous studies showed that some genome-stabilizing sequences resulted from problems related to DNA replication (Anand et al. 2012; Shah et al. 2012). Since the binding of sequence-specific proteins can affect the frequency of fork stalling, we analyzed mutants affecting telomere-binding proteins (Rif1p, Sir2p, Sir3p, and Tel1p). We studied mutations affecting the repair of DSBs by homologous recombination (HR) (Exo1p, Mre11p, Mus81p, Rad1p, Rad50p, Rad51p, Rad52p, Rad59p, and Sae2p) and by nonhomologous end-joining (Lig4p), since terminal inversions are likely to reflect a recombinational interaction between the ITS and the telomere. We also examined mutants that affect the ratio of error-free (Mms2p and Rad18p) to error-prone (Rev3p) postreplication repair, since such mutants could affect the relative frequencies of the class 1 and 2 events. Lastly, we examined strains with defective mismatch repair (Msh2p and Msh6p) to determine whether the class 1 mutations were subject to mismatch repair.

Genetic regulation of class 1 events

In some previous studies of mutations induced by trinucleotide repeats or palindromic sequences, the induction of mutants in nearby sequences was dependent on the error-prone DNA polymerase ζ (Shah and Mirkin 2015). We found that the rev3 mutation (eliminating the catalytic subunit of DNA polymerase ζ) reduced the rate of class 1 events ~10-fold (Table 1). DNA polymerase ζ is recruited to the sites of DNA damage as well as stalled replication forks that lack overt damage (Northam et al. 2010, 2014; Makarova and Burgers 2015). ITSs in yeast were previously associated with both stalled replication forks and a low frequency of DSB formation (Anand et al. 2012; Goto et al. 2015).

To determine whether the class 1 events were associated with repair of a DSB, we examined the length of the ITSs in strains with a class 1 event. The rationale for this approach is shown in Figure 2A. In our previous study of point mutations induced by a (GAA)n tract, we found that > 80% mutations were associated with a loss of repeats from the (GAA)n tract. Since these tracts also resulted in high levels of DSBs (Tang et al. 2011), to account for these findings, we suggested that the mutagenesis was initiated by a DSB within the GAA tract, followed by processing of the broken ends by 5′–3′ degradation (Tang et al. 2013). Reannealing of these broken ends, and repair of the two single-stranded gaps by DNA polymerase ζ, would result in an elevated rate of mutations in the linked reporter gene and a decrease in the length of the GAA tract (Figure 2A). To determine whether a similar mechanism explained the ITS-mediated class 1 events, we examined the length of the ITS in 37 strains that had a class 1 event. No alterations were observed in 34 strains, two gained one copy of the telomeric repeat, and one gained three copies. These results argue that class 1 events are not likely to reflect repair of an ITS-associated DSB. Alternatives are that the class 1 events result from the recruitment of error-prone DNA polymerase to ITS-associated single-stranded gaps (Figure 2B) or from the recruitment of the error-prone polymerase to a slow-moving or stalled replication fork in the absence of DNA damage (Figure 2C).

Of the other mutants examined, loss of the Tof1, Mre11, Rad50, and Sae2 proteins reduced class 1 events three-
Table 2 Functional roles of proteins examined in this study

