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Inhibiting translation elongation can aid genome duplication in *Escherichia coli*

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**ABSTRACT**

Conflicts between replication and transcription challenge chromosome duplication. *Escherichia coli* replisome movement along transcribed DNA is promoted by Rep and UvrD accessory helicases with Δrep ΔuvrD cells being inviable under rapid growth conditions. We have discovered that mutations in a tRNA gene, *aspT*, in an aminoacyl tRNA synthetase, AspRS, and in a translation factor needed for efficient proline–proline bond formation, EF-P, suppress Δrep ΔuvrD lethality. Thus replication-transcription conflicts can be alleviated by the partial sacrifice of a mechanism that reduces replicative barriers, namely translating ribosomes that reduce RNA polymerase backtracking. Suppression depends on RelA-directed synthesis of (p)ppGpp, a signalling molecule that reduces replication-transcription conflicts, with RelA activation requiring ribosomal pausing. Levels of (p)ppGpp in these suppressors also correlate inversely with the need for Rho activity, an RNA translocase that can bind to emerging transcripts and displace transcription complexes. These data illustrate the fine balance between different mechanisms in facilitating gene expression and genome duplication and demonstrate that accessory helicases are a major determinant of this balance. This balance is also critical for other aspects of bacterial survival: the mutations identified here increase persistence indicating that similar mutations could arise in naturally occurring bacterial populations facing antibiotic challenge.

**INTRODUCTION**

Cell survival from one generation to the next relies on efficient and faithful replication of the genome. However, the template for replication frequently harbours obstacles that have the potential to interfere with the progression of replisome complexes, the macromolecular machines responsible for driving genome duplication. Protein–DNA complexes are important sources of such obstacles and those associated with transcription are especially problematic, in part because of their abundance and in part because they present multiple different challenges to replisome movement (1,2). One major challenge is the very high free energy of binding of transcription complexes that creates a need to disrupt many RNA polymerase–nucleic acid interactions as replication proceeds. Transcribing RNA polymerases (RNAPs) also pause frequently either spontaneously or upon encountering DNA template damage (3). Paused RNAPs can also diffuse backwards along the template resulting in displacement of the transcript 3′ end from the RNAP active site. This causes an inability to resume transcription, creating barriers to replication that threaten genome stability (4,5).

Mechanisms exist that reduce the probability of a replication fork encountering paused transcription complexes. Upstream RNAPs on the DNA and ribosomes on the emerging transcript may inhibit backtracking of a paused transcription complex in bacteria (5–7). RNA translocases such as the bacterial Rho hexamer can also translocate along transcripts not coated with ribosomes and displace paused and blocked RNAPs (5,8,9). Rho translocation also disrupts R-loops, structures in which an RNA transcript hybridizes to the DNA template strand (9). Nucleases can also aid genome duplication by promoting the resumption of transcription by backtracked RNAP (5,10). GreA and
GreB bind to *E. coli* RNAP and stimulate cleavage of backtracked transcripts, creating a new RNA 3′ end that can be used to restart transcription (11). Conversely, the 3′-5′ helicase UvrD can increase rather than decrease backtracking of RNAP and this may increase access of nucleotide excision repair enzymes UvrABC to DNA damage thus helping to clear barriers to RNAP and aid genome duplication (12).

Signalling molecules such as guanosine tetraphosphate and pentaphosphate in bacteria (referred to collectively as (p)ppGpp) can also reprogramme transcription. (p)ppGpp is central to the stringent response in bacteria, identified originally as a response to amino acid starvation in which the presence of uncharged tRNA within the ribosomal A site stimulates ribosome-bound RelA to synthesize (p)ppGpp (13–15). Control of the stringent response also requires (p)ppGpp-specific pyrophosphohydrolase, SpoT, that also has low (p)ppGpp synthase activity (15). However, the synthase activity of SpoT is induced by other stresses such as depletion of fatty acids (16). Central to the stringent response is binding of (p)ppGpp to RNAP which leads to downregulation of stable RNA production and upregulation of stress response genes (15,17,18). Expression of ribosomal rRNA operons accounts for half of all transcription during rapid growth when (p)ppGpp levels are low and thus provides the majority of transcriptional obstacles to replisomes in vivo when cells are dividing quickly (19–22). Elevated (p)ppGpp therefore reduces conflicts between replication and transcription by decreasing the density of transcribing RNAPs on the genome. (p)ppGpp can also decrease the stability of transcription complexes blocked by template damage in vitro which may decrease the density of blocked RNAP on the genome (10). This destabilization has been questioned, though, and a role for (p)ppGpp in increasing the fidelity of transcription and hence reducing RNAP pausing and backtracking has been proposed (23). Other work implicates (p)ppGpp acting synergistically with UvrD in promoting backtracking of paused RNAP, one function of which could be to facilitate transcription-coupled repair of any pause-inducing DNA damage (24).

Thus, how (p)ppGpp might act on transcription elongation complexes, as opposed to transcription initiation, is still far from clear. Transcription is not the only target of (p)ppGpp, though, since (p)ppGpp also binds multiple other targets. Inhibited targets include DnaG primase (25–28) and the translation elongation factors EFG and EF-Tu (29), indicating the pleiotropic impact of elevated (p)ppGpp on replication, transcription and elongation. (p)ppGpp also affects other diverse aspects of metabolism such as phospholipid synthesis, oxidative metabolism and resistance to antibiotics (30). These signalling molecules also enhance the ability of *E. coli* to persist in the presence of antibiotics. Persistence is the non-heritable ability of a small fraction of a bacterial population to survive exposure to an otherwise lethal concentration of antibiotic (31). (p)ppGpp levels vary stochastically in a bacterial population and increased (p)ppGpp activates toxin–antitoxin systems that induce slow growth in a sub-population of cells, leading to antibiotic tolerance (32).

