Recovery of Arrested Replication Forks by Homologous Recombination Is Error-Prone

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Abstract
Homologous recombination is a universal mechanism that allows repair of DNA and provides support for DNA replication. Homologous recombination is therefore a major pathway that suppresses non-homology-mediated genome instability. Here, we report that recovery of impeded replication forks by homologous recombination is error-prone. Using a fork-arrest-based assay in fission yeast, we demonstrate that a single collapsed fork can cause mutations and large-scale genomic changes, including deletions and translocations. Fork-arrest-induced gross chromosomal rearrangements are mediated by inappropriate ectopic recombination events at the site of collapsed forks. Inverted repeats near the site of fork collapse stimulate large-scale genomic changes up to 1,500 times over spontaneous events. We also show that the high accuracy of DNA replication during S-phase is impaired by impediments to fork progression, since fork-arrest-induced mutation is due to erroneous DNA synthesis during recovery of replication forks. The mutations caused are small insertions/duplications between short tandem repeats (micro-homology) indicative of replication slippage. Our data establish that collapsed forks, but not stalled forks, recovered by homologous recombination are prone to replication slippage. The inaccuracy of DNA synthesis does not rely on PCNA ubiquitination or trans-lesion-synthesis DNA polymerases, and it is not counteracted by mismatch repair. We propose that deletions/insertions, mediated by micro-homology, leading to copy number variations during replication stress may arise by progression of error-prone replication forks restarted by homologous recombination.

Introduction
Maintenance of genome stability requires the faithful and accurate replication of the genetic material. Genome instability is a hallmark for most types of cancer and it is strongly associated with predisposition to cancer in many human syndromes (for a review, see [1,2]). Genome instability is manifest at two levels: at the nucleotide level, resulting in base-substitutions, frame-shifts or insertions in deletions (referred to herein as mutations); and at the chromosomal level, resulting in duplications, deletions, inversions and translocations (referred to herein as gross chromosomal rearrangements or GCRs).

Genome instability during cancer development and in other human genomic disorders may be consequences of failures in chromosome replication (for a review, see [3,4]). Origin spacing has recently been shown to cause chromosomal fragility at some human fragile sites [5,6]. Impediments to replication fork movements per se may also cause genome instability [7–9]. Indeed, both slowing down and blockages to fork progression can lead to chromosomal fragilities or GCRs in human cells and yeast models [10–14]. However, how a blocked replication fork leads to genetic instability remains poorly understood.

In eukaryotes, DNA replication is initiated at numerous origins along linear chromosomes, and impediments to fork progression appear unavoidable during each S-phase (for a review, see [9,15]). Impediments to fork progression can be caused by DNA lesions, by non-histone proteins tightly bound to DNA, by sequence-caused secondary structures such as cruciform structures and possibly G-quadruplexes, by nucleotide pool imbalance and by conflicts with transcription machinery (for a review, see [16,17]). In case of failures in fork progression, DNA replication can be completed either by the recovery of the arrested fork by fork-restart mechanisms, or as a result of the progression of a converging fork which can be ensured by activation of dormant origins [7,15,18]. Fork restart is presumably essential in unidirectional replication regions, such as the rDNA locus, in regions of low densities of origins, such as some human fragile sites, and when two converging forks are both impeded [5,19,20].

To ensure faithful and complete DNA replication, cells coordinate DNA synthesis restart with specific pathways, including DNA replication checkpoint and homologous recombination mechanisms [17]. The integrity of replication forks is guaranteed by the DNA replication checkpoint that maintains the replisome in a replication-competent state to keep DNA polymerases at the site...
Author Summary

The appropriate transmission of genetic material during successive cell divisions requires the accurate duplication and segregation of parental DNA. The semi-conservative replication of chromosomes during S-phase is highly accurate and prevents accumulation of deleterious mutations. However, during each round of duplication, there are many impediments to the replication fork machinery that may hinder faithful chromosome duplication. Homologous recombination is a universal mechanism involved in the rescue of replication forks by rebuilding a replication apparatus at the fork (by mechanisms that are not yet understood). However, recombination can jeopardize genome stability because it allows genetic exchanges between homologous repeated sequences dispersed through the genome. In this study, we employ a fission yeast-based arrest of a single replication fork to investigate the consequences of replication fork arrest for genome stability. We report that a single blocked fork favours genomic deletions, translocations, and mutations; and this instability occurs during fork recovery by recombination. We also report that a single arrested fork that resumes its progression by recombination is prone to causing replication slippage mediated by micro-homology. We propose that deletions/duplications observed in human cancer cells suffering from replication stress can be viewed as scars left by error-prone replication forks restarted by recombination.

of nucleotide incorporation [21]. It remains unclear how the DNA replication checkpoint modulates replisome activities to maintain its function [21,22]. The DNA replication checkpoint also regulates nuclease activities (e.g. Exo1 or Mus81) which contribute to preserving the integrity of stalled forks [23,24]. If replisome function is lost or the replisome dissociates at broken replication forks, the resumption of DNA synthesis appears to require the replisome to be rebuilt. In E. coli, restart of a collapsed fork involves homologous recombination and the PriA helicase that allows replisome components to be loaded de novo on joint-molecule structures [25,26]. In eukaryotes, the restart of collapsed or broken replication forks is dependent upon homologous recombination, but the mechanism of origin-independent loading of the replisome remains to be described [20,27–30]. It has been proposed that the repair of a double-strand break (DSB) by recombination (break-induced replication, BIR) in budding yeast similarly involves the assembly of a replication fork (for a review, see [30–32]). When BIR occurs outside S-phase, recombination-dependent replication fork assembly can synthesise hundreds of kilobases (Kb). However, this DNA synthesis is highly inaccurate due to frequent template switching of nascent-strands and frame-shift mutations [33,34].

We previously reported a system that displays replication fork arrest at a specific locus in the fission yeast S. pombe. The system is a polar replication fork barrier (RFB) regulated by the Rtf1 protein binding to its RTS1 binding site [35]. The RTS1-RFB causes fork arrest because of a non-histone protein complex binding to the DNA. As proposed for other polar RFBs, the RTS1-RFB is thought to block fork progression by directly (contact between proteins and the replisome) or indirectly (topological constraint) affecting the replicative helicase activity and thereby preventing DNA unwinding [36,37]. Recovery of the arrested fork occurs by a DSB-independent mechanism and involves the recruitment of recombination proteins at the RTS1-RFB site. We proposed that recombination proteins associate with unwound nascent strands that then anneal with the initial template to allow DNA synthesis to restart [11,20]. The causative protein barrier then has to be removed either by DNA helicase or by the recombination machinery itself to allow fork-progression to resume [38–40]. Occasionally, the unwound nascent strand can mistakenly anneal with a homologous template in the vicinity of the collapsed fork, resulting in the restart of DNA synthesis on non-contiguous template. This incorrect template switch of nascent strands results in inversions and iso-aentric and dicentric chromosomes in ~2–3% of cells/generation [11,20]. Error-free template switching between sister-chromatids provides an efficient mechanism for filling-in single-stranded gaps left behind damage-induced stalled forks [41]. Inverted chromosome fusions in yeast and rare-genome rearrangements in human genomic disorders, may both be consequences of template switching between ectopic repeats associated with impeded replication forks [8,14].