| Protein                  | Enzymatic activity: pathway involvement in mitosis                                                                 |
|--------------------------|------------------------------------------------------------------------------------------------------------------|
| Elg1p<sup>a,b</sup>     | Subunit of replication factor C complex: involved in stability of replication fork, HR, and telomere maintenance  |
| Rrm3p<sup>b</sup>        | DNA helicase: relieves replication fork pauses at G4 motifs, tRNA genes, and other fork-stalling sequences       |
| Sgs1p<sup>c</sup>        | DNA helicase: involved in unwinding of HR intermediates and intermediates formed during DNA replication, unwinds G4 structures; in complex with Dna2p, Top3p, and Rlp1 promotes 5’-3’ resection |
| Srs2p<sup>b</sup>        | DNA helicase: negative regulator of HR at replication fork, promotes resolution of HJs by Mss80p/Mms4p, unwinds triplet repeat hairpins |
| Tof1p<sup>c</sup>        | Subunit of replication-fork-pausing complex: stabilizes replication fork and facilitates progression of fork to prevent genomic alterations; S-phase checkpoint function |
| Top1p<sup>d</sup>        | Topoisomerase I: involved in DNA replication, recombination, and transcription                                       |
| Exo1p<sup>b</sup>        | 5’-3’ exonuclease, flap-endonuclease: involved in processing of broken ends, DNA mismatch repair, error-free postreplication pathway, and telomere maintenance |
| Mre11<sup>f</sup>        | Nuclease subunit of Mre11p/Rad50p/Xrs2p complex: processing of DNA ends resulting from DSB, roles in HR and NHEJ, telomere maintenance, and checkpoint functions. |
| Mus81p<sup>f</sup>       | Interacts with Mms4p to form structure-specific endonuclease: promotes reciprocal crossovers in HR pathway         |
| Rad10p<sup>c</sup>       | Single-stranded endonuclease: NER and SSA pathway of HR                                                         |
| Rad50p<sup>c</sup>       | Part of complex with Mre11p/Rad50p: processing of DNA ends resulting from DSB, roles in HR and NHEJ, telomere maintenance, and checkpoint functions. |
| Rad51p<sup>c</sup>       | Strand-exchange protein: most HR pathways except SSA                                                              |
| Rad52p<sup>c</sup>       | Stimulates Rad51p and anneals single-stranded DNA: all HR pathways                                               |
| Rad59p<sup>c</sup>       | Paralog of Rad52p: stimulates SSA                                                                                    |
| Sae2p<sup>f</sup>        | Endo- and exonuclease: roles in resection needed at telomeres and at broken DNA ends for DSB repair by HR, removal of MRX complex from DNA ends |
| Smi1p<sup>f</sup>        | Inhibitor of ribonucleotide reductase: DSB repair                                                                 |
| Lig4p<sup>f</sup>        | Specialized DNA ligase: NHEJ pathway                                                                                 |
| Mms2p<sup>p</sup>        | Ubiquitin-conjugating enzyme: involved in error-free postreparation repair                                         |
| Rad18p<sup>f</sup>       | E3 ubiquitin ligase: postrepetition repair                                                                         |
| Rev3p<sup>f</sup>        | Catalytic subunit of DNA polymerase ζ: error-prone bypass of lesions during DNA replication                       |
| Msh2p<sup>c</sup>        | Mismatch-binding protein: DNA mismatch repair, processing of DNA branches in SSA pathway                           |
| Msh6p<sup>c</sup>        | Mismatch-binding protein acting with Msh2p: Repair of base-base mismatches                                       |
| Rif1p<sup>e</sup>        | Rap1p-binding protein: mutant results in increased telomere silencing and elongated telomeres                     |
| Sir2p<sup>b</sup>        | Histone deacetylase: regulates silencing at HML, HMR, telomeres, and rDNA; regulates life span, negative regulator of DNA initiation |
| Sir3p<sup>b</sup>        | Silencing protein interacting with Rap1p and Sir2p: regulates silencing at HML, HMR, and telomeres, but not in DNA |
| Tel1p<sup>h</sup>        | Protein kinase: telomere length regulation and DNA damage checkpoint                                              |

DBS, double-strand break; HJ, Holliday junction; HR, homologous recombination; MRX, Mre11p/Rad50p/Xrs2p; NER, nucleotide excision repair; NHEJ, nonhomologous end-joining; SSA, single-strand annealing.

a The functions for these proteins were determined using data in the Saccharomyces Genome Database (https://www.yeastgenome.org).

b These proteins are involved in stabilization of the replication fork and mutants lacking these proteins have elevated rates of genome instability.

c The proteins directly or indirectly affect the repair of double-strand DNA breaks by homologous recombination.

d This enzyme is required for “classic” nonhomologous end-joining.

e These proteins are utilized in error-free postreparation repair.

f Rev3p is required in the error-prone pathway of postreparation repair.

g Both Msh2p and Msh6p are required for the repair of base-base mismatches.

h These proteins affect telomere length and/or telomere silencing.

Since the tof1 and mre11 deletions lowered the rate of class 1 events, we also examined the rate of class 1 events in tof1 mre11 double mutants. We found that the tof1 mre11 strain had a rate of class 1 events that was similar to the tof1 single mutant (Table 1). Although this result suggests that Mre11p and Tof1p may operate in the same pathway to promote class 1 events, this interpretation is complicated by the observation that mre11 mutants themselves are strong mutants, elevating mutation rates ~10-fold (Huang et al. 2003). In addition, since both Tof1p and the MRX complex have multiple cellular roles, other interpretations of this result are also possible.