Mechanisms also exist to increase the probability of continued replisome movement in the event of a collision with an RNAP. Accessory replicative motors provide a supply of additional helicases at the fork to aid protein displacement ahead of the replisome (33–37). In *E. coli* Rep helicase promotes movement of replisomes along protein-bound DNA in vitro and in vivo (20,34). Rep appears to be the main replisome anti-pausing factor in *E. coli* (38) and absence of Rep results in at least a two-fold increase in genome duplication time (39,40) and higher dependence on recombination-directed repair of damaged forks (41–44). However, Δrep cells remain viable since the homologous helicase UvrD can substitute for the absence of Rep at the replication fork (20,34). This substitution is only partial, though, and correlates with a physical and functional interaction between DnaB and Rep but not between DnaB and UvrD (34,45). This partial functional overlap is sufficient for single deletion mutants to be viable during rapid growth whereas Δrep ΔuvrD cells are not (46). Δrep ΔuvrD inviability can be suppressed by growth on minimal medium, conditions under which (p)ppGpp levels are high, and also by elevation of (p)ppGpp levels on rich medium via the spoT1 mutation encoding a (p)ppGpp pyrophosphohydrolase-defective SpoT (20,34). Mutations within rpo genes can also suppress Δrep ΔuvrD rich medium lethality (20,34,47). These mutant RNAPs display different phenotypes suggesting distinct mechanisms of suppression but some seem to pheno- copy elevated (p)ppGpp and/or reduce RNAP backtracking (10,47,48).

The above mechanisms reduce conflicts between replication and transcription but under rapid growth conditions forks are still blocked sufficiently frequently to require replisome reloading enzymes to maintain viability (49,50). Replisome reloading also often requires remodeling of the DNA at the blocked fork by recombination enzymes in order to generate a DNA structure suitable for replisome reloading (51). However, recombinational processing can lead to genome instability and is thus tightly controlled. One control is exerted by UvrD as it can disrupt RecA-ssDNA filaments and this disruption prevents excessive RecFOR-dependent loading of RecA onto ssDNA at blocked forks (52). Such excessive RecA loading contributes to Δrep ΔuvrD lethality, evinced by the weak suppression of Δrep ΔuvrD rich medium lethality via mutations in recF, recO or recR (34,53,54).

We have searched for suppressors of Δrep ΔuvrD rich medium lethality that are not within rpo, spoT, recF, O or R. We identified a spontaneous suppressor in a tRNA gene, *asp*T*18c*, that mutated a highly conserved residue that is structurally important in other tRNA species. Probing the basis of this suppression revealed that defective tRNA aminoacylation or insufficient peptide bond formation within the ribosome also suppressed Δrep ΔuvrD lethality. In all cases, suppression required RelA-directed (p)ppGpp synthesis indicating that stalling of ribosome translocation by uncharged tRNA in the A site of ribosomes underpinned suppression. Thus, although efficient translation elongation aids genome duplication (5,7), the partial inhibition of ribosome translocation is more effective at mitigating replication-transcription conflicts. These data illustrate the fine balance between the multiple mechanisms that promote simultaneous gene expression and genome duplication and reveal the importance of accessory replicative helicase activity in determining this balance.
MATERIALS AND METHODS

Plasmids and strains
pAM403 (55) and pAM407 (34) are pRC7 derivatives encoding rep and uvrD, respectively. p3LC-TL30-5P and p3LC-TL30-5D contain a cadC–lacZ gene fusion in which five tandem codons present within the linker encode proline or aspartate, respectively (56). Strains were constructed by P1 vir transduction and are listed in Supplementary Table S1.

Genome sequencing and analysis
DNA was extracted from 1 culture each of strain N7153 and N7182 grown in LB broth to stationary phase using Qiagen 100/G genomic tips from 5 ml cultures following the manufacturer’s protocol. Genome sequencing was performed using an Illumina GAIIx instrument with 100 bp paired end reads. Paired reads were trimmed to remove adapters and mapped against the E. coli K12 strain MG1655 NC_000913 genome using bwa (57), duplicates marked using Picard (http://broadinstitute.github.io/ picard) and variant analysis performed with SAMtools (58), followed by merging of variant tables using perl. Identified high quality synonymous and non-synonymous single nucleotide polymorphisms were annotated manually using the Integrative Genomics Viewer (59). The Illumina data were submitted in the form of fastq files to the European Nucleotide Archive (ENA) and are available under accession number PRJEB14483 at http://www.ebi.ac.uk/ena/data/view/PRJEB14483.

Growth assays
All steps in plasmid loss assays were carried out at 37°C except those shown in Figures 2B and 3B in which all steps were conducted at 30°C or 25°C as indicated. Colonies were grown in LB broth and agar except the assays in Figure 2B and Supplementary Figures S3B and S4 which were performed with a defined rich medium broth and agar containing 0.2% glycerol (Figure 2B) or 0.2% glucose (Supplementary Figures S3B and S4) as a carbon source (60). The plates used in Supplementary Figures S3B and S4 also had decreasing concentrations of aspartate or phenylalanine, as indicated. Strains carrying derivatives of pRC7 were grown in LB broth or defined rich medium with 100 μg ml⁻¹ of ampicillin overnight, diluted 10-fold into the same type of fresh liquid medium without ampicillin and grown to A600 0.4. Dilutions were then plated onto LB or defined rich medium agar plates containing 120 μg ml⁻¹ X-gal and 1 mM IPTG and incubated for 48 h except those shown in Figure 2B. Plates were then photographed and scored for blue/white colony formation.

For assays to assess colony forming ability, strains were grown in LB broth overnight at 37°C or, when temperature sensitive strains were involved, at 30°C. Serial 10-fold dilutions were made with 56/2 salts (61) on ice and then 5 μl of each dilution was spotted onto LB agar plates. Plates were then incubated at 37°C for 16 h unless otherwise stated. Ampicillin and bicyclomycin were included in LB agar plates where indicated at 100 and 25 μg ml⁻¹, respectively.

