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Vincent Pagès, Gerard Mazon, Karel Naiman, Gaëlle Philippin, Robert P Fuchs. Monitoring bypass of single replication-blocking lesions by damage avoidance in the *Escherichia coli* chromosome.. *Nucleic Acids Research*, 2012, 40 (18), pp.9036-9043. 10.1093/nar/gks675 . hal-01446625

HAL Id: hal-01446625

<https://hal.science/hal-01446625>

Submitted on 15 Mar 2017

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Monitoring bypass of single replication-blocking lesions by damage avoidance in the *Escherichia coli* chromosome

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Received May 23, 2012; Revised and Accepted June 19, 2012

ABSTRACT

Although most deoxyribonucleic acid (DNA) lesions are accurately repaired before replication, replication across unrepaired lesions is the main source of point mutations. The lesion tolerance processes, which allow damaged DNA to be replicated, entail two branches, error-prone translesion synthesis (TLS) and error-free damage avoidance (DA). While TLS pathways are reasonably well established, DA pathways are poorly understood. The fate of a replication-blocking lesion is generally explored by means of plasmid-based assays. Although such assays represent efficient tools to analyse TLS, we show here that plasmid-borne lesions are inappropriate models to study DA pathways due to extensive replication fork uncoupling. This observation prompted us to develop a method to graft, site-specifically, a single lesion in the genome of a living cell. With this novel assay, we show that in *Escherichia coli* DA events massively outweigh TLS events and that in contrast to plasmid, chromosome-borne lesions partially require RecA for tolerance.

INTRODUCTION

Replication of damaged deoxyribonucleic acid (DNA) is a universal problem faced by all organisms. DNA lesions arise continuously due to endogenous or environmental agents. Despite the efficient action of numerous repair systems, some lesions that escape these repair mechanisms are present when the genome is being replicated. To overcome the challenge of replicating damaged DNA,

cells have developed several lesion tolerance mechanisms that enable the replication machinery to bypass sites of damaged DNA. The conceptually simplest procedure of bypassing lesions encountered during DNA replication is translesion synthesis (TLS), whereby the replicative polymerase is replaced by a specialized polymerase that can synthesize the new DNA strand across the site of damage. This process, although not always mutagenic, is inherently error-prone [for a review, see (1)]. On the other hand, error-free bypass of DNA lesions is possible by using the information present in the undamaged sister chromatid (2–6). These processes, collectively referred as damage avoidance (DA), embrace several pathways related to homologous recombination and are still poorly defined to date. Failure to achieve TLS or DA will lead to incomplete replication of the genome and therefore to the cell death.

Until today, studies on the consequences of lesions in DNA *in vivo* are usually limited to the analysis of induced mutations. Indeed, mutations that result from TLS events represent the major biological consequence of the presence of lesions in DNA because DA events are considered to be error-free. However, the respective proportion of TLS versus DA events within DNA damage tolerance events is presently not known. Moreover, most induced-mutagenesis studies involve treatment of cells with damaging agents that introduce a variety of different lesions randomly distributed all over the genome. Consequently, the DNA lesion that causes a given mutation can only be guessed. Over the past recent years, many studies aimed at uncovering the mechanism of TLS by introducing single site-specific lesions within plasmids. This approach has been instrumental to unravel the complexity of TLS pathways and the genetics of mutagenesis in *Escherichia coli*, *Saccharomyces cerevisiae* and recently in human

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cells (7–9). In the present paper, we show that, during plasmid replication, a single blocking lesion triggers replication fork uncoupling (10) accompanied by full unwinding of the two sister chromatids. As DA events require the two sisters to be maintained in close proximity, plasmid systems are not suited for the analysis of DA events. To overcome the limitations inherent to plasmids, we developed a methodology that allows single lesions to be site-specifically introduced into the *E. coli* chromosome. Under these assay conditions we show that, in contrast to plasmid-borne lesions, tolerance of chromosomal DNA lesions by DA is partially dependent on a functional *recA* gene. Until now, it was essentially impossible to monitor the rescue of a blocked replication fork *in vivo* by lack of an appropriate methodology. Indeed, current assays involve lesions randomly distributed over the chromosome, making it impossible to determine the structure and the dynamic of a fork blocked at a specific lesion site.