Mutations in both sir2 and sir3 reduced the rate of class 1 events twofold and 17-fold, respectively (Table 1). Although there are several possible interpretations of this result, we suggest that the Sir2 and Sir3 proteins, which bind to telomeres, may also bind to the ITS, inducing partial
silencing. This partial silencing may contribute to fork stalling at the ITS. If fork stalling is reduced in the sir2 and sir3 mutants, the MRX complex and DNA polymerase ζ may be recruited less often, resulting in a lower level of ITS-associated mutations. We note that the postulated binding of Sir2p and Sir3p to the ITS is not sufficient to turn off expression of the URA3 reporter gene or TRP1 gene; however, the expansion of the ITS that occurs in class 2 strains does silence TRP1 expression (Aksenova et al. 2013).

Loss of the Rad52p resulted in a fourfold elevation of class 1 events. Previously, rad52 strains have been shown to have a strong global mutator phenotype (Huang et al. 2003), likely reflecting the channeling of the repair of DNA lesions from the error-free HR pathway to error-prone pathways (Kunz et al. 1989). As expected from this model, the rad52 mutation also substantially decreased the level of class 2 events, as will be discussed in detail in the next section.

None of the other mutants examined had a strong effect on the rate of class 1 alterations. Although the lack of effect of the mismatch repair (MMR) mutations msh2 and msh6 may seem surprising, the mutator phenotype of these strains in our genetic background at the CAN1 locus is small (sixfold elevation in rates in the msh2 strain AM24 compared to the wild-type strain PG329) compared to the 100-fold stimulation of mutations by the ITS. This result is consistent with the Rev3-dependence of class 1 events, since the mismatch repair system fails to efficiently correct mutations caused by DNA polymerase ζ (Lehner and Jinks-Robertson 2009). In summary, our analysis suggests that the ITS stimulates point mutations by a mechanism that may involve an endonucleolytic nick of the ITS, followed by gap formation and recruitment of the error-prone DNA polymerase ζ. A further description of this model, including various caveats, will be given in the Discussion.

Figure 2 Point mutations induced within (GAA)n repeats and by ITSs. The chromosomes containing the repeats are shown as double-stranded DNA molecules with arrows marking the 3’ ends. In all of the proposed mechanism, the mutant base is introduced into only one strand of the duplex, and replication of the resulting DNA molecule produces one mutant gene and one wild-type gene. (A) Point mutations induced by (GAA)n repeats (Tang et al. 2013). As described in the text, (GAA)n tracts are frequently broken in both replicating and nonreplicating yeast cells. DSB formation, followed by 5’–3’ resection of the broken ends, results in products that can reanneal. Reannealing, followed by gap repair with an error-prone DNA polymerase ζ (Rev3p), could produce a mutation in one strand of the linked URA3 gene. Replication of this product would result in one Ura+ and one Ura– cell. This mechanism would result in loss of (GAA) repeats. (B) Mutations introduced by single-stranded gap formation. We suggest that the ITS can be nicked by the Mre11p-associated endonuclease. This nick could be extended into a gap by the 3’–5’ Mre11p-associated nuclease and the 5’–3’ activities of Exo1p or Dna2-Sgs1-Top3p-Rmi1 complex. The resulting gap is filled in by the Rev3p-associated error-prone DNA polymerase. This process could occur outside of the S-period or be associated with a stalled replication fork. No alteration in the size of the ITS is expected from this mechanism. (C) Mutations introduced in the absence of a DNA lesion. In this model, the error-prone DNA polymerase is recruited to a stalled replication fork in the absence of a DNA lesion. This type of recruitment has been observed to be associated with certain DNA polymerase mutations (Northam et al. 2010). DSB, double-strand break; ITS, interstitial telomeric sequence.
**Genetic regulation of class 2 events**

Class 2 events represent HR between the ITS and the left telomere. Most mitotic recombination events are likely initiated by DSBs, although some events appear to be initiated by single-strand nicks or gaps (Symington et al. 2014). The genetic regulation of class 2 events is likely to be more complicated than for class 1 events for several reasons. First, these events reflect the interaction between two substrates, the ITS and the telomere, rather than an event triggered solely by the ITS. Second, most HR pathways involve multiple steps (end resection, strand invasion, and junction resolution), each step requiring multiple proteins (Symington et al. 2014). For these reasons, we expected to find many genes that would reduce or elevate the rate of class 2 events.