For the colony formation assays in Supplementary Figure S3A, strains were grown in defined rich medium broth (60) containing 0.2% glucose, all amino acids and 100 μg ml⁻¹ ampicillin overnight at 37°C. Tenfold serial dilutions were made in 56/2 salts and then 5 μl of each dilution spotted onto defined rich medium plates containing 0.2% glucose, 100 μg ml⁻¹ ampicillin and with all amino acids or missing either aspartate or phenylalanine. These plates were incubated at 37°C for 16 h. Minimal medium agar plates (61) were used in Supplementary Figure S3C without and with 100 μg ml⁻¹ ampicillin as indicated and incubated at 37°C for 72 h.

Colony-forming ability at increasing doses of UV light was assessed as described (62). Mismatch repair capacity was measured as the fraction of cells in a culture that acquired spontaneous mutations leading to rifampicin resistance. Briefly, overnight LB cultures were washed once in 56/2 salts and serially diluted 10-fold. 100 μl of the neat and the 10⁻¹ dilution were spread on LB agar containing 15 μg ml⁻¹ rifampicin. To estimate the total cfu ml⁻¹ in the overnight culture, 10 μl of the 10⁻⁶ dilution was spotted in triplicate on LB agar without rifampicin. Plates were then incubated at 37°C for 16 h and the colonies then counted.

The liquid growth assays in Figure 2A were performed using a Tecan Infinite M200 Pro plate reader. Overnight LB cultures were diluted to A400 0.005 in LB and 150 μl of each culture was transferred into 20 wells of a 96-well flat bottom plate (Corning). The microplate was incubated at 37°C for 24 h and A400 measurements were performed every 30 min preceded by plate shaking.

Persistence
Single colonies were inoculated into 5 ml LB broth and grown with shaking at 37°C overnight. 50 μl of overnight culture was then inoculated into 5 ml of LB broth and grown at 37°C with shaking until 1–2 × 10⁸ colony forming units ml⁻¹ reached. 500 μl was then transferred into two 1.5 ml microcentrifuge tubes. The first tube was centrifuged at 6000 rpm for 5 min at room temperature, resuspended in 500 μl 56/2 salts, serially diluted tenfold in 56/2 salts and 10 μl volumes of the 10⁻⁴ and 10⁻³ dilutions spotted onto LB agar plates containing 20 mM MgSO₄ in triplicate. To the second tube 5 μl of 10 μg ml⁻¹ ciprofloxacin was added and the tube was then inserted into a 50 ml Falcon tube and placed in a shaking incubator at 37°C for 5 h. After the 5 h incubation, this tube was centrifuged at 6000 rpm for 5 min at room temperature, resuspended in 500 μl 56/2 salts, recentrifuged and resuspended in 500 μl 56/2 salts. Colony-forming units were then assayed as for the first tube by serial dilution into 56/2 salts and plating in triplicate onto LB agar containing 20 mM MgSO₄. MgSO₄ was included to inhibit the activity of any traces of ciprofloxacin not removed by washing of the cells (63). All plates were incubated at 37°C overnight and then the fraction of colony forming units surviving ciprofloxacin exposure calculated.
Flow cytometry

Analysis of chromosome content after completion of all ongoing rounds of replication, so-called run-out conditions, was performed on mid-log phase cultures after treatment with rifampicin and cephalixin as described using a Becton Coulter CyAn ADP cytometer with 488 nm excitation and a 530/540 nm bandpass filter. Analysis of (p)ppGpp-dependent formation of RpoS-mCherry by flow cytometry was performed by growing the indicated strains overnight in LB, transferring 100 μl into 10 ml LB in a 125 ml flask followed by incubation at 37°C for 2h. Then 1 ml of the culture was centrifuged, resuspended in 2 ml of 10% LB in M9 medium and analysed by flow cytometry on a Becton Dickinson LSRFortessa X-20 flow cytometer using 561 nm laser excitation and detection using a 610/620 nm bandpass filter.

Translation assays

β-Galactosidase activity assays monitoring the relative levels of translation of cadC–lacZ fusions were performed in LB broth as described (56).

RESULTS

A tRNA mutation suppresses the requirement for accessory replicative helicases

pRC7 is a very low copy plasmid which encodes β lactamase and can therefore be maintained in E. coli cells by the inclusion of ampicillin in the medium (65). However, the inefficient origin of replication within pRC7 results in rapid loss of the plasmid in the absence of ampicillin. This rapid loss can be detected if the host strain has a chromosomal deletion of lacIZYA since pRC7 encodes the lac operon and hence cells with and without the plasmid appear blue and white, respectively, on plates containing IPTG and X-gal (65). This retention or loss of pRC7 can be used to assess whether a gene is essential for colony formation by cloning of the test gene into pRC7, transformation of a ΔlacIZYA strain with the pRC7 derivative and subsequent deletion of the test gene from the chromosome. Plating of the strain onto medium containing IPTG and X-gal results in formation of only blue colonies if the test gene is essential or white and segregating colonies if the test gene is not essential (65). Rapid growth of E. coli requires accessory helicase activity and so pRC7 encoding uvrD can be lost rapidly from ΔlacIZYA repΔuvrD cells on rich medium but pRC7 ΔuvrD cannot be lost from ΔlacIZYA Δrep ΔuvrD cells as monitored by blue/white colony colour (34) (see also Figure 1A, compare i and ii). Spontaneous mutations that suppress this requirement for an accessory helicase can be isolated by exploiting the ability of Δrep ΔuvrD cells to grow on minimal medium in the absence of a complementing pRC7 plasmid and subsequent plating of plasmid-less cells onto rich medium (34). Rare survivors on rich medium can then form colonies and the mutation(s) responsible for allowing Δrep ΔuvrD cells to grow under rapid growth conditions can be analysed. We identified one such suppressor, the strain designation of which is N7182 (Supplementary Table S1). Potential linkage of the suppressor mutation to rep was tested by transducing Δrep::cat from the suppressor strain into pRC7 ΔuvrD repΔ ΔuvrD::dfr (N6639). Eleven chloramphenicol-resistant transductants were tested for loss of pRC7 ΔuvrD on rich medium. Four transductants could not lose pRC7 ΔuvrD but seven could, indicating close linkage of the suppressor mutation with Δrep::cat (see also Figure 1A, compare ii and iii).