MATERIALS AND METHODS

Plasmids construction

pVP135 expresses the integrase and excisionase (*int-xis*) genes from lambda under control of a *trc* promoter that has been weakened by mutations in the –35 and the –10 region (11). Transcription from *P_{trc}* is regulated by the *lac* repressor, supplied by a copy of *lacI_q* on the plasmid. The vector is derived from pDSW206 (11) by replacing chloramphenicol resistance by a kanamycin resistance cassette. A polymerase chain reaction (PCR) fragment containing the *int-xis* operon from pTSC29cxi (12) was cloned into the *NcoI*-*PstI* restriction sites.

pVP146 is derived from pACYC184 plasmid where the chloramphenicol resistance gene has been deleted by *Bsa*AI digestion and re-ligation. This vector, which carries only the tetracycline resistance gene, serves as an internal control for transformation efficiency.

pVP141-144 and pGP1-2 are derived from pLDR9-attL-*lacZ* plasmid (12). These plasmid vectors contain the following genes characteristics: the ampicillin resistance gene, the R6K replication origin that allows plasmid replication only if the recipient strain carries the *pir* gene (13), and the 5' end of *lacZ* gene in fusion with the attL site-specific recombination site of phage lambda. The P'3 site of attL has been mutated (AATCATTAT to AATTATTAT) to avoid the excision of the plasmid once integrated (14). pVP141-144 and pGP1-2 are produced in strain EC100D *pir*-116 (from Epicentre Biotechnologies—cat# EC6P0950H) in which the *pir*-116 allele supports higher copy number of R6K origin plasmids.

Construction of vectors carrying a single lesion

Vectors for integration

Duplex plasmids carrying a single lesion were constructed following the gap-duplex method previously described (15). A 13-mer oligonucleotide, 5'-GCAAGTTAACAC G, containing no lesion, a cyclobutane pyrimidine dimer (TT-CPD) or a thymine-thymine pyrimidine(6-4) pyrimidone photoproduct [TT(6-4)] lesion (underlined)

was inserted into the gapped-duplex pGP1/2 leading to an in frame *lacZ* gene. Because the G-AAF (N-2-acetylaminofluorene) lesion can be bypassed by two distinct pathways (7), two vectors were constructed to monitor all TLS events. A 15-mer oligonucleotide containing or not a single G-AAF adduct (underlined) in the *NarI* site (ATCACCGGCGCCACA) was inserted into a gapped duplex pVP141/142 or pVP143/144, leading respectively to an in frame *lacZ* gene, and a +2 frameshift *lacZ*. Therefore, the construct pVP141/142 *Nar3^{AAF}/Nar+3* monitors TLS0 events, whereas pVP143/144 *Nar3^{AAF}/Nar+3* monitors TLS–2 events.

Replicating vectors

pCUL– series and pCUL+ series contain the G-AAF lesion, respectively, in the lagging and leading strand. pCUL–*Nar0/Nar+3*, pCUL–*Nar3^{AAF}/Nar+3*, pCUL+*Nar0/Nar+3* and pCUL+*Nar3^{AAF}/Nar+3* were constructed as previously described (10).

