Novel *Escherichia coli* RF1 mutants with decreased translation termination activity and increased sensitivity to the cytotoxic effect of the bacterial toxins Kid and RelE

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Summary

Novel mutations in *prfA*, the gene for the polypeptide release factor RF1 of *Escherichia coli*, were isolated using a positive genetic screen based on the parD (*kis, kid*) toxin–antitoxin system. This original approach allowed the direct selection of mutants with altered translational termination efficiency at UAG codons. The isolated *prfA* mutants displayed a ~10-fold decrease in UAG termination efficiency with no significant changes in RF1 stability *in vivo*. All three mutations, G121S, G301S and R303H, were situated close to the nonsense codon recognition site in RF1:ribosome complexes. The *prfA* mutants displayed increased sensitivity to the RelE toxin encoded by the *relBE* system of *E. coli*, thus providing *in vivo* support for the functional interaction between RF1 and RelE. The *prfA* mutants also showed increased sensitivity to the Kid toxin. Since this toxin can cleave RNA in a ribosome-independent manner, this result was not anticipated and provided first evidence for the involvement of RF1 in the pathway of Kid toxicity. The sensitivity of the *prfA* mutants to RelE and Kid was restored to normal levels upon overproduction of the wild-type RF1 protein. We discuss these results and their utility for the design of novel antibacterial strategies in the light of the recently reported structure of ribosome-bound RF1.

Introduction

Translation termination is an important stage in protein synthesis which leads to the release of the newly synthesized polypeptide chain from the ribosome. This is initiated when a termination codon in mRNA enters the ribosomal A site and is recognized by a class 1 release factor (RF). In bacteria, translation termination requires the active contribution of two protein release factors, RF1 and RF2 (Scolnick *et al*., 1968); these factors recognize, respectively, and with different affinity UAA/UAG or UAA/UGA termination codons at the decoding centre of the ribosomal A site. A conserved tripeptide motif GGQ in RF1 or RF2 interacts with the peptidyl transfer centre (PTC) (Frolova *et al*., 1999) and activates the hydrolysis of the ester bond between the polypeptide chain and the tRNA at the PTC, thereby releasing the newly synthesized polypeptide chain.

Specific recognition of termination codons by RF1 and RF2 involves specific tripeptide motifs, PxT for RF1 or SPF for RF2 (Ito *et al*., 2000), which are present in a central domain of the proteins. The tripeptide motifs were first suggested to be responsible for recognizing the last two nucleotides of the three stop codons (Ito *et al*., 2000), but recent structural studies indicate that they interact with the first two nucleotides (Laurberg *et al*., 2008) and explain in part the remarkable accuracy with which the first stop codon nucleotide, an invariant uridine, is recognized (Freistroffer *et al*., 2000). Discrimination against purines in the first stop codon position implicates residues near the tip of a helical finger (α-helix 5) present in the central domain (Laurberg *et al*., 2008). Two residues near the PxT motif, Q185 and T198 in *Escherichia coli* RF1, discriminate against pyrimidines in the third stop codon position (Laurberg *et al*., 2008). Interactions of RF1 or RF2 with a third factor, RF3-GTP or GDP, at the GTP-associated centre of the 50S ribosomal subunit, are also required for the final detachment of the class 1 RFs from the ribosome (Freistroffer *et al*., 1997; Zavialov *et al*., 2001; 2002; Gao *et al*., 2007).
A more detailed understanding of the role of the GGQ motif conserved in class I RFs has been provided by crystal structures of RF1 and RF2 in complex with the ribosome (Petry et al., 2005; Laurberg et al., 2008), and by molecular dynamics simulations of peptide release (Trobro and Aqvist, 2007). The GGQ motif, together with A76 of the P-site tRNA, is instrumental in co-ordinating the water molecule that initiates a nucleophilic attack on the peptidyl-tRNA ester bond. Following the detachment of the RF from the ribosome the subsequent action of the ribosomal releasing factor RRF and the elongation factor EF-G (Karimi et al., 1999; Gao et al., 2007) leads to the de-assembling of the ribosomal subunits, which can then be recycled and reassembled in a new translation initiation event.