Here, we used the RTS1-RFB to investigate the consequences of fork collapse on genome instability. We report that recovery from a collapsed fork is associated with a high frequency of instability, with a single fork arrest increasing the rates of mutation, deletion and translocation by 10, 40 and 5 fold, respectively. We show that genetic instability associated with fork arrest is dependent on homologous recombination. Fork-arrest-induced GCRs (deletion and translocation) result from inappropriate ectopic recombination at the site of the collapsed fork. We also demonstrate that restoration of fork progression by homologous recombination results in error-prone DNA synthesis due to frequent replication slippage between short tandem repeats. We investigated the molecular mechanisms of this replication slippage and found that post-replication repair, including ubiquitination of PCNA or translesion-synthesis (TLS) DNA polymerases, is not involved in fork-arrest-induced replication slippage. Micro-deletions/insertions flanked by micro-homology associated with copy number variations (CNVs) in cancer cells or in response to replication stress may therefore be scars left following the restoration of forks progression by homologous recombination.

Results

The conditional replication fork barrier RTS1

We generated fork arrest constructs by manipulating the polar RTS1-RFB (Figure 1A). We introduced the RTS1 sequence on the centromere-proximal (eco-proximal) side of the ure4 locus, 5 kb away from the strong replication origin (ori) 3006/7 on chromosome III. This created the t-ure4<on locus, in which “t-” and “on” refer to the telomere and the origin 3006/7, respectively; and “<” and “>” refer to the RTS1-barrier and its polarity that is whether it blocks replication forks travelling from the ori 3006/7 towards the telomere or forks travelling from the telomere towards the ori 3006/7, respectively. We previously confirmed that forks moving from ori 3006/7 towards the telomere (tel) are efficiently blocked by the RTS1-RFB at the t-ure4<con locus [35]. In this model system, fork arrest is activated by inducing the expression of rif7” gene that is under control of the thiamine repressible promoter nmt41. Thus, the RTS1-RFB is inactivated by adding thiamine to the media and it is activated in thiamine-free media. Efficient induction of Rif1 expression requires incubation for 12–16 hours in thiamine-free media. Replication intermediates were analysed by native 2-dimensional gel electrophoresis (2DGE). In conditions of Rif1 expression, more than 95% of replication forks were blocked by the RTS1-RFB at the t-ure4<con locus [see black arrow on Figure 1B, t-ure4<con ON]. Arrested forks were not detected without Rif1 induction [Figure 1B, t-ure4<con OFF] [20].
Conditional fork-arrest constructs to investigate fork- 
arrest-induced genome instability

The RTS1 sequence was inserted in the tel-proximal side of the ora4 locus. 2DGE analysis of this construct revealed a strong fork arrest signal on the descending large Y arc (Figure 1A and 1B, <ora4-ori ON>). The ora4 sequence, used in this system as a reporter to score genetic instability, is located behind the arrested fork when the RTS1-RFB is active at the <ora4-ori locus and ahead of the arrested fork at the t-ura4-ori locus. This explains the distinct position of the arrested fork signal on the Y arc.

Introducing a second RTS1 sequence, such that the two RTS1 sequences are inverted repeats (IRs), created t-ura4<ori and t<ura4<ori loci (Figure 1A and 1B, t<ura4<ori ON). Given the orientation of the two RTS1-RFB in the t<ura4<ori strain, converging forks cannot be blocked.

A single fork arrest induces genomic deletions

We investigated fork-arrest-induced genome instability by selecting for cell resistance to 5-FOA, the result of loss of ora4 function. Inducing fork-arrest at t-ura4<ori increased ora4 loss 3-fold (Table 1). Rtf1 expression in the t-ura4-ori and t<ura4-ori strains did not cause site-specific fork-arrest at ora4 as assessed by 2DGE and did not increase the rate of ora4 loss. Thus, ora4 loss results from the RTS1-RFB activity and not simply from the presence of RTS1 and/or Rtf1 expression (Table 1). To investigate the nature of this genetic instability, primers were designed to amplify the ora4 coding sequence and, as a control, the essential mg3 gene, mapping 30 kb tel-proximal to ora4, that should not be rearranged (Figure 2A and 2B) [35]. The absence of ora4 amplification was classified as a deletion event; sequencing of amplified ora4 sequence was used to identify point mutation events (Figure 2B).

A single arrested fork at the t-ura4<ori locus was sufficient to increase the rate of genomic deletion up to 40 times over spontaneous events (i.e. in the t-ura4-ori strain, p = 0.006) (Figure 2C and Figure S1A). Fork-arrest-induced deletion was recombination-dependent. Spontaneously (i.e. when the RTS1-barrier was inactivated), the rate of genomic deletion in rad22-d or rhp51-d strains was higher than that in the wild-type strain (Figure S1B).

A single fork arrest is not restarted by homologous recombination, the RTS1-RFB behaves as a hot spot for replication termination [20].
indicating that fork-arrest-induced deletion was also mediated by inter-chromosomal recombination (Figure 2C and t-ura4,ori RTS1-d on Figure S1A). Thus, these data are consistent with the view that homologous recombination makes a major contribution to suppressing genome instability, but can occasionally drive non-allelic recombination events leading to GCRs [35,45].

We detected no fork-arrest-induced deletion in the t-ura4,ori strain, in contrast to the t-ura4,ori RTS1-d on Figure S1A and Figure 2C. The ura4 marker is located behind and ahead of collapsed forks in the t-ura4,ori and t-ura4<ori strains, respectively (Figure 1A). Therefore, replicated regions, located behind collapsed forks, do not display instability, and fork-arrest-induced deletion occurs within unreplicated regions immediately in front of arrested forks. Overall, our data establish that genomic deletion at collapsed forks results from inappropriate recombination between ectopic sequences during the process of fork recovery by recombination proteins.

Inverted repeats stimulate fork-arrest-induced deletion by promoting inter- and intra-chromosomal recombination

Inverted repeats (IRs) are structural elements often associated with genome rearrangements [11,14,46,47]. We investigated the
effects of IRs in the vicinity of the RTS1-RFB on fork-arrest-induced genomic deletion. We first compared the \( \geq u44 \) strain (IRs flanking \( u44 \)) to the \( \leq u44 \) strain (no IRs near the RTS1-RFB). The rate of fork-arrest-induced genomic deletion was 200 times higher in the \( \geq u44 \) than that in the \( \leq u44 \) strain (\( p = 0.009 \), Figure 2C and Figure S1A). Thus, intrachromosomal ectopic recombination permitted by the RTS1 sequence on the tel-proximal side of \( u44 \) accounted for 99.5% of the genomic deletions observed in the \( \geq u44 \) strain (Figure 2C, compare with \( \leq u44 \)). Preventing interchromosomal recombination by deleting RTS1 from the chromosome II (\( \geq u44 \) \( \leq u44 \) RTS1-d) abolished 90% of deletion events (Figure 2C and Figure S1A). Thus, genomic deletions induced by fork-arrest near IRs are due to inter- and intrachromosomal recombination events. In support of this, stimulation of fork-arrest-induced deletion by IRs is mediated by homologous recombination. Indeed, the rate of genomic deletion was not increased upon induction of the RTS1-RFB in the surviving population of \( \geq u44 \) \( \leq u44 \) rad22-d and rhp51-d strains (Figure S1B). These data indicate that IRs favour genomic deletion at collapsed forks by promoting inappropriate inter- and intrachromosomal recombination during fork recovery by recombination proteins.