Strains

All strains used in the present study for site-specific recombination are derivative of strain FBG152 (16). Strain FBG152 is derived from strain MG1655 in which the original λ *attB* site was replaced by an artificial promoterless operon carrying *attR* fused to the 3' part of *lacZ* upstream of the *aadA* gene and between *ybhC* and *ybhB* (at ~17th min) (16). Gene disruptions of *recA*, *mutS*, *uvrA* and *phrB* were achieved by the one-step PCR method (17). The following FBG152-derived strains were constructed by P1 transduction: EVP23 (FBG152 *uvrA::f_{rt}* *mutS::f_{rt}*), EVP123 (FBG152 *uvrA::f_{rt}* *mutS::f_{rt}* *recA::f_{rt}*), EVP184 (FBG152 *uvrA::f_{rt}* *mutS::f_{rt}* *phrB::f_{rt}*) and EVP206 (FBG152 *uvrA::f_{rt}* *mutS::f_{rt}* *phrB::f_{rt}* *recA::f_{rt}*). All strains carry plasmid pVP135 that allows the expression of the *int-xis* under the control of IPTG. Following the site-specific recombination reaction, the lesion [G-AAF, TT-CPD or TT(6-4)] is located in the leading strand of strain FBG152.

Strand segregation assay protocol

After transformation of plasmid pCUL-*Nar3^{AAF}/Nar+3* or pCUL+*Nar3^{AAF}/Nar+3* in strain JM103*uvrAmutS*, and plating on X-gal indicator plates, individual blue colonies (i.e. colonies in which TLS has occurred) are collected and grown in Luria Broth (LB) media containing ampicillin. Plasmids extracted from a single colony are re-transformed into JM103 and plated on X-gal indicator plates to score blue and white clones.

Integration protocol—measurement of lesion tolerance pathways: TLS and DA

LB (50 ml) containing kanamycin to maintain plasmid pVP135 that expresses the *int-xis* operon and 200 μ M of IPTG (to induce the expression of *int-xis*) are inoculated with 500 μ l of an overnight starter. When the culture reaches OD⁶⁰⁰ \approx 0.5, cells are washed twice in water, once in 10% glycerol and finally re-suspended in 200 μ l of 10% glycerol and frozen in 40 μ l aliquots.

To the 40 μ l aliquot of cells, 1 ng of the lesion carrying vector mixed with 1 ng of the internal standard (pVP146) were added, and electroporated (in a GenePulser Xcell from BioRad, 2.5 kV, 25 μ F, 200 Ω). 1 ml of SOC medium containing 200 μ M IPTG is then added, and cells are incubated 1 h at 37°C. Part of the cells are plated on LB + 10 μ g/ml tetracycline to measure the transformation efficiency of plasmid pVP146 (internal transformation standard), and the rest is plated on LB + 50 μ g/ml Ampicillin and 80 μ g/ml X-gal to select for integrants (Amp^R) and TLS events (blue colonies). The integration rate is about 2000 clones per picogram of non-damaged vector for a wild-type (WT) strain.

Following the integration of the damaged vector, blue colonies represent TLS events, whereas white colonies represent DA events. The relative integration efficiencies of damaged vectors compared with their non-damaged homologues, and normalized by the transformation efficiency of pVP146 plasmid in the same electroporation experiment, allow the overall rate of tolerance of the lesion to be measured. DA events are estimated by subtracting TLS events from the total lesion tolerance events.

RESULTS AND DISCUSSION

Evidence for strand uncoupling during replication of a plasmid containing a single replication-blocking lesion

We developed an assay to analyse the products that are generated during replication of a plasmid vector that contains a single replication-blocking lesion. For this purpose, we used an adduct formed by covalent binding of the chemical carcinogen *N*-2-acetylaminofluorene (AAF) to the C-8 position of guanine (G-AAF) (18). This adduct blocks replication similarly to the ultraviolet (UV)-induced lesions such as TT-CPDs or 6-4 photoproducts (19). The plasmid construct also contains a short sequence heterology in the complementary strand across from the single adduct (18). The lesion is located in the *lacZ* gene such that TLS events give rise to lac⁺ plasmids, while the complementary strand is out of frame and thus gives rise to lac⁻ plasmids. These constructs are transformed into a recipient cell (primary transformation) and plated on X-gal indicator plates to identify colonies that have experienced TLS (TLS colonies) as blue colonies (*lacZ*⁺) (Figure 1A). Individual 'TLS colonies' are further analysed to determine the average proportion of blue and white plasmids within the cells by extracting plasmid DNA and retransforming it into an indicator strain to determine the proportion of white and blue plasmids (secondary transformation) (Figure 1A). When a lesion-free construct is analysed, essentially all primary transformants are blue because the blue phenotype is dominant. Analysis of the plasmid content of individual blue colonies by secondary transformation shows that, as expected, the progeny of the two strands are about equally represented in the majority of colonies analysed (although there is a slight, as yet unexplained, bias towards <50% of blue colonies, Figure 1B). When a construct that monitors TLS events across a single G-AAF adduct is introduced in a non-SOS induced WT strain, only 2–3% of colonies are