Pausing of a bacterial ribosome due to limitations in tRNA can lead to cleavage of mRNA at a site within or immediately adjacent to the A-site codon; the ribosome participates in this cleavage in an, as yet, undefined way (Hayes and Sauer, 2003; Sunohara et al., 2004). Breaks at UAG and UGA stop codons induced by ribosome stalling due to the reduced activity of RF1 have been reported (Li et al., 2007). Breaks in the mRNA at UAG termination codons at the A site can also be induced by the RelE toxin encoded by the bacterial toxin–antitoxin (TA) system relBE. The cleavage occurs preferentially between the second and third nucleotides of UAG termination codons in the ribosomal A site implying the concerted action of the RelE toxin and the ribosome (Pedersen et al., 2003). The RelE toxin is activated under amino acid or carbon source limitation by a mechanism which involves the enhanced decay of the RelB antitoxin protein (Gerdes et al., 2005). Consistent with this, RF1, which preferentially recognizes this termination codon at the A site, can protect against the RelE cleavage in vitro (Pedersen et al., 2003). Cleavage of mRNA in all the above cases leads to the stalling of the ribosome at the end of the non-stop mRNA, and can eventually inhibit protein synthesis as in the case with RelE. Stalled ribosomes are released by the intervention of transfer-messenger RNA (tmRNA), which provides a new stop codon and tags potentially harmful truncated peptides for rapid degradation (Moore and Sauer, 2007); this, in turn, feeds amino acid pools [see Discussions by Gerdes et al. (2005) and by Condon (2006)].

The kis–kid antitoxin–toxin system of plasmid R1 codes for an unstable antitoxin Kis and a stable toxin Kid. Kid, like RelE, is also a specific endoribonuclease and an inhibitor of translation, but unlike RelE, Kid can cleave RNA in the absence of ribosomes and does so at the core sequence UA(A/C) preferentially but not exclusively in single-stranded regions (Zhang et al., 2004; Munoz-Gomez et al., 2005). Cleavage occurs by a mechanism similar to the RNase A and RNase T1, involving the Uracyl 2′-OH group and producing two fragments, with a 2′:3′-cyclic phosphate group at the uracil and a free 5′-OH group at the adenosine respectively. Cleavage of RNA in the absence of ribosomes can also introduce non-stop mRNA in the translation machinery, a proposal that is consistent with the increased sensitivity to the Kid toxin observed in tmRNA-deficient strains (Munoz-Gomez, 2004).

While searching for E. coli mutations affecting the toxic activity of the Kid protein, we developed a genetic screen that was based on the selection of bacteria that could survive in the presence of a mutant parD locus encoding a functional Kid toxin and a truncated and inactive antitoxin resulting from an amber termination mutation in the kis antitoxin gene (Bravo et al., 1988). As a consequence, the isolation of suppressors leading to the efficient bypass of the amber mutation in kis was one anticipated response. Unexpectedly, however, these mutations were found to be novel and to be exclusively targeted to the prfA gene encoding the RF1 protein. Our work reported here shows that these mutations reveal novel functional connections between the activity of bacterial toxins and translation termination factors.

**Results**

*Isolation of novel prfA mutants in E. coli*

Our work was initially focused on a search for E. coli mutants resistant to the cytotoxic activity of Kid. For this purpose, we used a double selection approach inspired by the work by Bernard and Couturier (1992), which led to the isolation of mutations in the gyrA gene encoding the A subunit of DNA gyrase, the bacterial target of the CcdB toxin. Two compatible plasmids were employed, pKK1120 and pAB1120, that confer resistance to tetracycline and kanamycin, respectively, and which both contain the mutant parD operon kis74 (amber), kid+ (Table 1). In the absence of suppression of the kis74 mutation, an inactive form of the Kis antitoxin is produced, thus leading to a Kid-dependent inhibition of colony formation. In a first round of selection, E. coli cells treated with the mutagen 2-aminopurine were transformed with the pKK1120 plasmid and cultured in medium containing tetracycline. To enrich for potential Kid-resistant mutants, a second round of selection was carried out by transforming the cells with the pAB1120 plasmid and plating on medium supplemented with kanamycin. To discard possible mutations affecting either the kid gene itself or its regulatory elements present in the plasmid, the toxicity of the resident plasmids was assessed by extracting the plasmid DNA and retransforming fresh wild-type E. coli cells. Mutants that exhibited growth despite containing a toxic plasmid were kept for further characterization.