We verified that our data were not influenced by the orientation of IRs or by rare blocking of converging forks in the \( \geq u44 \) strain. We analysed the \( \geq u44 \) construct in which RTS1 repeats are in the opposite orientations relative to the \( \geq u44 \) construct, such that forks converging towards \( u44 \) cannot be blocked (Figure 1). The rate of fork-arrest-induced genomic deletion was 1,000 times higher in the \( \geq u44 \) than that in the \( \leq u44 \) strain, that does not contain IRs near the RTS1-RFB (\( p = 0.008 \), Figure 2C and Figure S1A). Thus, interchromosomal recombination, permitted by the RTS1-RFB sequence on the centrominal side of \( u44 \), accounted for nearly 100% of the genomic deletions observed in the \( \geq u44 \) strain (Figure 2C, compare with \( \leq u44 \)). Preventing interchromosomal recombination by deleting RTS1 from the chromosome II \( \geq u44 \) \( \leq u44 \) RTS1-d abolished 90% of deletion events (Figure 2C and Figure S1A). Importantly, the deletion rates for the \( \geq u44 \) and \( \leq u44 \) strains were not significantly different (Figure 2C), showing that IRs cause genomic deletion at collapsed forks irrespective of their orientations and independently of blockage of converging forks.

A single collapsed fork induces translocations that are stimulated by IRs

Fork-arrest at \( \geq u44 \) results in translocations between ectopic RTS1 repeats on chromosomes II and III [35]. We investigated the influence of IRs on fork-arrest-induced translocation. We designed primers to amplify the predicted translocation junction between chromosomes II and III (TLII and TLIII on Figure 2A and 2B). A single arrested fork at the \( \leq u44 \) locus was sufficient to increase the translocation rate to 5 times higher than the spontaneous rate (\( p = 0.002 \), Figure 2D and Figure S1C). The translocation rate for the \( \geq u44 \) construct (containing IRs) was 1,500 fold higher than that for the \( \leq u44 \) strain that does not contain IRs near the RTS1-RFB (\( p = 0.009 \), Figure 2D and Figure S1C). Thus, intra-chromosomal recombination accounted for nearly 99% of translocation events observed in the \( \geq u44 \) construct (Figure 2D and Figure S1C, compare with \( \leq u44 \)). No translocation events were detected when interchromosomal recombination was prevented by deleting RTS1 from the chromosome II (\( \geq u44 \) \( \leq u44 \) RTS1-d on Figure 2B). Therefore, as reported for genomic deletions, fork-arrest-induced translocation associated with IRs is due to inter- and intrachromosomal ectopic recombination. No translocations were detected in the \( \leq u44 \) construct (data not shown), so we cannot formally exclude the possibility that fork-arrest-induced translocations in the \( \leq u44 \) strain was caused by blocking of converging forks. However, as no translocation event occurred in the absence of Rad22\( ^{R S} \) or Rhp51\( ^{R S} \), it is most likely that translocations occur during fork recovery by recombination (Figure 2B and [35]). Overall, our data indicate that recovery of a single collapsed fork causes translocations and IRs near the site of fork-arrest stimulate translocations by promoting inappropriate inter- and intrachromosomal recombination.

Table 1. Rates of \( u44 \) loss (including genomic deletion, translocation, and mutation events), calculated using the method of the median.

| strains* | Rate of \( u44 \) lossb | Fold induction by Rtf1 expression (−/+ thiamine)c | Fold induction by the RFB (relative to the \( t-u44\)-ori construct) |
|----------|-------------------------|-----------------------------------------------|-----------------------------------------------|
| \( t-u44\)-ori | 4.6±1 4.3±1.8 0.9 | 3.4 (p = 0.002) 2.6 |
| \( t-u44\)-ori | 4.1±0.9 13.9±2.8 | 0.6 (p = 0.03) 0.1 |
| \( t-u44\)-ori RTS1-d | 4.9±1.4 11.4±4.1 | 0.6 (p = 0.004) 0.1 |
| \( t-u44\)-ori | 3.9±1.3 4.8±2.5 | 1.2 | 1.1 |
| \( t-u44\)-ori | 5.8±2.1 97.4±51.5 | 16.8 (p = 0.004) 22.6 |
| \( t-u44\)<ori RTS1-d | 4.9±2.2 19.5±3.3 | 4 (p = 0.006) 4.5 |
| \( t-u44\)-ori | 3.2±1.7 3.6±1.8 | 1.1 0.8 |
| \( t-u44\)-ori | 6.5±1.7 50.3±14.2 | 7.6 (p = 0.014) 11.7 |

* the following nomenclature is used: “t” and “ori” refer to the telomere and the replication origin 3006/7, respectively; “<” and “>” indicate the RTS1-RFB and its polarity (< blocks replication forks moving from the ori3006/7 towards the telomere, and > blocks replication forks moving from the telomere towards the origin 3006/7).

b event/cell/division \( \times 10^{-8} \) ± standard error. Values are means of at least 3 independent rates.

c statistical significance was determined using the nonparametric Mann-Whitney U test.

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Fork-recovery by Recombination is Inaccurate

Fork-arrest-induced GCRs are caused by inter- and intra-chromosomal recombination. We noticed a slightly greater contribution of intra- than inter-chromosomal recombination (Figure 2C). This is consistent with ectopic recombination preferentially occurring at the most proximal homologous sequence, as previously reported [49]. Nonetheless, the rate of fork-arrest-induced deletion in the $t$-ura4 strain ($8.4 \times 10^{-7}$) was not simply the sum of the rates of intra-chromosomal recombination events ($9.9 \times 10^{-8}$ in the $t$-ura4 strain) and inter-chromosomal recombination events ($4 \times 10^{-9}$ in the t-ura4 strain). Similar reasoning can be applied for the $t$-ura4 strain. Thus, independent intra- and inter-recombination events cannot themselves explain high rate of GCRs induced by fork arrest near IRs. Therefore, we infer that there is interplay between inter- and intra-chromosomal recombination such that fork-arrest-induced GCRs may involve recombination between three homologous sequences (tri-parental recombination).