Before discussing the effects of individual mutations, we should emphasize that the rate of class 2 events can be regulated either by affecting the rate of DSB formation or by affecting the efficiency with which a DSB is processed to yield a terminal inversion. In regard to the second point, a DSB within the ITS has at least four potential fates (Figure 3): (a) repair of the DSB to produce a terminal inversion, (b) rejoining of the broken ends by reannealing the ITs, thus restoring the original arrangement of chromosomal sequences, (c) repair of the DSB utilizing the sequences on the sister chromatid, or (d) failure to repair the DSB, leading to cell death. Since we can assay only the first of these four pathways, a mutation that alters the rate of one of the three undetectable pathways may indirectly affect the rate of the terminal inversions. This issue will be discussed further below.

The *lig4* mutation that eliminates nonhomologous end-joining had no effect on the rate of class 2 events, whereas mutations in the HR pathway (*MRE11*, *RAD50*, *RAD52*, and *RAD59*, but not *RAD51*) reduced the rate of these events (Table 1). The lack of an effect of the *rad51* mutation, in contrast to the effects of the *rad52* and *rad59* mutations, strongly argues that the terminal inversions reflect SSA (Symington et al. 2014). SSA is usually assayed by inducing a DSB in single-copy sequences located between two direct repeats (Figure 4A). Following end resection, the complementary sequences of these repeats anneal. Since this process involves annealing, but not strand invasion, Rad52p, but not Rad51p, is required (Ivanov et al. 1996). The single-stranded branches of the intermediate are removed by the Rad1, Rad10, Msh2, and Msh3 proteins (Symington et al. 2014). We suggest that class 2 events are generated by a similar process (Figure 4B). As expected from this model, the rate of class 2 events is significantly reduced by *rad1* or *msh2* mutations (Table 1). The structure-specific nuclease Mus81p has no role in the SSA pathway and, as expected, had no effect on the rate of class 2 events (Table 1).

The 7–10-fold reduction in rates of class 2 events in the *mre11* and *rad50* strains could reflect either a role of the MRX complex in DSB formation at the ITS or the role of these proteins in end resection. However, since the nuclease-dead *mre11-H125N* allele does not reduce levels of class 2 events, the MRX complex is unlikely to be directly involved in making the DSB. Similarly, the lack of effect of the *sae2* mutation suggests that the nuclease activity of Mre11p does not contribute to DSB formation at the ITS. An alternative possibility is that the MRX complex involves in a subsequent step in class 2 events. The MRX complex has a structural role in bridging DNA ends, and this function is deficient in *mre11Δ* strains but unaffected in *mre11-H125N* strains (Lobachev et al. 2004). We suggest that the MRX complex may stabilize the intermediate in which the ITS and telomeric sequences are paired, facilitating the subsequent steps in SSA. Since the SSA event that generates class 2 events involves a very small (≤ 120 bp) heteroduplex relative to the gene-sized regions of heteroduplex analyzed in most studies (Figure 4), the MRX-mediated stabilization of the intermediate may be important. A second relevant observation is that the Mre11p has DNA unwinding activity that is independent of its nuclease activity (Ghodke and Muniyappa 2013). This unwinding activity may facilitate the SSA pathway.

Mutants lacking Tof1p had about a twofold reduction of class 2 events. As described for the class 1 events, since *tof1* mutants reduce stalling caused by proteins bound at ITSs (Anand et al. 2012), there may be a lower level of ITS-associated DSBs, leading to a reduced rate of class 2 events. We also examined a *tof1 mre11* double-mutant strain. 0/41 5-FOA<sup>+</sup> strains had class 2 events in this strain (Table S4). This low rate likely reflects a lower rate of DSB formation (because of *tof1*) coupled with a defect in stabilizing the interaction between the telomere and the ITS (because of *mre11*), as described above.