blue (TLS colonies) in the primary transformation, showing that TLS across G-AAF is a rare event. Analysis of individual blue colonies shows that most 'TLS colonies' contain <20% of blue plasmids (and thus >80% white plasmids) (Figure 1B). The same distribution is seen whether the lesion is located in the leading or in the lagging strand. These results illustrate the 'replisome uncoupling' phenomenon as initially identified by molecular analysis of replication intermediates (10). It was concluded from these experiments that when a replisome encounters a replication-blocking lesion, synthesis in the lesion strand is transiently arrested, whereas replication of the undamaged strand continues (10) (Figure 1C). It is estimated that following a leading strand block the replicative helicase can unwind the parental strands beyond the blocking lesion over a distance of at least 2–3 kb (10). Therefore, the commonly used single-adducted plasmid constructs become fully unwound leading to unbalanced amplification of the non-damaged strand with respect to the damaged strand. Replication of the damaged strand lags behind due to the time required to perform TLS (Figure 1C) (10).

In conclusion, these data suggest that due to their limited size, plasmids are inappropriate tools for the analysis of DA pathways that require the two sister chromatids to be maintained in close proximity to undergo a recombination-based transfer of information that characterize DA events (20). When the lesion is located in a larger genome such as a chromosome, even if the helicase travels over a certain distance beyond the replication blocking lesion, the fork will eventually come to an arrest as observed by electron microscopy in *S. cerevisiae* after UV irradiation (21). In an attempt to investigate DA pathways, we developed a method that allows a single lesion to be located site-specifically in a genome.

Site-specific insertion of a single lesion in the *E. coli* chromosome

To overcome the intrinsic limitations of plasmid systems we developed a methodology to insert a single lesion at a precise chromosomal location in *E. coli*. The experimental system (Figure 2A), based on phage lambda site-specific recombination, entails the following two major components: a recipient *E. coli* strain with a single *attR* site and a non-replicating plasmid construct containing the single lesion of interest and an *attL* site and an ampicillin resistance gene. The recombination reaction between *attL* and *attR* is controlled by ectopic expression of phage lambda *int-xis* proteins (12), and leads to the integration of the lesion-containing vector into the chromosome. Integrants are selected on the basis of their resistance to ampicillin. The chromosomal integration region carries the 3'-end of the *lacZ* gene fused to *attR*, whereas the remaining 5'-end is located on the incoming fragment in fusion with *attL*, so that precise integration restores a functional β -galactosidase gene (*lacZ*). The non-damaged opposite strand contains a short sequence heterology that inactivates the *lacZ* gene and serves as a genetic marker that allows strand discrimination (22). Precise integration of the vector at nucleotide resolution was shown by the