The RecQ helicase Rqh1 prevents GCRs at collapsed forks

To confirm that fork-arrest-induced GCRs are the result of inappropriate ectopic recombination during fork recovery, we analysed the involvement of the RecQ helicase Rqh1. We previously reported that Rqh1 limits inappropriate template switching of stalled nascent strands without affecting the efficiency of fork restart [20]. In the $t$-ura4 strain (in which only inter-chromosomal recombination is possible), fork-arrest-induced deletion and translocation rates were 31 and 109 times higher in the rqh1-d strain than that in the wild-type control, respectively ($p = 0.0003$, Figure 2D–2E and Figure S1C). For the $t$-ura4 strain construct (containing IRs near fork-arrest), fork-arrest-induced deletion and translocation rates were 5 times higher in the rqh1-d strain than that in the wild-type control ($p = 0.0007$, Figure 2D–2E and Figure S1C). Thus, Rqh1 limits GCRs at collapsed forks by preventing inappropriate ectopic recombination during the process of fork recovery by recombination proteins.

A single fork arrest induces mutations

We analysed the effects of collapsed forks on the mutation rate. We sequenced the ura4 coding sequence from 5-FOA+ isolated cells and identified base-substitutions, frame-shifts and small insertions and duplications between short tandem repeats (Table 2). A single collapsed fork in the $t$-ura4 strain increased the overall mutation rate up to 10 times over spontaneous events (Figure 3A, $p = 0.0003$). Similar increases in the overall mutation rate were found for the strains with IRs near the arrested fork and those with RTS1 deleted from chromosome II (Figure 3A and Figure S2A). Thus, fork-arrest-induced mutation is not mediated by inappropriate ectopic recombination. Induction of the RTS1-RFB in the $t$-ura4 strain did not increase the mutation rate of the ura4 gene. Thus, as for GCRs, replicated regions behind arrested forks are not prone to mutation. This observation rules out the hypothesis that fork-arrest-induced mutation is a consequence of the accumulation of damaged single-stranded DNA behind collapsed forks (see discussion). Our data suggest that recovery from collapsed forks results in error-prone DNA-synthesis.

Collapsed forks result specifically in replication slippage

We then analysed the spectra of mutations found in the ura4 ORF by sequencing the PCR products. The rates of base-substitutions and frame-shifts were not significantly increased by the RTS1-RFB activity over spontaneous events (i.e. compare to t-ura4 strain, Figure 3C and Table 2). In contrast, the rate of deletions and duplications (Del/Dup) flanked by short homology was increased by 7 times over spontaneous events in the t-ura4 strain, but not in the t-ura4 strain (Figure 3C and Table 1). These data further confirm that fork-arrest does not promote mutation events behind collapsed forks.

We used reversed mutation assays to test if fork-arrest at the RTS1-RFB specifically induced Del/Dup mutations. We made use of strains harbouring a single mutation within the ura4 ORF: either a single base-substitution or a $-1$ frame-shift in homonucleotide (Figure S2B). We also studied strains harbouring either a duplication of 20 or 22 nt flanked by 5 or 4 bp of micro-homology, respectively (defined as ura4-dup20 and ura4-dup22, Figure S2B). These non-functional ura4 alleles were inserted in front of the RTS1-RFB in the t-ura4 strain configuration and we then tested whether fork arrest could restore a functional ura4 gene. Activation of the RTS1-RFB at ura4 increased the frequency of Ura+ revertants up to 15 and 7 times in strains harbouring ura4-dup20 and ura4-dup22, respectively (Figure 3D and Figure S2B). Thirty Ura+ colonies were studied by PCR and all gave a product of the same size as the wild-type ura4 gene: they had therefore lost the duplication (Figure 3E and data not shown). Sequencing the full ura4 ORF confirmed that Ura+ revertants contained an intact ura4 sequence, showing that the reversion of these alleles was due solely to the precise deletion of 20 or 22 nt (Figure 3F and data not shown). In contrast, activation of the RTS1-RFB did not increase the frequency of Ura+ revertants of strains harbouring ura4 alleles with a single base-substitution or a $-1$ frame-shift (Figure 3D and Figure S2B). Thus, collapsed forks tend to induce deletion events between short tandem repeats rather than base-substitution or frame-shift mutations. Among Del/Dup events, deletions represented the two-third of events in the t-ura4 strain (Table 2). The median size of Del/Dup events was 24 and 22 nt respectively, and Del/Dup occurred between short direct repeats 1 to 10 nt long (Figure S3). Thus, the ura4-dup20 and ura4-dup22 alleles used in the reverse mutation assay were representative of the Del/Dup events observed. Del/Dup flanked by micro-homology result from intra-molecular template switching mechanisms in which nascent strands dissociate from the template and misalign with the template when restarting the elongation step. This leads to loop formation, either in the nascent strand or in the template, resulting in duplication or deletion events, respectively [49]. Consequently, we will hereafter refer to Del/Dup as replication slippage. Replication slippage was observed all along the ura4 ORF and up to 1.2 kb ahead of the arrested fork, even if a hot spot of deletion was present 500 bp away from the RTS1-RFB (Figure 3G and Figure S3B). Thus, our data suggest that the DNA synthesis is prone to replication slippage at least for the first 1,200 nt synthesized during the recovery of collapsed forks. Inaccuracy of DNA synthesis on further distances was not directly addressed.

Replication slippage results from error-prone DNA synthesis during fork recovery

To confirm that replication slippage occurs as forks recover, and not behind the fork in the DNA already replicated, we inserted the ura4-dup20 or the ura4-dup22 allele either behind (in the t-ura4 configuration) or in front of the RTS1-RFB (in the t-ura4 configuration) (Figure 4). This allows the analysis of the same event of replication slippage behind and ahead of collapsed forks. In the t-ura4 configuration, induction of the RTS1-RFB resulted in a 8 and 16 fold increases in the replication slippage frequency for the ura4-dup20 and ura4-dup22 alleles, respectively (Figure 4A and 4B). Similar increases in the rate of replication slippages were observed (Figure 4C). In contrast, in the t-ura4 background, the frequency of replication slippage was induced by only 2–3 fold
Figure 3. Fork-arrest induces mutations. A. Effect of intra- and inter-chromosomal recombination between RTS1 repeats on fork-arrest-induced mutation rates (base-substitutions, frame-shifts and small insertions or deletions between short tandem repeats). RTS1-RFB activity and ura4 location with respect to the RFB are given for each construct. The % of mutation events, as determined by the PCR assay and sequencing, was used to balance the rate of ura4 loss. Then, the RFB-induced mutation rate was calculated by subtracting the rate obtained in the presence of thiamine (Rtf1 being repressed) from the rate obtained in the absence of thiamine (Rtf1 being expressed). The values reported are means of at least 3 independent median rates. Error bars correspond to SE. (Refer to Figure S2 for corresponding rates of mutation when Rtf1 is expressed or not). B. Rate of mutation for indicated strains; ON and OFF refers to the RTS1-RFB being active or not, respectively. The % of mutation events, as determined by the PCR assay and sequencing, was used to balance the rate of ura4 loss. The values reported are means of at least 3 independent median rates. Error bars correspond to SE. Statistically significant fold differences in mutation rates between the "OFF" and "ON" conditions are indicated with an *. C. Spectra of mutation events in indicated strains upon RFB induction (refer to Table 2 for exact numbers and to Figure S3 for mapping of deletions/duplications and their features). D. Strains harbouring the ura4 alleles with a single base-substitution or frame-shift or duplication of 20 or 22 nt, together with the RTS1-RFB in the t-ura4-dup20-ori configuration were streaked onto the indicated media after cell growth with (RFB "OFF") or without (RFB "ON") thiamine. The bottom diagram indicates strain positions and the mutation events required to obtain Ura+ revertants. PCR analysis of Ura+ revertants isolated from the t-ura4-dup20-ori strain (duplication of 20 nt in ura4) after RFB induction. With the primers used, a 106 bp fragment is amplified from the ura4+ strain and a 126 bp fragment is amplified from the t-ura4-dup20-ori strain. Sequence alignments of ura4-dup22, ura4-dup20, ura4+ alleles and corresponding Ura+ revertants. Micro-homologies are indicated in grey and duplicated sequences are underlined in black. The phenotype of each allele is indicated on the figure. G. Map of deletion and duplication events within the ura4 ORF.