As described above, loss of Sir2p or Sir3p resulted in a reduction in class 1 events. In contrast, loss of Sir2p elevated the rate of class 2 events threefold; loss of Sir3p had no effect on class 2 events. One difference between these two silencing proteins is that Sir2p, unlike Sir3p, affects silencing in the ribosomal DNA (Gartenberg and Smith 2016). It is likely that the elevation in class 2 events is not ITS-specific. Foss et al. (2017) reported that loss of Sir2p resulted in increased replication gaps in the genome and suggested that the activation of ribosomal DNA replication origins in *sir2* strains results in less efficient replication of other genomic sequences. They found a three- to fivefold elevation in mitotic recombination in non-rDNA sequences, consistent with the threefold elevation of class 2 events in the *sir2*, but not the *sir3*, strain.

Strains lacking Rrm3p, Sgs1p, or Srs2p had rates of class 2 events that were three- to fivefold less than wild-type strains; mutants lacking Mms2p or Rad18p [components of the postreplication repair (PRR) pathway] had no significant effect on the rates of class 1 or class 2 events (Table 1). Sgs1p, Srs2p, and Rrm3p are all helicases, and Sgs1p and Srs2p are antirecombinases (Symington et al. 2014). One interpretation of our observations is that loss of these proteins elevates the frequency of sister-chromatid exchange (Figure 3C), thereby reducing DSB repair in the competing terminal inversion pathway (Figure 3A) responsible for class 2 events. Since the SSA pathway does not involve Rad51p or formation of a Holliday junction, the
antirecombinase activities of Sgs1p and Srs2p would not be likely to affect the rate of terminal inversions directly.

The lowered rate of class 2 events in the *rrm3* strain was quite unexpected. Since Rrm3p promotes replication fork progress through ITSs and other hard-to-replicate sequences (Ivessa et al. 2002), we expected that loss of Rrm3p would lead to elevated DSB formation in the ITS and increased formation of the terminal inversion. However, Muñoz-Galván et al. (2017) showed recently that Rrm3p is required for the repair of broken replication forks. Although most of these studies concerned sister-chromatid recombination, it is possible that SSA is also stimulated by Rrm3p. Thus, loss of Rrm3p would result in elevated cell death (Figure 3D) and a reduced rate of class 2 events (Figure 3A).

Loss of Tel1p elevated terminal inversions about threefold. It is possible that recruitment of telomerase to telomeric repeats results in less efficient SSA. Since Tel1p is involved in recruiting telomerase to the telomeres, loss of Tel1p could allow more efficient SSA. It is unlikely that the short telomeres characteristic of *tel1* strains (Lustig and Petes 1986) contribute directly to elevated rates of class 2 events. The *mre11* and *rad50* strains have equally short telomeres and a reduced level of class 2 events (as described above), and the *rif1* mutation that results in extended telomeres (Hardy et al. 1992) had no significant effect on class 2 events (Table 1). In addition, mutants that lacked Elg1p (involved in stabilizing the replication fork), Top1p (topoisomerase I), and Sml1p (an inhibitor of ribonucleotide reductase) had no substantial effects on either class 1 or 2 events (Table 1).

**Effects of hydroxyurea on the rates of ITS-induced genomic alterations**

ITSs are associated with replication fork stalling and *tof1* mutants reduce fork stalling (Anand et al. 2012); we found that the *tof1* mutation reduced the rates of both class 1 and 2 events. Hydroxyurea (HU) reduces the pools of deoxyribonucleotides, resulting in an accumulation of single-stranded DNA at the replication fork and triggering of a Mec1p-dependent DNA damage checkpoint (Signon 2018). Although the reduction in DNA fork movement caused by HU might be expected to increase the rate of fork stalling, we previously showed that HU reduced fork stalling at ITSs (Anand et al. 2012).
To determine whether HU affected genomic rearrangements, we compared the rates of class 1 and 2 events in the wild-type PG329 strain grown in the presence (100 mM) and absence of HU. The results are shown in Table S5.