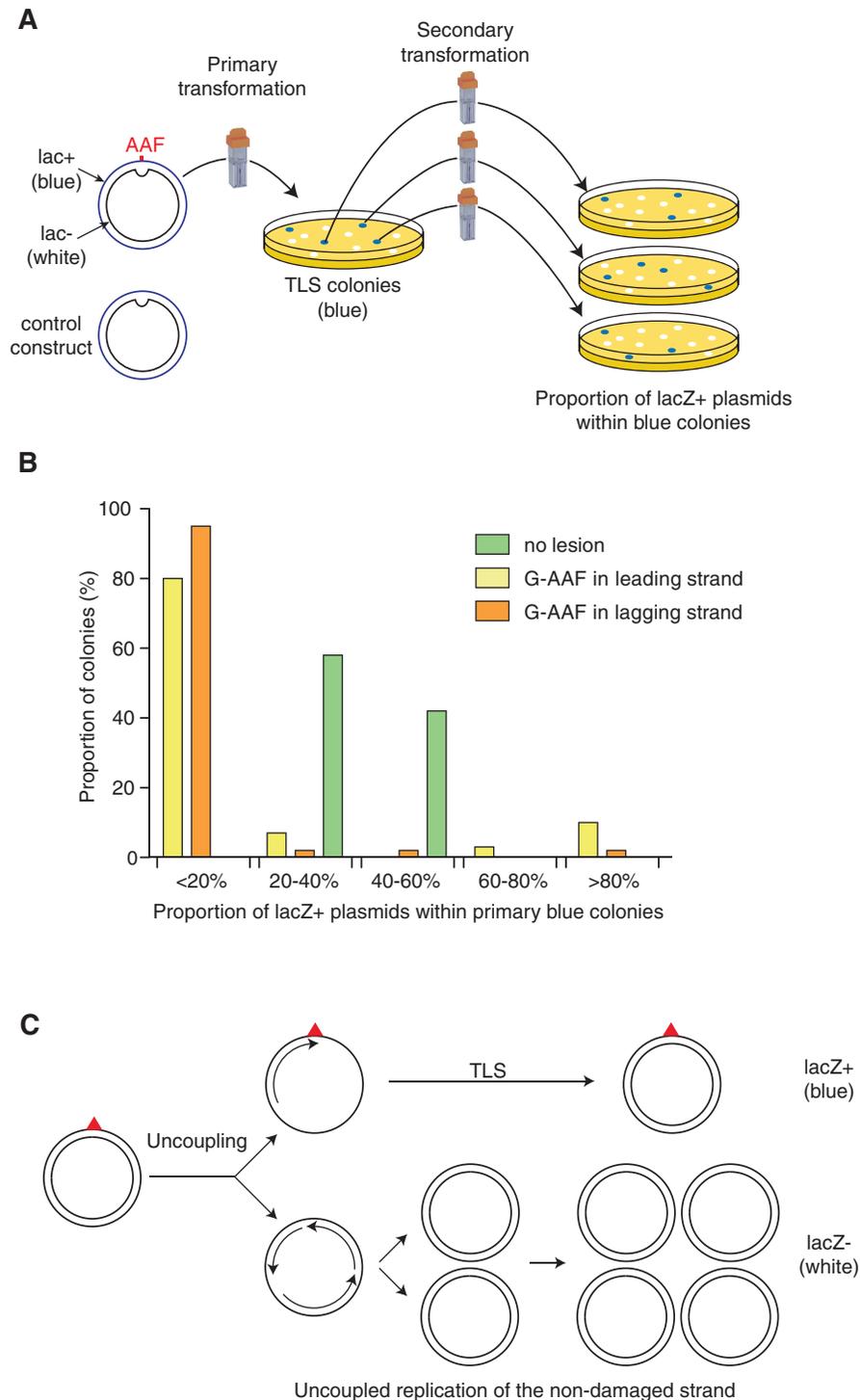


Figure 1. Strand segregation analysis using plasmids with a single replication-blocking lesion. (A) Outline of the strand segregation analysis assay (see the text). (B) Histogram showing the distribution, within individual primary blue colonies, of the proportion of lacZ+ versus lacZ- plasmids. For a lesion-free construct, most primary colonies are blue and carry a similar proportion of blue and white plasmids (although there is a slight, as yet unexplained, bias towards <50% of blue colonies). In contrast, most primary blue TLS colonies obtained with a plasmid carrying a single G-AAF adduct contain a small proportion of lacZ+ plasmids: 80–100% of primary TLS colonies contain <20% of lacZ+ plasmids. (C) Scheme representing full replication fork uncoupling in plasmids (sister-chromatid separation) following the encounter by the fork of a single replication-blocking lesion in one of the template strands: while replication is blocked at the damage site awaiting for TLS to occur, the non-damaged strand is fully replicated and amplified, leading to a high proportion of lacZ- (white) over lacZ+ (blue) plasmid progeny as revealed in the secondary transformation experiment.