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by the RTS1-RFB (Figure 4B–4C). These data confirm that DNA located ahead of collapsed forks is more prone to replication slippage than replicated DNA adjacent to arrested forks, further evidence that replication slippage arises during fork recovery.

Replication slippage results from forks restarted by recombination

Replication slippage occurs in DNA in front of (and not behind) the arrested fork, this DNA being replicated only after restart of the fork. Thus, a defect preventing fork recovery would be expected to abolish the error-prone DNA synthesis during restart. We analyzed fork-arrest-induced mutation in recombination mutants in which collapsed forks at the RTS1-RFB cannot recover, resulting in cell death. Induction of the RTS1-RFB did not increase the overall mutation rate in the surviving populations of \( t \geq \text{ura4} < \text{ori} \) or \( t \leq \text{ura4} < \text{ori} \) rad22-d and rhp51-d strains (Figure 3B). In addition, only 7% of mutation events in the survivors of the rad22-d \( t \geq \text{ura4} < \text{ori} \) strain were Del/Dup mutations, compared to 40% in the wild-type strain (Figure 3C and Table 1). We currently cannot assess mutation events associated with defects in fork recovery because this appears to be lethal in the absence of recombination. Nevertheless, our data are consistent with fork-arrest-induced replication slippage being dependent on homologous recombination. The rad22-d and rhp51-d strains are themselves spontaneously mutagenic. Consequently, any small increase in the fork-arrest-induced mutation rate might be masked by the high frequency of spontaneous 5-FOA\(^6\) cells in rad22-d and rhp51-d strains. We therefore used a more specific mutation assay, based on the \( \text{ura4} \)-dup20 allele, to determine the rate of replication slippage induced by the RTS1-RFB over spontaneous events.

Strains carrying mutations in recombination genes grow slowly, so replication slippage was scored as a function of the number of generations following thiamine removal (i.e. generations subject to fork arrest at \( \text{ura} \)) (Figure 4D and 4E). In the wild-type strain, fork arrest at the RTS1-RFB resulted in a 10 fold-increase in the frequency of replication slippage, as expected. In recombination mutants (\( \text{rad50-d} \), \( \text{rhp51-d} \) and \( \text{rad22-d} \)), fork-arrest at the RTS1-RFB increased the frequency of replication slippage by only 2 times over spontaneous events: therefore, replication slippage occurs less frequently in survivors from recombination mutants than those from the wild-type strain (Figure 4D–4F). Based on 2DGE analysis, fork-restoration is severely impaired in the absence of

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**Table 2. Mutations spectra in the indicated strains.**

|                     | \( t \geq \text{ura4} < \text{ori} \) | \( t \leq \text{ura4} < \text{ori} \) | \( t \leq \text{ura4} < \text{ori} \) | \( t \leq \text{ura4} < \text{ori} \) | \( t \leq \text{ura4} < \text{ori} \) | \( t \geq \text{ura4} < \text{ori} \) rad22-d |
|---------------------|--------------------------------------|--------------------------------------|--------------------------------------|--------------------------------------|--------------------------------------|--------------------------------------|
| **Rtf1 expression** | -                                    | +                                    | -                                    | +                                    | -                                    | +                                    |
| **Fork arrest at \( \text{ura4} \)** | -                                    | -                                    | -                                    | +                                    | -                                    | +                                    |
| **Transitions**     |                                      |                                      |                                      |                                      |                                      |                                      |
| \( \text{GC} \rightarrow \text{AT} \) | 5                                    | 1                                    | 7                                    | 14                                   | 6                                    | 2                                    |
| \( \text{AT} \rightarrow \text{GC} \) | 3                                    | 2                                    | 4                                    | 9                                    | 1                                    | 4                                    |
| **Transversions**   |                                      |                                      |                                      |                                      |                                      |                                      |
| \( \text{GC} \rightarrow \text{TA} \) | 3                                    | 1                                    | 2                                    | 1                                    | 0                                    | 5                                    |
| \( \text{GC} \rightarrow \text{CG} \) | 1                                    | 3                                    | 1                                    | 2                                    | 0                                    | 2                                    |
| **Rate of base substitutions** | 4.6\( \pm 1.1 \) | 3.7\( \pm 1.5 \) | 3.3\( \pm 1.6 \) | 4.2\( \pm 1.3 \) | 4.2\( \pm 0.9 \) | 6.4\( \pm 1.2 \) |
| + Frame-shift\(^2\) | 0.6\( \pm 0.25 \) | 0.5\( \pm 0.3 \) | 0.5\( \pm 0.1 \) | 0.2\( \pm 0.06 \) | 1\( \pm 0.2 \) | 0.1\( \pm 0.06 \) |
| - Frame-shift\(^2\) | 0.8\( \pm 0.25 \) | 0.7\( \pm 0.2 \) | 0.7\( \pm 0.1 \) | 0.3\( \pm 0.06 \) | 0.2\( \pm 0.06 \) | 0.2\( \pm 0.06 \) |
| **Deletions**       | 9                                    | 27                                   | 4                                    | 1                                    | 2                                    |                                      |
| **Duplications**    | 5                                    | 12                                   |                                      |                                      |                                      |                                      |
| **Other (complex)\(^3\)** | 1                                    |                                      |                                      |                                      |                                      |                                      |
| **Total**           | 15                                   | 14                                   | 14                                   | 10                                   | 87                                   | 99                                   |

\( a \) the following nomenclature is used: “\( \text{t} \)” and “\( \text{ori} \)” refer to the telomere and the replication origin 3006/7, respectively; “\( \text{t} < \)” and “\( \text{t} > \)” indicate the RTS1-RFB and its polarity (\( \text{t} < \) blocks replication forks moving from the origin 3006/7 towards the telomere, and \( \text{t} > \) blocks replication forks moving from the telomere towards the origin 3006/7).

\( b \) number of events.

\( c \) event/cell/division \( \times 10^{-9} \) ± standard error. The % of each mutation events was used to balance the rate of mutation.