The mutations were mapped to the region between 22 and 32 min on the E. coli chromosome using a collection
Table 1. Plasmids used and constructed in this work.

| Plasmid          | Relevant features                                      | Reference                      |
|------------------|-------------------------------------------------------|--------------------------------|
| pAB1120          | R1, copB, parD [kid74, kid[wt]], kan                 | Bravo et al.                   |
| pkK1120          | pACYC184-derivative plasmid, parD [kid74, kid[wt]], tet | This work                     |
| pBAD18           | pBR322-derivative expression vector, amp              | Guzman et al. (1995)           |
| pELI01           | pBAD18, Para::prfA121, amp                           | This work                     |
| pELI02           | pBAD18, Para::prfAwt, amp                            | This work                     |
| pELI03           | pBAD18, Para::prfA301, amp                           | This work                     |
| pELI04           | pBAD18, Para::prfA304, amp                           | This work                     |
| pELI05           | pFUS2, Para::kis, kan                                | This work                     |
| pELI06           | pFUS2, Para::kis74, kan                              | This work                     |
| pELI07           | pNDM220, Plac::relE, amp                             | This work                     |
| pELI08           | pFUS2, Para::prfAwt, kan                             | This work                     |
| pELI09           | pFUS2, Para::relB, kan                               | This work                     |
| pET80            | parD [kid[wt], kid[wt]], kan                          | Bravo et al. (1988)           |
| pMLM1            | mini-F, repFIA-, sop-, cat                            | Lemonnier et al. (2000)       |
| pFUS2            | pFUS-derivative expression vector, amp                | Lemonnier et al. (2000)       |
| pNDM220          | miniR1-derivative expression vector, amp              | Gottfredsen and Gerdes (1998) |
| pSS100           | pNDM220, Plac::kid, amp                              | Santos, Sierra et al. (2002)  |

of different Hfr-Tn10 strains (data not shown). To refine the genetic mapping of the mutations, the mutant strains were used as recipients in P1 transduction experiments performed using different donor strains bearing Tn10 transposon insertions at regular intervals between minutes 22 and 32 on the chromosome. For one of the mutants we observed that the pAB1120-resistance phenotype was lost at high frequency upon transduction from a donor containing the Tn10 at position 26.61 min. In order to isolate the mutant gene, fragments of the chromosome of the mutant strain (hereafter the KR19 strain), generated by partial Sau3A restriction digestion, were cloned into a low-copy-number mini-F, pMLM1 (Lemonnier et al., 2000). E. coli cells were transformed with this library, pooled and subsequently transformed with the pAB1120 plasmid. Mini-F plasmids carrying fragments that conferred resistance to pAB1120 were isolated and their chromosomal insert sequenced. All the sequenced fragments consisted in a chromosomal region enclosing the prfA gene encoding polypeptide release factor RF1. The localization of this gene at 27.2 min in the E. coli chromosome was consistent with the results of the Hfr mapping. Subsequently, the sequencing of the prfA region of KR19 and of several independent mutants isolated by this approach revealed that in all cases mutations in the prfA gene could be detected. Four independent mutants, hereafter called KR2, KR4, KR17 and KR19, were kept for subsequent studies. Sequencing analysis revealed that the mutations consisted of single-base G-C or A-T transitions, as expected from a 2-aminopurine mutagenic treatment (Ronen, 1980). Strains KR2 and KR19 were found to bear the same mutation, prfA301, which led to a Gly to Ser amino acid change at position 301 of RF1. Strain KR4 carried the mutation prfA303, in which an Arg has changed to Ser at position 303 of RF1. Finally, KR17 strain carried the mutation prfA121, a Gly to Ser change at position 121 of RF1.

The prfA mutants are unable to complement a prfA(ts) mutation unless they are overexpressed

To assess whether the isolated prfA mutants were affected in their translation termination activity in vivo, we performed a complementation assay using a conditional temperature-sensitive E. coli mutant, MRA8 prfA1(ts) (Ryden and Isaksson, 1984; Zhang et al., 1994). The strain was transformed with the pBAD18 plasmid derivatives pELI01, pELI02, pELI03 and pELI04, which carried the alleles prfA121, prfA wt, prfA301 and prfA303, respectively, under the control of the arabinose-inducible promoter pBAD (Guzman et al., 1995). MRA8 was also transformed with pBAD18 as a control. At basal expression levels (i.e. in the absence of arabinose), wild-type prfA expressed from pELI02 was able to complement the prfA1(ts) mutation at the non-permissive temperature of 42°C, whereas none of the mutants was able to do so (Fig. 1, left). When expression of the prfA alleles was induced by adding arabinose to the medium, the growth of strain MRA8 at 42°C was significantly recovered for all the mutants tested (Fig. 1, right). Since overexpression of the mutant proteins was able to achieve complementation, these results indicated that the mutations significantly decreased, but did not abolish, the efficiency of translational termination by RF1.