We sequenced the genome of this suppressor strain and compared it with the genome of N7153, a Δrep ΔuvrD strain that contains a well-characterized suppressor mutation rpoBΔ35 (5,10,34,66). Use of the Δrep ΔuvrD ΔrpoBΔ35 strain as a reference genome avoided the need for the reference strain to retain a complementing plasmid for viability. Only one mutation in N7182 that was not present in N7153 was located sufficiently close to Δrep::cat to explain the above linkage (see Supplementary File 1). This mutation resulted in replacement of T with C at position 8 within the aspT gene, one of three identical tRNA^Np genes in E. coli. Sequencing of the aspT gene from the 11 transductants obtained in the above cross revealed that all seven strains able to lose pRC7 ΔuvrD contained aspT[t8c] whereas all four that could not lose pRC7 ΔuvrD retained a wild type copy of aspT. Suppression of the viability of Δrep ΔuvrD cells on rich medium was therefore associated with the aspT[t8c] allele, with suppression comparable to that of ΔrpoBΔ35 in a plasmid loss assay (Figure 1A, compare iii and iv).

Δrep ΔuvrD^+ cells have a growth defect defects since UvrD can compensate only partially for the absence of Rep accessory helicase activity, resulting in slower movement of replication forks in Δrep cells (39,40). Thus, the median number of copies of oriC in Δrep ΔuvrD^+ cells is twice that of repΔ ΔuvrD^+ cells in rich medium due to an extended cell cycle in Δrep cells and hence more replication initiation events per cell cycle (38). This doubling in oriC numbers results in a doubling of chromosome content when cells are treated with cephalixin and rifampicin to inhibit cell division and reinitiation of replication, so-called run-out conditions (38) (see also Figure 1B, compare i and iii). ΔrpoBΔ35 suppressed the increased chromosome copy number in Δrep ΔuvrD^+ cells, reducing the median number of chromosomes from eight to four (Figure 1B, compare iii and iv). These data support the conclusion that this tRNA mutation reduces the need for accessory helicase activity. Furthermore, ΔaspT[t8c] had no detectable impact on other UvrD-mediated processes. Defects in nucleotide excision repair, mismatch repair and control of recombination in repΔ ΔuvrD cells (52,67,68) were not suppressed by ΔaspT[t8c] (Supplementary Figure S1A, B and C, respectively), providing further support for the specific suppression of the accessory helicase defect in Δrep ΔuvrD cells by ΔaspT[t8c].

The requirement for accessory helicase activity is driven primarily by transcriptional barriers to replication (20,34). Backtracking of paused RNA polymerases results in formation of particularly stable replicative barriers and multiple factors have evolved to reduce the numbers of backtracked complexes (5). Cells lacking two homologous anti-backtracking factors, GreA and GreB, display a temperature-sensitive growth defect that is a consequence of more frequent collisions between replisomes and backtracked transcription complexes (5,10,69). ΔaspT[t8c] suppressed the temperature-sensitive phenotype of ΔgreA
A mutation in an aspartyl tRNA gene suppresses the need for accessory replicative helicases and for anti-backtracking factors. (A) Retention or loss of pRC7uvrD (pAM407) from strains without or with Δrep ΔuvrD deletions as judged by blue/white colony color on LB plates containing X-gal and IPTG. Fractions of white colonies are indicated below each image with actual numbers of white versus total colonies counted in parentheses. (B) DNA content of the indicated strains as monitored by flow cytometry under run out conditions in LB. The number of chromosome equivalents per cell is shown below. (C) The viability of greA+ greB+ versus ΔgreA ΔgreB cells without and with aspT[t8c] as monitored by serial dilutions of liquid cultures grown at 30°C and plated subsequently on LB agar at 30°C and 42°C.

ΔgreB cells (Figure 1C), similar to the suppression seen with rpoB*35 (48). We conclude that aspT[t8c] reduces the need for GreA/GreB-dependent rescue of backtracked RNA polymerase.

Taken together, these data indicate that aspT[t8c] reduces the need for Rep and UvrD to underpin replication and for anti-backtracking factors to resuscitate transcription complexes.

Suppression by aspT[t8c] does not occur via growth rate restriction

The uridine encoded at position 8 within the wild type aspT gene is post-transcriptionally modified to 4-thiouridine and this modified nucleotide is conserved across all kingdoms of life (70). This residue is involved in a triple non-Watson–Crick pairing interaction and is important in coordination of magnesium within tRNA (71,72). This central structural role is reflected in the temperature-dependent destabilisation of human mitochondrial tRNA\textsubscript{Met} structure by the same T to C mutation as found in aspT[t8c]. This transition mutation in tRNA\textsubscript{Met} inhibits aminoacylation and any mutated tRNA\textsubscript{Met} that is aminoacylated fails to form a stable ternary complex with elongation factor EF-Tu (70).

Given the conservation of tRNA structure, aspT[t8c] may result in similar structural destabilisation of the encoded tRNA\textsuperscript{Asp} and consequent inhibition of interactions with aspartyl tRNA synthetase and EF-Tu. However, aspT[t8c] did not have a major impact on growth of rep\textsuperscript{+} uvrD\textsuperscript{+} cells in liquid culture (Figure 2A, compare i and ii). This absence of a significant growth defect in aspT[t8c] strains may be due to the presence of two other identical tRNA\textsuperscript{Asp} genes in E. coli, aspU and aspV.