\( d \) value not significantly different from those for the \( t \leq \text{ura4} < \text{ori} \) strain (spontaneous events).

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Rad22<sup>Rad52</sup> (Figure 1B and [20]), such that even the two-fold induction in replication slippage by fork arrest in the rad22<sup>d</sup> strain was surprising. The rad22<sup>d</sup> strain accumulates suppressors involving the Fbh1 helicase that limits Rhp51Rad51-dependent recombination at blocked replication forks [50,51]. Therefore, we analyzed replication slippage in the indicated strains and conditions. The rate of Ura<sup>+</sup> revertants was calculated using the rate of the median from at least 11 independent cultures. Values in brackets indicate the 95% confidence interval. Statistical significance was detected using the nonparametric Mann-Whitney U test. C. Rate of replication slippage in the indicated strains and conditions. The rate of Ura<sup>+</sup> revertants was calculated using the rate of the median from at least 11 independent cultures. Values in brackets indicate the 95% confidence interval. Statistical significance was detected using the nonparametric Mann-Whitney U test. D. Serial tenfold-dilutions from the strains indicated spotted onto the media indicated after cell growth with (Rtf1 repressed) or without (Rtf1 expressed) thiamine. RTS1-RFB activity “−” refers to the strain t-ura4-dup20-ori and “+” refers to the strain t-ura4-dup20<sup>−</sup>. E. Kinetics of Ura<sup>+</sup> revertants frequency for the strains indicated as a function of the number of generations after thiamine removal. RTS1-RFB activity “−” refers to the strain t-ura4-dup20-ori and “+” refers to the strain t-ura4-dup20<sup>−</sup>. The values reported are the means of two experiments. F. The rate of replication slippage/generation for the strains indicated (t-ura4-dup20<sup>−</sup>) or without (t-ura4-dup20-ori) the active RTS1-RFB. The rate was calculated from the slope of the curves presented in panel F. The values reported are means of three independent experiments and error bars correspond to SE.

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agents or UV-C-induced DNA damages and the frequency of Ura⁺ revertants was scored. Three hours of treatment with either the topoisomerase I inhibitor camptothecin (CPT) or mitomycin C (MMC), an inter-strand cross-linking agent (ICls), increased the frequency of Ura⁺ revertants by 3 to 4 fold in the ura4-dup20 strain (Figure 5A and 5B). At equivalent survival (70–90%), DNA-damages induced by a dose of 100 J/m² of UV-C did not increase the frequency of Ura⁺ revertants in the ura4-dup20 strain. Increasing the UV-C dose (150 J/m²) resulted in an increased reversion effect.

The other ura4 alleles exhibited an opposite behaviour pattern. As expected, UV-C-induced DNA damages, but not CPT or MMC treatment, increased the frequency of Ura⁺ revertants of the base-substitution and the frame-shift mutants (Figure 5A). Thus, replication slippage, unlike other point mutations, appears to be a mutation event specifically induced by replication stress.

Hydroxyurea (HU) that prevents the bulk of dNTP synthesis during S-phase by inhibiting the ribonucleotide reductase, results in a slow-down of fork progression which did not induce replication slippage (Figure 5A). In contrast, CPT and MMC treatments that lead to replication stress by causing fork collapse induced replication slippage. Homologous recombination is repressed during HU treatment and recombination proteins are recruited to collapsed but not stalled forks [52–54]. Consistent with this, we found that the rad22-d mutant is highly sensitive to acute exposure to CPT, but not to HU (Figure S4). Thus, acute exposure to HU results in stalled forks that recover without...
recombination, whereas recombination may be required for restarting forks that have collapsed due to CPT or MMC treatment. We confirmed that CPT-induced replication slippage results from collapsed forks and was S-phase specific: the ura4-dup20 strain was synchronized in early S-phase by HU treatment and released into S-phase with or without CPT. HU-synchronization and release into DMSO (used as vehicle for CPT) did not induce replication slippage. In contrast, the release of cells into S-phase in the presence of CPT stimulated replication slippage up to 12 fold (Figure 5C). These data indicate that CPT-induced fork collapse results in error-prone DNA synthesis characterized by replication slippage. These experiments further support the view that replication slippage results from recovery of collapsed forks by recombination and point out that the RTS1 barrier is representative of collapsed forks restarted by homologous recombination.

TLS–DNA polymerases are not involved in fork-arrest-induced replication slippage

To investigate the inaccuracy of the DNA synthesis occurring immediately following the restart of collapsed forks, we analysed the involvement of TLS-DNA polymerases. In fission yeast, TLS pathways require either mono- or poly-ubiquitination of the clamp loader PCNA on lysine 164 [55]. We found that mutating this lysine to arginine residue did not affect replication slippage induced by the RTS1-RFB. None of Rev1, Rev3 or DinB DNA polymerases were required for fork-arrest-induced replication slippage (Figure S5). Therefore, the error-prone DNA synthesis associated with fork recovery by recombination does not rely on TLS DNA polymerases activity.

The mismatch repair pathway does not counteract fork-arrest-induced replication slippage

The mismatch repair (MMR) pathway is temporally coupled to DNA replication, and MMR components are associated with replication centres [56]. The heterodimer Msh2/Msh6 recognises mispaired DNA and Msh2/Msh3 recognises small DNA loops up to 31 bases long, arising from replication slippage [57]. The failure to repair small DNA loops results in more frequent insertions and deletions [58]. Therefore, MMR activity could potentially lead to an underestimation of the extent of fork-arrest-induced replication slippage. However, replication slippage induced by the RTS1-RFB activity was as frequent in msh2-d, msh6-d and msh3-d strains as in wild-type control. Also, spontaneous replication slippage at ura4 (without RTS1-RFB) was unaffected by the absence of MMR proteins (Figure S5). Therefore, there is no evidence that MMR repairs small DNA loops (20 nt) in fission yeast and fork-arrest-induced replication slippage is not counteracted by MMR in our model system.

Discussion

Using conditional fork arrest constructs, we studied the consequences for genome instability of impediments to replication forks progression. A single fork arrest results in large-scale genomic changes and mutations that occur during recombination-mediated fork recovery (Figure 6). Inappropriate ectopic recombination at arrested forks results in GCRs, whereas appropriate restarting of the fork on the initial template results in error-prone DNA synthesis. GCRs and mutations at collapsed forks are genetically separable: Rqh1 limits fork-arrest-induced GCRs but not mutations (Figure 2D and Figure 3B). We demonstrate here that collapsed forks whose progression resumes by recombination lose accuracy during DNA synthesis, resulting in frequent intra-template switches. Thus, homologous recombination contributes to completion of DNA replication when forks progression is impeded but also fuels genome modifications both at the chromosomal and nucleotide level.
Non allelic homologous recombination (NAHR) between low copy number repeats (LCR) contributes to recombination-mediated GCRs in mitosis and meiosis. NAHR is responsible for translocations, deletions, inversions and loss of heterozygosity [59]. Ou et al. predicted 1143 LCR pairs in the human genome liable to mediate recurrent translocations [60]. In budding yeast, a single DSB is sufficient to mediate recombination-dependent translocation [61]. Here, we report that a single collapsed fork increases the rate of genomic deletion 40 fold, and that of translocation 5 fold. Fork-arrest-induced GCRs are mediated by NAHR between heterologous chromosomes. It is not clear whether fork arrest on both homologous repeats contributes to fork-arrest-induced GCRs. We could not address this question in our model system, because the RTS1 sequence close to the matl locus on chromosome II has a low RFB activity [62]. Also, the recruitment of recombination proteins at the RTS1 sequence near the matl locus is not regulated by the level of Rtf1 expression, showing that regulating Rtf1 expression was in itself insufficient to regulate the RTS1-RFB activity at the matl locus [35].