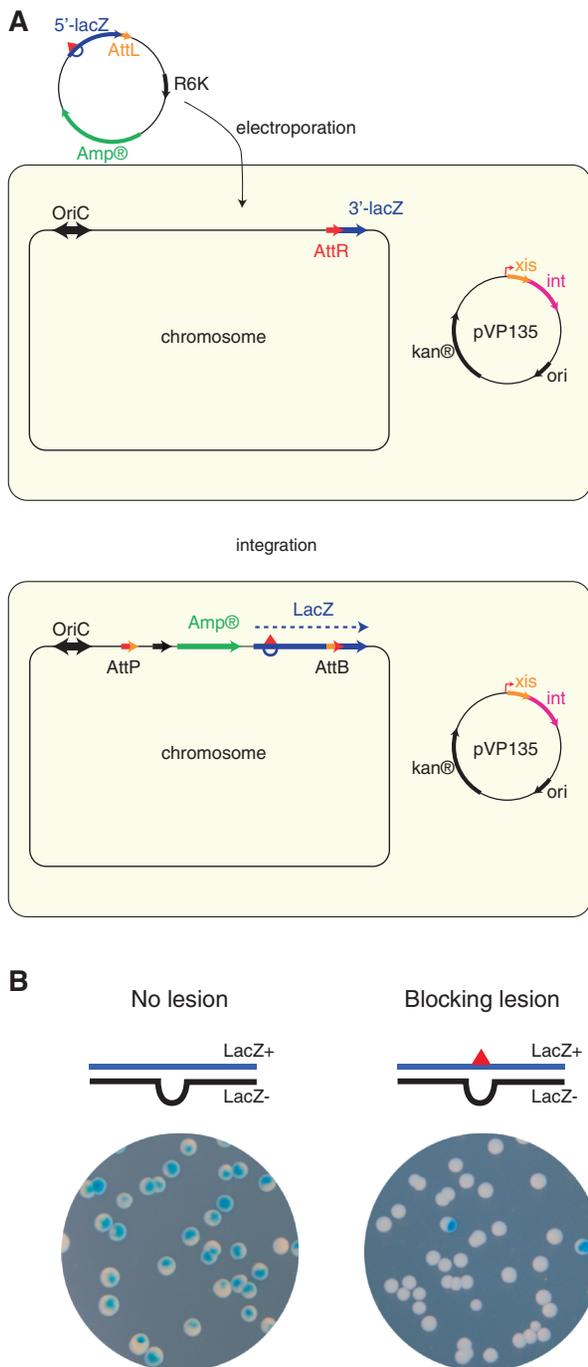


Figure 2. Grafting a single lesion into the chromosome. (A) The recipient strain contains a single attR integration site in fusion with the 3' end of *lacZ* gene at min 17 in the *E. coli* chromosome. Following ectopic expression of phage lambda int-xis, the lesion-carrying construct is introduced by electroporation. Its attL site will recombine with the chromosomal attR, leading to integration of the entire lesion-containing construct. Integration events are selected on the basis of their resistance to ampicillin. Integration at nucleotide level resolution restores a functional *lacZ* gene allowing these events to be monitored on X-gal indicator plates. (B) Chromosomal integration of *lac*⁻/*lac*⁺ heteroduplex constructs. As expected, integration of a lesion-free heteroduplex gives rise to essentially only sectorial colonies on X-gal plates. Replication of the G-AAF heteroduplex yields a *lac*⁺ event when the lesion is bypassed by TLS, whereas complementary strand replication yields a *lac*⁻ event. TLS events are characterized as sectorial colonies illustrating the coupling during replication of both strands even in the presence of a replication-blocking lesion.

presence of only LacZ⁺ colonies following the integration of a lesion-free construct. Individual integration events were analysed by PCR and found to be precisely located at the expected chromosomal position. The present experimental approach allows a single replication-blocking lesion to be located in the leading strand template at min 17. The way this block is tolerated either by TLS or via DA pathways can thus be studied genetically and at the molecular level.