Other means of growth restriction did not suppress Δrep ΔuvrD inviability. Growth on defined rich medium containing all 20 amino acids but with a poor carbon source, glyc-
tRNAAsp aminoacylation was therefore tested for suppression of Δrep ΔuvrD lethality at 30°C on low salt medium, conditions under which viability is similar to wild type (73) (see also Figure 3A, compare i and ii). pRC7uvrD could be lost from Δrep ΔuvrD aspSP555S at 30°C on low salt medium but not on high salt medium (Figure 3Bi). Suppression of Δrep ΔuvrD lethality correlates therefore with a defect in aspartyl tRNA synthetase.

We also tested whether an alternative means of inhibiting tRNA<sup>Asp</sup> aminoacylation could suppress Δrep ΔuvrD inviability. A pRC7uvrD/Δrep ΔuvrD strain defective in aspartate biosynthesis could lose the complementing pRC7uvrD plasmid on defined rich medium upon restriction of aspartate availability (Supplementary Figure S3). Thus a defect in the enzyme needed to synthetize aspartyl tRNA, AspRS, or a limiting concentration of one of the substrates needed for formation of aspartyl tRNA, aspartate, can suppress the need for accessory replicative helicase activity. Moreover, suppression was not specific to aspartate starvation as restriction of availability of phenylalanine also allowed Δrep ΔuvrD cells to survive in the absence of a complementing plasmid (Supplementary Figure S4). As expected, restricting amino acid availability also resulted in very poor growth (Supplementary Figures S3 and S4) but a restricted growth rate does not by itself provide suppression of Δrep ΔuvrD lethality (Figure 2B). These data support the conclusion that inhibition of aminoacylation of tRNA<sup>Asp</sup>, and of other tRNAs, reduces the need for accessory helicase activity.

Translational pausing suppresses Δrep ΔuvrD inviability

Suppression of Δrep ΔuvrD inviability via inhibition of tRNA<sup>Asp</sup> or tRNA<sup>Phe</sup> aminoacylation might occur directly via pausing of ribosomes but could also occur via decreased production of one or more specific proteins containing both aspartate and phenylalanine. We tested therefore whether increased translational pausing by a well-defined mechanism that does not rely on decreased tRNA aminoacylation also suppresses the need for accessory helicase activity. Peptide bond formation by ribosomes occurs with low efficiency within polyproline tracts as compared with other amino acids and this low efficiency is compensated for by interaction of elongation factor P (EF-P) with the ribosome (56,75). When EF-P is absent (Δefp) ribosomes pause for extended periods at tandem proline codons (56,75) (see also Supplementary Figure S2) resulting in reduced growth rates (76) (see also Figure 2A). Despite its negative effect on growth rate, introduction of Δefp clearly suppressed Δrep ΔuvrD lethality (Figure 4, compare A and B). yjeA and yjeK encode enzymes needed for post-translational modification of EF-P to form functional enzyme (56,75–78) and deletion of either yjeA or yjeK also resulted in suppression (Figure 4C and D). Enhancement of ribosomal pausing at polyproline sequences can therefore reduce the need for accessory helicase activity.

Figure 2. The impact of aspT<sup>18c</sup> on growth and its relevance to Δrep ΔuvrD inviability. (A) Growth of (i) asp<sup>T</sup> <sup>+</sup> efp+ (TB28), (ii) asp<sup>T</sup> <sup>18c</sup> efp+ (KM231) and (iii) asp<sup>T</sup> <sup>+</sup> efp (MH299) in LB at 37°C as monitored by absorbance at 600 nm. (B) Assessment of the ability of pRC7uvrD (pAM407) to be lost from Δrep ΔuvrD cells by reducing growth rates via culturing at 37°C and 25°C for the indicated times on defined rich medium containing glycerol as a carbon source.

Defective tRNA aminoacylation suppresses Δrep ΔuvrD inviability

As reduced growth rate was not the cause of suppression by aspT<sup>18c</sup>, we probed whether defective translation was responsible. We could not detect significant defects in translation in vivo at tandem aspartate codons in an aspT<sup>18c</sup> strain (Supplementary Figure S2). Either this allele does not result in translational pausing or such pausing is below the limits of detection using this assay. We therefore used alternative approaches to probe the aspT<sup>18c</sup> suppression mechanism. The same 18c mutation in human mitochondrial tRNA<sup>Met</sup> inhibits aminoacylation (70). Inhibition of tRNA<sup>Asp</sup> aminoacylation was therefore tested for suppression of Δrep ΔuvrD lethality. The E. coli tls-1 allele encodes a P555S mutation in aspartyl tRNA synthetase that reduces the thermal stability of the synthetase and causes a severe reduction in growth rate at 42°C under low salt conditions (73,74) (see also Figure 3A). Given that AspRS<sup>P555S</sup> is less stable than wild type enzyme regardless of the temperature (74), we tested whether asp SP555S suppressed Δrep ΔuvrD lethality at 30°C on low salt medium, conditions under which viability is similar to wild type (73) (see also Figure 3A, compare i and ii). pRC7uvrD could be lost from Δrep ΔuvrD aspSP555S at 30°C on low salt medium but not on high salt medium (Figure 3Bi). Suppression of Δrep ΔuvrD lethality correlates therefore with a defect in aspartyl tRNA synthetase.

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Suppression by aspT<sup>18c</sup> and Δefp requires (p)ppGpp synthesis

One consequence of ribosomal pausing is increased synthesis of (p)ppGpp by RelA upon binding of RelA to a ribo-
Figure 3. A mutation in aspartyl tRNA synthetase suppresses ∆rep ∆uvrD lethality. (A) The indicated strains were grown overnight in high salt medium at 30°C and then serial dilutions plated onto low and high salt plates containing 100 µg mL⁻¹ ampicillin and incubated at either 30°C or 42°C. (B) Suppression of ∆rep ∆uvrD lethality by aspS¹⁸⁵⁵⁵ on rich medium was analysed by monitoring retention or loss of pRC7uvrD(pAM407) from the indicated strains at 30°C on either low or high salt LB medium.

some containing a non-acylated tRNA in the A site (13,14). Elevated (p)ppGpp is known to suppress ∆rep ∆uvrD inviability (34) and so aspT[t8c] and ∆efp might both therefore suppress via elevation of (p)ppGpp concentration.