Inverted repeats (IRs) are structural elements that contribute to genome instability. Impediments to replication forks progressing through IRs favour their fusion to generate acentric and dicentric inverted inverted chromosomes [11,14]. IRs in humans can also trigger the formation of inverted genomic segments and complex duplication rearrangements by a replication-based mechanism [47]. Here, we report that IRs near collapsed forks can increase the rate of GCRs by up to 1,500 fold. This high level of GCRs cannot be explained merely by the addition of independent inter- and intra-chromosomal recombination events. Rather, our analyses suggest that IRs may stimulate tri-parental recombination events induced by template switching of nascent strands at collapsed forks, such that three homologous sequences are involved. Similarly, recombination-dependent translocations induced by a single DSB in budding yeast is proposed to be the consequence of tri-parental recombination [63,64]. One possible mechanism is that IRs-induced GCRs result from successive template switches initiated by nascent strands at collapsed forks, reminiscent of the multiple template switches during break-induced-replication (BIR) in budding yeast [33]. Interestingly, Rqh1 prevents fork-arrest-induced GCRs by limiting both inter- and intra-chromosomal recombination without affecting fork restart efficiency. Thus, tri-parental recombination might correspond to multiple and successive template switches between homologous repeats.

The high accuracy of DNA replication is compromised by impediments to fork progression, and recombination-mediated fork recovery results in decreased processivity of DNA synthesis (Figure 6). Recombination-induced mutations associated with DSBs or impeding DNA replication have been described previously. The formation of damaged single-stranded DNA during the resection of DSBs favours base-substitutions [65]. We detected the recruitment of the single-strand binding protein RPA up to 1.4 kb behind, but not ahead of the RTS1-arrested fork (data to be published), showing that fork-arrest-induced mutation is not correlated with damaged single-stranded DNA exposed behind collapsed forks. Nevertheless, there were rare replication slippage occurred behind RTS1-arrested forks (in the in- and on-construct), suggesting that resumption of DNA synthesis can in some cases occur at a position behind the site of the collapsed fork. Recombination-dependent DNA synthesis occurring outside S-phase is also highly inaccurate during gene conversion and BIR, resulting in either template switches, base-substitutions or frame-shifts [33,34,66]. Elevated dNTP pools, due to activation of the DNA-damage checkpoint in G2 cells, contributes to the generation of mutations when hundreds of kbs are synthesised during BIR [34]. Here, we demonstrate that during normal S-phase progression, a single collapsed fork, restored by recombination, results in replication slippage up to 1.2 Kb away from the initial resuming point.

Recombination-induced replication slippage has been reported previously. In fission yeast, a defect in pol alpha (swi7-44 mutant) is associated with a recombination-mediated mutator phenotype characterized by an increased frequency of base-substitutions and Del/Dup between short direct repeats [67]. DNA-polymerase kappa (DnB) and zeta (Rev3) are responsible for the increased base-substitution rate, but the DNA-polymerases involved in Del/ Dup mutations were not identified [69]. In budding yeast, a defect in polymerase delta (pol3-4 mutant) results in an increased level of replication slippage, mediated by homologous recombination [49]. In contrast, the increased rate of replication slippage in the absence of the accessory subunit of polymerase delta, Pol32, does not depend on a functional recombination pathway [69]. Here, we report that recovery of collapsed forks by recombination is specifically associated with replication slippage. Nonetheless, spontaneous replication slippage events are also increased in strains mutated for recombination genes (Figure 4D–4F), Pol32 is required for BIR and replication-induced template switches leading to segmental duplication [70,71]. Recombination is responsible for only half of these segmental duplications. Thus, it is possible that fork-restart mechanisms dependent on Pol32 and homologous recombination are prone to replication slippage and that in the absence of these pathways alternative micro-homology mediated mechanisms are revealed.

We suggest that at least two steps of the recombination-based fork recovery mechanism can compromise genome stability (Figure 6). At an initial stage, recruitment of recombination proteins on stalled nascent strands favours both fork recovery and ectopic template switches leading to GCRs. At a later stage, once the replisome has been reconstituted and the fork has resumed its progression, the nascent strands are prone to intra-template switching leading to replication slippage. The reasons for the inaccuracy of DNA synthesis associated with restarted forks during scheduled DNA replication (i.e. in S-phase) remain to be determined. One possibility is that one or more factors are missing in the rebuilt replisome during recovery by recombination. Oncogene-induced replication stress results from unbalanced DNA replication that contributes to genome instability in precancerous cells [12]. Completion of DNA replication in such stress conditions is likely to rely on recombination-mediated fork recovery that in turn generates genome instability. Insertions/deletions flanked by micro-homology, responsible for copy number variations (CNVs), have been identified both in cancer cells and also in response to replication inhibition [72,73]; their reported sizes are between 1 Kb and several tens of mega bases, but the analysis of these features has been limited by the resolution of array-based genomics approaches. Sub-microscopic insertions/deletions flanked by micro-homology have been also described at loci in which replication origins are scarce, including the human fragile site FRA3B, the instability of which is a consequence of replication stress [74]. Interestingly, homologous recombination contributes to the stability of fragile sites by facilitating complete replication or by repairing gaps and breaks at these sites. Thus, we propose that micro-homology-mediated CNVs could be viewed as scars left by error-prone replication forks restarted by recombination.

Materials and Methods

Standard genetic and molecular biology

Strains used were constructed by standard genetic techniques and are listed in Table S1. 2DGE was performed as previously.
described [20]. To create *ura4-dup20* and *ura4-dup22* alleles associated or not with the *RTS1*-RFB, genomic DNA was isolated from selected 5-FOA<sup>R</sup> cells containing duplications identified by sequencing. A PCR fragment containing duplications within the *ura4* ORF was amplified using the following primers: TGTTCACCAGCACCATTTCCTTCCACG and TCAAGCGTTTATTTTCCAGTCTCAGG. The PCR products were purified and used to sequence. A PCR fragment containing duplications within the *RTS1* ORF, TGAATCCTCCGTTCAGTAGG and TTTAACGAGGCTCTTCTCTAGC to amplify the translocation junction.