HU significantly elevated the rate of class 1 alterations and significantly reduced the rate of class 2 alterations. The elevated rate of class 1 alterations could be explained by an increased efficiency of recruitment of DNA polymerase ζ to the large single-stranded regions at the replication fork in HU-treated cells. Alternatively, the reduced pools of deoxyribonucleotides could result in the increased misincorporation of nucleotides by the replicative DNA polymerases or by DNA polymerase ζ. To determine whether the elevation in point mutations in HU-treated cells was specific to the reporter gene containing the ITS, we examined the rate of class 1 events in a strain (MD737) in which the URA3 reporter gene lacked the ITS. HU resulted in a fourfold elevation in mutation rates, indicating that the effects of HU are independent of the ITS. Northam et al. (2010) previously reported that HU (100 mM) elevated the frequency of CAN1 mutations by about eightfold, and that the increase was largely dependent on Rev3p. Thus, the elevation of class 1 events reflects mutagenic effects that are independent of the ITS.

Although HU was reported to elevate the rate of reciprocal mitotic crossovers ~40-fold in one assay (Barbera and Petes 2006), the rate of class 2 events was reduced about eightfold by HU (Table S5). This reduction is consistent with the fivefold decrease in fork stalling at the ITS in the presence of HU observed by Anand et al. (2012). It is also possible that the lower pools of deoxyribonucleotides in cells treated with HU reduce the efficiency of some steps in SSA. Finally, since the HU-induced DNA damage could result in elevated global cohesion of sister chromatids (Unal et al. 2007), this cohesion would be expected to produce more efficient repair of the DSB by sister-chromatid recombination. As shown in Figure 3, more efficient repair of a DSB by sister-chromatid recombination might then reduce the observed rate of terminal inversions, although other possibilities cannot be excluded.

Discussion

Below, we will summarize the mechanisms by which the ITSs produce class 1 and 2 events, as well as caveats associated with these proposed mechanisms. We will also compare our results with previous studies of ITSs in yeast and mammalian cells.
**Molecular mechanisms that result in class 1 and 2 events**

Although the ITS stimulates both point mutations and terminal inversions, these two types of events are regulated quite differently. The point mutations are dependent on the error-prone DNA polymerase \( \xi \). Since the size of the ITS did not decrease in strains with the point mutation, these mutations are not likely to be a consequence of the mutagenic repair of a DSB. Instead, we suggest that the mutations are introduced during repair of a single-stranded gap (Figure 2B and Figure 5A). Since class 1 events are reduced in strains lacking Mre11p, Rad50p, and Sae2p (which activates the Mre11p nuclease), or containing the nuclease-deficient \( mre11-H12SN \) allele, the proposed gap may be initiated by a nick caused by the Mre11p endonuclease that is expanded into a gap by the 3′–5′ nuclease of Mre11p, and the 5′–3′ activities of Exo1p or Sgs1p/Top3p/Rmi1p/Dna2p (Symington et al. 2014). We found that mutations in neither \( \text{EXO1} \) nor \( \text{SGS1} \) reduced the rate of class 1 events (Table 1), possibly due to the redundancy of the enzymes involved in resection.

Unlike the class 1 events, the class 2 events are likely initiated by a DSB. In a previous study with an ITS that was about twice as long as that used in our study, Goto et al. (2015) found an ITS-associated DSB. In contrast to class 1 events, class 2 events occurred independently of the nuclease activity of Mre11p. As with most other spontaneous mitotic recombination events, the enzymes that initiate DSB formation at the ITS are not known. In addition, although the partial dependence of class 2 events on Tof1p argues that about half of the DSBs may be associated with stalled replication forks, about half of these events may be independent of DNA replication. Previously, we showed that more than half of spontaneous mitotic crossovers in yeast are associated with DSB formation in unreplicated chromosomes (Lee et al. 2009; St. Charles and Petes 2013). In addition, Zhang et al. (2012) showed that tracts of GAA/TTC repeats simulated DSBs in both S-phase and in nondividing cells.

Following the initiating DNA lesion, the subsequent steps in the generation of class 2 events have a leaky Ura− phenotype and grow weakly on medium containing 5-FOA. However, strains with only eight copies of the telomere repeat [\( \text{URA3::Int (ACACACCC)}_8 \)] have a leaky Ura− phenotype and grow weakly on medium containing 5-FOA. Fast-growing 5-FOA− derivatives have expansions of the telomeric tract. These expansions are dependent on two recombination pathways that both require Rad56p: a gap-repair pathway requiring the HR proteins Rad51p and Rad52p, and a PRR pathway utilizing Rad5p. We previously suggested that telomeric proteins bound to the ITS result in fork stalling and a Tof1p-mediated pause. This pause then facilitates template switching (PRR pathway) or gap filling (HR pathway). Since class 1 and class 2 events in our study are independent of Rad51p and the PRR proteins Mms2p and Rad18p, the mechanisms that generate class 1 and 2 events are clearly quite different from those that produce expansions of the ITS.