Direct measurement of (p)ppGpp using 3²-P cannot be performed on cells grown in rich media (79) which prevents direct assessment of (p)ppGpp levels in aspT[t8c] and ∆efp strains under conditions relevant to suppression of ∆rep ∆uvrD lethality. An indirect assay was therefore used that employs an RpoS–mCherry translational fusion (32). This reporter provides a fluorescence signal that correlates with intracellular (p)ppGpp concentration due to (p)ppGpp both stimulating rpoS transcription and inhibiting RpoS degradation (32). Fluorescence was assayed by flow cytometry of cells grown to mid-logarithmic phase in rich medium. relA⁺ spoT⁺ cells lacking the RpoS–mCherry fusion and rpoS–mCherry ΔrelA ΔspoT cells which are unable to synthesize (p)ppGpp gave similar levels of background fluorescence (Figure 5Ai and ii). An increase in fluorescence was observed in wild type cells carrying the fusion but this increase was modest (Figure 5A, compare iii with i and ii), consonant with low (p)ppGpp levels in wild type cells growing in nutrient-rich environments (32,80). mCherry fluorescence increased substantially in ∆efp cells but not in aspT[t8c] cells with respect to wild type (Figure 5A, compare iv and v with iii). We conclude that absence of EF-P results in elevation of (p)ppGpp concentration. Higher (p)ppGpp concentration in ∆efp cells as compared with wild type or aspT[t8c] cells is consistent with the significant retardation of growth of ∆efp cells (Figure 2A). In contrast, aspT[t8c] either does not increase (p)ppGpp levels as compared with wild type cells or any increase is
Figure 5. (p)ppGpp synthesis is needed for suppression of Δrep ΔuvrD lethality by aspT[t8c] and Δefp. (A) Flow cytometric detection of in vivo levels of RpoS-mCherry fluorescence in the indicated strains. All strains contain the rpoS-mCherry fusion with the exception of (i) MG1655. (B) Survival after challenge with ciprofloxacin. The strains are (i) TB28, (ii) KM241, (iii) N5777, (iv) KM231 and (v) MH372. (C) Retention and loss of pRC7uuvD (pAM407) on LB X-gal IPTG agar in relA+ and ΔrelA strains.

below the limits of detection using this assay. To gauge the sensitivity of this assay we monitored fluorescence in spoT1 cells. spoT1 encodes a SpoT enzyme that retains (p)ppGpp synthase activity but lacks (p)ppGpp pyrophosphorylase activity, resulting in elevated (p)ppGpp (81) and the ability to suppress Δrep ΔuvrD lethality (34). spoT1 rpoS-mCherry cells did not result in increased fluorescence as compared with spoT+ rpoS-mCherry cells indicating that (p)ppGpp levels sufficient to reduce the need for accessory replicative helicases could go undetected using this assay (Figure 5Aiii and vi).

To address this potential sensitivity problem we employed a second assay to determine whether either Δefp or aspT[t8c] results in elevated (p)ppGpp. Cells with elevated (p)ppGpp display elevated levels of persistence and thus spoT1 enhances whereas ΔrelA ΔspoT alleles reduce persistence (32,82) (see also Figure 5Bi–iii). aspT[t8c] and Δefp both increased persistence with the increase being higher for Δefp (Figure 5Biv and v). The large increase in persistence in Δefp cells correlates with the enhancement of fluorescence in Δefp rpoS-mCherry cells. The level of persistence in aspT[t8c] cells is consistent with a smaller increase in
(pppGpp levels as compared with 3Δefp) that is below the limits of detection in the mCherry fluorescence assay.

We tested whether increased (pppGpp) contributed to suppression of 3Δrep ΔuvrD inviability by aspT[18c] and 3Δefp by deleting the gene encoding the primary (pppGpp) synthase, relA. Deletion of relA prevented loss of pRC7uvrD from 3Δrep ΔuvrD aspT[18c] and 3Δrep ΔuvrD 3Δefp cells, indicating that RelA was required for suppression by both mutant alleles (Figure 5C, compare iii with iv and v with vi). In contrast, pRC7uvrD could be lost from 3Δrep ΔuvrD 3ΔrelA rpoB*35 cells indicating that RelA was not required for the viability of suppressed 3Δrep ΔuvrD strains under all circumstances (Figure 5C, compare vii and viii). This lack of dependence of 3Δrep ΔuvrD rpoB*35 on RelA likely reflects the recapitulation by rpoB*35 of many phenotypes associated with elevated (pppGpp) even in the absence of RelA (48,66).

Taken together, these data indicate that RelA-directed synthesis of (pppGpp) is important for suppression of 3Δrep ΔuvrD lethality by both aspT[18c] and 3Δefp.

### aspT[18c] and 3Δefp confer differing requirements for Rho activity

Translocation 5'–3' by Rho along untranslated and unstructured nascent transcripts can displace transcription complexes thus reducing both RNA polymerase occupancy on the chromosome and R-loop formation (9,83,84). Rep and Rho therefore provide two different mechanisms that reduce the impact of transcription on replication. Consequently, cells need either Rep or wild type levels of Rho activity to maintain genome duplication in the face of transcriptional barriers (8,85). One manifestation of this requirement is the hypersensitivity of 3Δrep ΔuvrD cells to low concentrations of the Rho-specific inhibitor bicyclomycin (8) (see also Figure 6A and B, compare i and iii).