**ura4** loss assay

A minimum of 11 independent single colonies from appropriate strains growing with or without thiamine were inoculated in 10 ml of non-selective media (with or without thiamine) and grown to stationary phase. Appropriate dilutions were plated on supplemented YEA to determine plating efficiency and on 5-FOA-containing plates. Colonies were counted after 3–7 days of incubation at 30°C. Rate of *ura4* loss was determined with the method of the median and data are presented on Table 1. Statistical significance was detected using the nonparametric Mann-Whitney U test.

PCR assays and sequencing to determine the rates of genomic deletion, translocation, and mutation

At least 200 5-FOA<sup>R</sup> colonies per strain and condition were subjected to PCR analysis with the following primers: AAAAAACCGGCAACACGGGCA and GTTTAAGCTAGGTGTCGGTCCG and AAGGACTGTTGTTCTTCTAGC to amplify *ura4* ORF, TGAATCCTCCAGTCTCAGG and AAGGACTGTTGTTCTTCTAGC to amplify *ura4* and TGTTCACCAGCACCATTTCCTTCCACG and TCAAGCGTTTATTTTCCAGTCTCAGG and AAGGACTGTTGTTCTTCTAGC to amplify the translocation junction. The amplified *ura4* fragments were then sequenced on both strands, with the primers used to amplify the *ura4* ORF. Only mutations present on both strands were considered to determine mutation spectra. Deletions, mutations and translocations were scored as percentages of all events and these values were used to balance the rates of *ura4* loss to determine the respective rates of deletion, mutation and translocation (see Figure S1 for deletion and translocation rates and Figure S2 for mutation rates).

The fork-arrest-induced deletion, translocation and mutation rates (Figure 2C–2E and Figure 3A) were calculated by subtracting the rate obtained in presence of thiamine (Rtf1 being repressed, OFF) from the rate obtained in absence of thiamine (Rtf1 being repressed, ON). This method allows the spontaneous fork arrestance instability of IRs and the leakiness of the *RTS1*-RFB activity to be disregarded to determine strictly the rate of events induced by fork-arrest. The nonparametric Mann-Whitney U test was used to test for statistically significant differences.

Reverse mutation assay

Exponentially growing cells were treated with 20 mM of HI, 40 µM of CPT or 1 mM of MMC. At indicated times, samples were taken and appropriate dilutions were plated on supplemented minimal media to determine plating efficiency and on uracil-free plates. Colonies were counted after incubation at 30°C for 5–7 days and the frequency of *Ura<sup>+</sup>* colonies was determined.

Replication slippage assay using *ura4-dup20* and *ura4-dup22* strains

For strains showing a slow growth phenotype (recombination mutants), the frequency of *Ura<sup>+</sup>* revertants was determined as a function of the number of generations experiencing fork arrest at *ura4*. Exponentially growing 5-FOA<sup>R</sup> cells were washed twice in water and used to inoculate uracil-containing media without thiamine. Every 24 hours, cells were counted to determine the number of generations, and appropriate dilutions were plated on supplemented minimal media and on uracil-free plates. Colonies were counted after incubation at 30°C for 5–7 days and the frequency of *Ura<sup>+</sup>* colonies was determined. The slope of the curves presented on Figure 4F corresponds to the rate of replication slippage/generation. For strains showing similar growth to wild-type cells, a single 5-FOA<sup>R</sup> colony was grown on uracil-containing plates with or without thiamine for 2–3 days, and then grown in uracil-containing media with or without thiamine for 2 days at 30°C. Appropriate dilutions were plated on supplemented minimal media and on uracil-free plates. Colonies were counted after incubation at 30°C for 5–7 days and the frequency of *Ura<sup>+</sup>* colonies was determined.

**Supporting Information**

**Figure S1** Fork-arrest results in GCRs in a recombination-dependent manner. A. The rate of deletion for indicated strains, in the presence (Rtf1 repressed) and in the absence (Rtf1 expressed) of thiamine. Numbers of *RTS1* repeats present in the *S. pombe* genome and the presence of a visible fork arrest (based on 2DGE presented on Figure 1B) are given for each strain. The % of deletion events, as determined by the PCR assay, was used to balance the rate of *ura4* loss. The values reported are means of at least 3 independent median rates. Error bars correspond to the standard error (SE). Statistically significant fold differences in the rate of deletion events between the Rtf1 “repressed” and “expressed” conditions are indicated with an *, B and C. Rate of deletion (B) and translocation (C) for the strains indicated; ON and OFF refers to the *RTS1*-RFB being active or not, respectively. The % of deletion and translocation events, as determined by the PCR assay, was used to balance the rate of *ura4* loss. The values reported are means at least 3 independent median rates. Error bars correspond to SE. Statistically significant fold differences in the rates of deletion or translocation events between the “OFF” and “ON” conditions are indicated with an *. Translocation events (based on the detection of the *TLII/TLIII* PCR product) were not detected in *rad22-d* or *rhb51-d* strains, whatever the conditional fork arrest construct.

**Figure S2** Fork-arrest induces replication slippage. A. The rate of mutation for indicated strains, in the presence (Rtf1 repressed) and in the absence (Rtf1 expressed) of thiamine. Numbers of *RTS1* repeats present in the *S. pombe* genome and the presence of a visible fork arrest (based on 2DGE presented on Figure 1) are given for each strain. The % of mutation events, as determined by the PCR assay and sequencing, was used to balance the rate of *ura4* loss. The reported values are means of at least 3 independent median rates. Error bars correspond to SE. Statistically significant fold differences in the rate of mutation events between the Rtf1 “repressed” and “expressed” conditions are indicated with an *, B and C. The frequency of *Ura<sup>+</sup>* revertants for the indicated strains and expected reverted mutations are indicated in the table. #1 and #2 correspond to two independent mutated strains for each type of mutation.
Figure S3 Features of replication slippage induced by fork arrest. A. Table of deletion/duplication and micro-homology features. B. Map of deletion and duplication events observed within the *ura4* ORF in the *t-ura4-dup20* construct upon fork arrest. Del and Dup stand for deletion and duplication, respectively. (TIF)

Figure S4 Sensitivity of *rad22-d* strain to acute exposure to 20 mM of HU or 20 μM of CPT. The values reported are means of two to four independent experiments. Error bars indicate the standard error of the mean (SEM). (TIF)

Figure S5 Fork-arrest-induced replication slippage is independent of the post-replication repair and mismatch repair, A−C. Left panels: Serial tenfold-dilutions of indicated strains cultured in thiamine-free medium spotted onto the medium indicated. *RTS1-RFB* activity “−” refers to the strain *t-ura4*, and “+” refers to the strain *t-ura4-dup20*. Right panels: The frequency of Ura+ revertants from the strains indicated (*t-ura4-dup20* associated or not with the *RTS1-RFB*) in the conditions indicated. The values reported are means of at least three independent experiments and error bars correspond to SEM. (TIF)

Table S1 Strains used in this study. (DOCX)

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Author Contributions
Conceived and designed the experiments: II AC SAEL. Performed the experiments: II VP YC NJ. Analyzed the data: II YC NJ KF AG SAEL. Wrote the paper: SAEL.

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