**Caveats about the proposed mechanisms of class 1 and 2 events**

Although the Rev3p-dependence of class 1 events and the reliance of class 2 events on the SSA pathway are clear, there are a number of mutations that had effects with multiple possible interpretations. For example, we suggested that the elevation of class 2 events in strains lacking Sir2p might be an indirect effect of the replication stress imposed by the unsilencing of replication origins in the ribosomal DNA, leading to elevated DSB formation. However, since Sir2p is a histone deacetylase with multiple potential targets, other possibilities also exist. Aksenova et al. (2015) noted that the sir2 mutation resulted in a 26-fold elevation in tract expansions of the ITS.

In our study of ITS-induced alterations, most of the mutations had relatively small effects, affecting the rate of the events < 10-fold. In contrast, in studies of recombination events induced by site-specific meganucleases such as HO or I-SceI, mutations often have larger quantitative effects. For example, loss of Rad52p reduces the efficiency of HO-induced recombination by 100-fold (Malkova et al. 1996) but has only a 10-fold effect on class 2 events. One difference between spontaneous events and those induced by meganucleases is that meganucleases produce a single type of DNA lesion, whereas spontaneous events likely reflect multiple types of DNA lesions. Thus, there are likely to be a large number of pathways that are relevant to the repair of spontaneous DNA damage. Some of these pathways may be redundant such that eliminating any single pathway has a relatively small effect.

**Genetic regulation of ITS-induced alterations in other studies**

In the current study, the ITS is oriented such that the G-rich telomeric repeats are on the Watson strand, as defined in the \( \text{Saccharomyces} \) Genome Database (Figure 1). In another study, we analyzed genomic alterations in strains with the reverse orientation of the ITS (Aksenova et al. 2015). In this orientation, strains with the \( \text{URA3::Int (ACACACCC)}_{15} \) allele are 5-FOA−. However, strains with only eight copies of the telomere repeat [\( \text{URA3::Int (ACACACCC)}_8 \)] have a leaky Ura− phenotype and grow weakly on medium containing 5-FOA. Fast-growing 5-FOA− derivatives have expansions of the telomeric tract. These expansions are dependent on two recombination pathways that both require Rad56p: a gap-repair pathway requiring the HR proteins Rad51p and Rad52p, and a PRR pathway utilizing Rad5p. We previously suggested that telomeric proteins bound to the ITS result in fork stalling and a Tof1p-mediated pause. This pause then facilitates template switching (PRR pathway) or gap filling (HR pathway). Since class 1 and class 2 events in our study are independent of Rad51p and the PRR proteins Mms2p and Rad18p, the mechanisms that generate class 1 and 2 events are clearly quite different from those that produce expansions of the ITS.

As discussed in the Introduction, ITSs in mammalian cells are often associated with fragile sites and with breakpoints of translocations observed in tumor cells. Although these observations suggest that ITSs stimulate DNA breaks, the alternative possibility is that ITSs get inserted at regions that are prone to breakage for other reasons. Our results, as well as several previous yeast studies (Aksenova et al. 2013, 2015; Goto et al. 2015), show that ITSs in yeast are themselves prone to generating chromosome rearrangements. Although the genetic regulation of ITS-caused instability in mammalian systems has not been examined in detail, Bosco and de
Lange (2012) showed that deletion of the telomere-binding TRF1 protein elevated the fragility of an ITS in human cells.

**Conclusions**

In conclusion, our data strongly argue several points. First, the ITS-induced point mutations are initiated by a mechanism that is different from point mutations initiated by the trinucleotide tract GAA/CTT. Although the mutagenic effects of both types of sequence require Rev3p, the GAA/CTT tracts induce mutations by repair of a DSB, whereas the ITS induces mutations independently of a DSB, likely utilizing a Mre11p-generated single-strand gap. Second, class 2 events occur through the SSA pathway by a mechanism that requires the MRX complex but does not require the Mre11p-encoded nuclease.

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Interstitial Telomeric Sequences

437
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