We assessed whether 3Δefp or aspT[18c] could suppress 3Δrep bicyclomycin hypersensitivity. Neither 3Δefp nor aspT[18c] altered bicyclomycin sensitivity in a rep+ background (Figure 6A and B, compare i and ii). In 3Δrep cells hypersensitivity was suppressed by 3Δefp but not by aspT[18c] (Figure 6A and B, compare iii and iv). These data demonstrate that 3Δefp but not aspT[18c] can reduce the requirement for Rho activity in the absence of Rep.

We also tested whether this differential need for Rho activity was reflected in cells lacking both rep and uvrD by exploiting rho [A243E] which encodes a partial loss of function enzyme (86). Rho dependency was assessed by the ability of 3Δrep ΔuvrD strains to lose a complementing pRC7rep rather than pRC7uvrD plasmid, allowing construction of pRC7rep/3Δrep ΔuvrD rho [A243E] strains regardless of reprho double mutant lethality (85). The similar numbers and sizes of white plasmidless colonies formed by 3Δrep ΔuvrD 3Δefp rho’ and 3Δrep ΔuvrD 3Δefp rho [A243E] cells indicate that wild type Rho function was not required for suppression by 3Δefp (Figure 6C, compare v and vi). In contrast, 3Δrep ΔuvrD aspT[18c] rho [A243E] gave much smaller white plasmidless colonies as compared with the isogenic rho’ strain indicating that wild type Rho function was important for growth of aspT[18c] 3Δrep ΔuvrD cells (Figure 6C, compare iii and iv).

The data in Figures 5 and 6 indicate that the viability of aspT[18c] 3Δrep ΔuvrD cells requires both the major (pppGpp) synthase in the cell, RelA, and wild type Rho activity. In contrast, 3Δefp 3Δrep ΔuvrD cells require RelA but not wild type Rho activity. 3Δefp cells have higher levels of (pppGpp) as compared with aspT[18c] (Figure 5A) and these elevated levels might explain the differential requirement for Rho, given the ability of (pppGpp) to reduce replication/transcription conflicts (10,34,66). Such a model implies that (pppGpp) synthesis is critical not only for the viability of 3Δefp 3Δrep ΔuvrD cells (Figure 5C) but also for suppression of bicyclomycin sensitivity of 3Δrep cells (Figure 6A). Absence of the primary (pppGpp) synthase RelA did not hypersensitize otherwise wild type cells to bicyclomycin (Figure 7Aii). However, the suppression of 3Δrep bicyclomycin hypersensitivity by 3Δefp was abolished upon deletion of relA (Figure 7A, compare vii with vii). Thus RelA-dependent (pppGpp) synthesis in 3Δrep 3Δefp cells is essential for survival with lowered Rho activity, supporting the hypothesis that elevated (pppGpp) can reduce the need for Rho.

We tested this possible link between (pppGpp) concentration and requirement for Rho by analysing the impact of other means of increasing (pppGpp) concentration on 3Δrep bicyclomycin hypersensitivity. The spoT1 allele suppressed 3Δrep bicyclomycin hypersensitivity supporting the conclusion that elevated (pppGpp) can reduce the need for Rho (Figure 7B, compare iv with ii).

The data in Figures 5–7 indicate that both aspT[18c] and 3Δefp require RelA-directed synthesis of (pppGpp) to maintain the viability of 3Δrep ΔuvrD cells. In contrast, 3Δefp has a much lower dependence on Rho activity as compared with aspT[18c]. This differential requirement for Rho activity correlates with higher (pppGpp) levels in 3Δefp cells as compared with aspT[18c]. Resolving conflicts between replication and transcription involves therefore a fine balance between accessory replicative helicases, levels of (pppGpp) and Rho activity.

### DISCUSSION

We have discovered three new types of suppressors of 3Δrep ΔuvrD rich medium lethality: mutations in tRNA genes, in aminoacyl-tRNA synthetases and in translation elongation factors. Whilst the mechanistic consequences of the aspT[18c] mutation are unclear, the inhibition of tRNA aminoacylation by aspSP5555 and of translation elongation by 3Δefp are well-characterized (56,74,75), indicating that partial inhibition of translation can suppress 3Δrep ΔuvrD lethality. Previous studies have demonstrated that this lethality is caused primarily by the conflict between replication and transcription (20,34,54). Thus partial inhibition of translation can compensate for the impact of transcription on DNA replication in the absence of Rep and UvrD. Suppression requires RelA-directed synthesis of (pppGpp) with (pppGpp) concentration being higher with 3Δefp than with aspT[18c] (Figure 5). Wild type levels of Rho activity are also needed in aspT[18c] 3Δrep ΔuvrD but not 3Δefp 3Δrep ΔuvrD cells indicating that the relative importance of (pppGpp) and Rho function depends upon the nature of the translational mutation (Figure 6).
Figure 6. Δefp but not aspT[t8c] can bypass the need for wild type Rho activity. (A) and (B) The indicated strains were grown in liquid culture in the absence of bicymycin and their ability to continue to divide with reduced Rho activity was assessed after serial dilution onto plates without and with 25 μg ml⁻¹ bicymycin. (C) Loss of pRC7rep (pAM403) on LB X-gal IPTG agar in rho⁺ (i, iii, v) and rho[A243E] (ii, iv, vi) strains.