Proper strand segregation is maintained during replication of a single chromosomal replication block

To observe what happens when a lesion escapes the repair mechanisms and is encountered by the replication fork, we conducted the following experiments in a nucleotide excision repair (*uvrA*) deficient strain. For experiments involving UV lesions, the photolyase gene (*phrB*) was additionally inactivated. Also, the recipient strain carried mutations in mismatch repair (*mutS*) to prevent correction of the heteroduplex. To assess normal chromosomal strand segregation we first integrated into the chromosome a lesion-free heteroduplex in which one strand carries a *lacZ*⁺ allele and the other strand a *lacZ*⁻ allele. Selection of individual integration events on *lac* indicator plates (X-gal plates) shows mostly blue:white sectorial colonies (Figure 2B). This is expected because chromosomal replication is semi-conservative and daughter chromosomes segregate in a one to one ratio. Molecular analysis of the sectorial colonies reveals that blue and white sectors contain the bacterial progeny that stems from the replication and subsequent segregation of the *lac*⁺ and *lac*⁻ strands, respectively. Occasional non-sectorial colonies are observed probably reflecting a low extent of heteroduplex correction before replication despite inactivation of mismatch repair.

Next, we integrated a *lac*⁺/*lac*⁻ heteroduplex containing a single G-AAF adduct in the *lac*⁺ strand. A majority of integration events form non-sectorial pure white colonies on the X-gal indicator plate. Molecular analysis of these events suggests that they result from the processing of the single adduct via a DA pathway. Indeed, these colonies only contain the specific *lac*⁻ allele carried in the non-damaged strand in the initial construct. However, it is presently not possible to eliminate the possibility that the damaged strand is degraded and that some white colonies originate from the sole replication of the non-damaged strand. A small amount of colonies are sectorial blue:white colonies (Figure 2B). Molecular analysis of the sectorial colonies reveals that the blue and white sectors contain the bacterial progeny that stems from the replication and subsequent segregation of the damaged and undamaged strands, respectively. The blue sectors contain the sequence of the lesion-containing strand and thus represent TLS events. The white sectors carry the *lac*⁻ allele initially present in the complementary strand (Figure 3). This observation illustrates 'coupled replication' of both strands even in the presence of a replication-blocking lesion in contrast to the situation of strand uncoupling encountered with plasmid probes (see previous text). These observations fully validate the use of the present