Δrep ΔuvrD cells are inviable on rich medium primarily because of lack of accessory replicative helicase function (20,34,54). Suppression of this growth defect by the above translation mutations (Figures 1A, 3B and 4) therefore suggest that it is the lack of accessory helicase activity in Δrep ΔuvrD cells that is being suppressed. Suppression by aspT[t8c] of the chromosome content defect of Δrep cells, a direct consequence of slower fork movement in the absence of Rep accessory helicase activity (34,38,39), supports this view (Figure 1B). UvrD also inhibits RecFOR-dependent loading of RecA at blocked replication forks (52,68) and absence of this function makes a minor contribution to Δrep ΔuvrD lethality. aspT[t8c] also does not suppress the sensitivity of ΔuvrD cells to UV light indicating that aspT[t8c] does not suppress nucleotide excision repair defects (Supplementary Figure S1A). Furthermore, aspT[t8c] suppression of the ΔgreA ΔgreB growth defect indicates that aspT[t8c] can suppress the absence of an anti-backtracking activity, making it difficult to explain how aspT[t8c] could also suppress the absence of a UvrD pro-backtracking function (Figure 1C).
RelA-directed synthesis of (p)ppGpp plays a central role in maintaining Δrep ΔuvrD viability by aspT[8c] and by Δefp (Figure 5C). (p)ppGpp dramatically inhibits transcription of rrr operons (17) which are the primary transcriptional barriers to genome duplication (19,22), together with other highly expressed operons (18). (p)ppGpp may also destabilize stalled RNAP (10) or increase the fidelity of transcription (23), both of which could decrease the impact of transcription on fork movement. aspT[8c]- and Δefp-directed suppression of Δrep ΔuvrD lethality via (p)ppGpp is therefore likely to occur by impacting on transcription initiation, stalled RNAP stability and/or decreased pausing due to enhanced fidelity.

RelA is stimulated to synthesize (p)ppGpp when an uncharged cognate tRNA is bound to the ribosomal A site (13,14). aspT[8c] and Δefp mutations may therefore increase the probability of A site-bound uncharged tRNA either directly or indirectly. The uridine at position 8 is highly conserved in tRNA species and the equivalent 8c mutation within human mitochondrial tRNAMet results in inhibition of tRNA aminoacylation (70). If aspT[8c] also resulted in inhibition of aminoacylation then the resulting increase in non-acylated tRNA8c could lead to an increased probability of uncharged tRNA8c occupying the ribosomal A site. However, whether the mutant tRNA8c can still bind to the A site is unknown. Alternatively tRNA8c might poison the aspartyl tRNA synthetase by binding to it and forming a dead end complex with respect to aminoacylation. Formation of such a dead end complex might titrate out aspartyl tRNA synthetase and increase levels of uncharged wild type tRNA8c leading to increased non-acylated tRNA8c occupying the ribosomal A site. For cells lacking EF-P, YjeA or YjeK it is difficult to conceive how a reduced rate of proline–proline bond formation within the ribosome could lead directly to increased occupancy of the A site by uncharged tRNA. Absence of any one of these three factors, though, does impact on expression of many genes and so altered expression of one or more genes in Δefp, ΔyjeA or ΔyjeK cells could result in an increased probability of A site-bound uncharged tRNA. For example, Δefp cells have reduced expression of valyl tRNA synthetase (87) which could lead to accumulation of uncharged tRNAVal and triggering of the stringent response. Suppression of Δrep ΔuvrD lethality by aspT8c demonstrates that suppression via partial loss of tRNA synthetase function can occur (Figure 3).

In contrast to the requirement for RelA to sustain suppression of Δrep ΔuvrD lethality by both aspT[8c] and Δefp, the requirement for Rho activity is reduced in Δefp as compared with aspT[8c] cells (Figures 6 and 7). This difference correlates with the higher concentration of (p)ppGpp in Δefp as compared with aspT[8c] cells (Figure 5). A role for (p)ppGpp in reducing the need for Rho activity is supported by the ability of spoT7 to suppress Δrep bicyclicin hypersensitivity (Figure 7B). Suppression of Δrep bicyclicin hypersensitivity by Δefp also depends on RelA which supports a critical balance between (p)ppGpp concentration and Rho activity in maintaining viability (Figure 7A). Elevated (p)ppGpp can therefore reduce the need for Rho.

The greater dependence of aspT[8c]-directed suppression on Rho might reflect not just lower (p)ppGpp levels in aspT[8c] versus Δefp cells but possibly also enhancement of Rho binding on nascent transcripts by aspT[8c]. In other words, suppression via aspT[8c] might occur at least partly via the increased generation of Rho binding sites on emerging transcripts. Aspartate codons are more frequent than proline tracts and hence aspT[8c] has the potential to impact on ribosome translocation more frequently than Δefp. Enhancement of Rho binding would also be dependent only on ribosomal pausing and not specifically require ribosomal A site occupancy by a non-ribosomal tRNA (88), in contrast to stimulation of RelA activity (13,14). However, any aspT[8c]-directed increase of transcription complex displacement by Rho cannot be sufficient by itself to suppress Δrep ΔuvrD lethality since RelA is also needed (Figure 5C).

Our data indicate that whilst translation helps prevent RNAP backtrackng (5,7) the reduction of conflicts between replication and transcription can be achieved more effectively by partial inhibition of translation. Of course, the growth defect in Δefp cells (Figure 2A) indicates why the balance between replication, transcription, translation and (p)ppGpp synthesis is poised as it is in wild type cells. Thus for the maintenance of rapid growth there is a very fine balance to be struck between gene expression and accurate, rapid genome duplication. Furthermore, accessory replicative helicases play a major role in determining this balance. In the absence of both Rep and UvrD the wild type balance between various other mechanisms that reduce the impact of transcription on replication is unable to effectively counter the adverse effects of transcription on completion of genome duplication. There may also be circumstances under which mutations such as those identified here confer a selective advantage even in rep+ uvrD+ cells. Elevated (p)ppGpp is a key factor in determining bacterial persistence in the face of antibiotic challenge (32) and both aspT[8c] and Δefp mutations enhance persistence (Figure 5B). It remains possible therefore that mutations such as aspT[8c] that have only a modest inhibitory effect on
growth could arise in bacterial populations continually exposed to antibiotics, especially given the many mutations known to affect translation (89). Increased persistence in strains lacking EF-P function also imply that EF-P and its unique post-translational modification pathway are poor potential targets for antibiotics.

ACCESSION NUMBERS

Sequencing data were submitted in form of fastq files to the European Nucleotide Archive (ENA) and are available under accession number PRJEB14483 at http://www.ebi.ac.uk/ena/data/view/PRJEB14483.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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REFERENCES