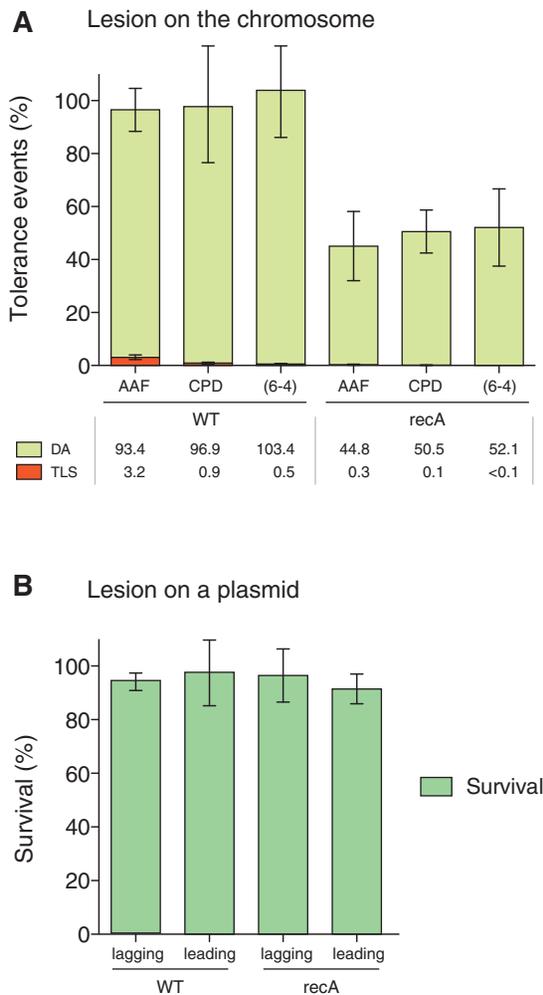


Figure 4. Survival of chromosomal and plasmid-borne replication-blocking lesions in WT and *recA* strains. **(A)** Relative integration efficiency (survival) and TLS events of a damage-containing construct relative to the corresponding damage-free construct, is measured in WT and *recA* strains. Three lesions, namely G-AAF, TT-CPD and TT(6-4), are introduced in the leading strand at position 17min in the chromosome by site-specific recombination. The data represent the average value and standard deviation of at least three independent experiments. **(B)** Relative transformation efficiency (survival) of a ColE1-based plasmid containing a single G-AAF adduct in the leading or lagging strand. In both orientations, for a G-AAF adduct located in a plasmid there is no loss of survival in a *recA* versus *rec+* strain. In contrast to plasmid-borne lesions, tolerance of chromosomal DNA lesions by DA is partially dependent on a functional *recA* gene.

discussed earlier, damaged plasmid replication exhibits uncoupled amplification of the two strands, a process that appears not to be impeded in a *recA*– strain in contrast to the observation made for the chromosome-borne lesions. This observation reinforces the notion that plasmids are not suitable tools for the analysis of genuine DA events as discussed in the previous section.

CONCLUSION

The present work provides the first experimental system to study TLS and DA pathways *in vivo*, in a quantitative

manner. Depending on the nature of the replication block, tolerance pathways occur either directly at the fork (on the fly) or at gaps left opposite lesion, i.e. behind the fork (26). Indeed, when encountering a strong replication block in the leading strand, the fork can restart by means of a priming event that occurs downstream of the lesion, leaving a gap that is subsequently processed by TLS or DA (31,32). We show that for common replication-blocking lesions, such as UV-induced photoproducts or G-AAF adducts, DA events massively outweigh TLS events (97% versus 1–3%) in non-SOS induced cells. The present methodology opens new ways to explore the metabolism of defined replication fork blocks at the chromosomal level. We highlight a major drawback of plasmid-borne lesion assays by showing that the extensive uncoupling between sister chromatids that occurs during plasmid replication precludes DA events to be monitored. This point is illustrated by the fact that, in contrast to plasmid-borne lesions, tolerance of chromosomal DNA lesions by DA is partially dependent on a functional *recA* gene. Until now, it was essentially impossible to monitor the repair of blocked replication forks *in vivo* by lack of an appropriate methodology.

This approach offers an experimental system able to open up the field of replication fork recovery in a way comparable with the development of double-strand break repair by the introduction of single double-strand breaks by HO and I-SceI endonucleases (33,34). In the present article, lesion tolerance pathways have been monitored using only genetic read-outs. However, given the efficiency of integration, i.e. 5–10% of all cells, we intend to develop molecular methods able to detect early replication intermediates that should allow us to monitor replication fork rescue in real time.

FUNDING

LIGUE Contre le Cancer (labellization 2011, in part); ANR grant ForkRepair [ANR 11 BSV8 017 01, in part]; fellowship from the Fondation ARC pour la recherche sur le cancer (to V.P. and K.N., in part). Funding for open access charge: LIGUE Contre le Cancer.

Conflict of interest statement. None declared.

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