Decreased translation termination in vivo leads to readthrough of stop codons

Since the mutations affected RF1, which specifically recognizes UAG (amber) stop codons, a likely explanation...
for the phenotype of resistance displayed by the mutants was the readthrough of the \textit{kis74} amber mutation, which would allow the production of an active, full-length antitoxin. Wild-type \textit{E. coli} cells and KR19 mutant bacteria were transformed with either the plasmid vector pFUS2, or its derivatives pELI05 and pELI06 bearing the wild-type \textit{kis} and the mutant \textit{kis74} alleles, respectively, placed under the control of the arabinose-dependent pBAD promoter (Guzman \textit{et al.}, 1995). Following overexpression of the \textit{kis} alleles by addition of arabinose to the cultures, cell lysates were prepared and analysed by Western blot using anti-Kis antibodies (Fig. 2). A band corresponding to the full-length Kis protein was detected in the lysates obtained from wild-type and KR19 cells carrying the pELI05 (\textit{kis}+) plasmid. This band was not present in lysates of wild-type cells containing pELI06, which produced a Kis protein of a lower size resulting from the truncation of the protein at site 74. Note that this Kis74 protein could not be well detected because it co-migrated with a lower non-specific band. In strain KR19 containing pELI06, readthrough of the stop codon at position 74 yielded a Kis protein of higher molecular weight, most likely a full-length protein, indicating a decreased \textit{in vivo} efficiency of translation termination by the mutant RF1 protein at UAG stop codons. Similar results were also obtained for the KR17 mutant (data not shown).

\textit{RF1 mutations strongly reduce the specific termination activity of RF1 at UAG codons}

Altogether, these results suggested that the \textit{prfA} mutant strains KR4, KR17 and KR19 were deficient in translation termination activity at UAG codons. This hypothesis could be tested using a system described by Curran and Yarus (1989) and developed into a convenient and precise assay system by Poole \textit{et al.} (1995). This system is based on competition between translational termination and ribosomal frameshifting at the frameshift site present in \textit{E. coli} \textit{prfB}. At this site, an in-frame UGA stop codon is bypassed by a \textit{+1} ribosomal frameshift, stimulated by the presence of a Shine–Dalgarno sequence six nucleotides upstream of the stop codon. In the assay system, the RF2 frameshift window is fused to the \textit{malE} gene, expressed from a plasmid. Termination and \textit{+1} frameshifting lead to proteins of 43.9 and 52.0 kDa, respectively (Fig. 3A), that are separated by SDS gel electrophoresis and detected by Western blotting, binding of anti-MalE antibody and interaction with 125I-labelled Protein A. The ratio of the termination product to the

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\textbf{Fig. 1.} Complementation of a temperature-sensitive \textit{prfA} mutation. MRA8 [\textit{prfA1(ts)}] strain was transformed with the plasmids pELI01 (\textit{prfA121}), pELI02 (\textit{prfAwt}), pELI03 (\textit{prfA301}) and pELI04 (\textit{prfA303}). Expression of the \textit{prfA} alleles in these plasmids was placed under the control of a promoter whose activity could be induced by adding arabinose into the medium.

A. Uninduced cultures were grown at 42°C for up to 160 min.

B. Same experiment but arabinose (1% final) was present in the medium to induce the expression of the different \textit{prfA} alleles.

\textbf{Fig. 2.} \textit{In vivo} readthrough of the \textit{kis74} amber mutation in KR19 strain. Western blot analysis was performed using different cell extracts as described in \textit{Experimental procedures}. The black arrow indicates full-length Kis protein. Lanes 1, 3 and 5, cell extracts from strain MC1061 bearing pFUS2, pELI05 (\textit{kis}) and pELI06 (\textit{kis74}) plasmids, respectively; lanes 2, 4 and 6, cell extracts from strain KR19 carrying pFUS2, pELI05 and pELI06 plasmids, respectively; lane 7, purified His-tagged Kis protein. The thick band at the bottom of the gel is a non-specific product present in whole-cell lysates.
The frameshift product is a measure of relative termination efficiency at different stop signals or in different strains. The activity of the prfA mutant strains was tested at UAG stop codons in two contexts: UAGA and UAGU, both less favourable for termination than the most common UAG stop signal, UAGG. Western blots of this assay are shown in Fig. 3B, and the termination rate constants relative to frameshifting in Table 2. Expression of the fusion

**Table 2.** Termination rate at UAG codons and quantities of RF1 in prfA mutant strains.

|          | MC1061 | KR4    | KR17    | KR19    |
|----------|--------|--------|---------|---------|
| RF1 mutation | –      | R303H  | G121S   | G301S   |
| Termination rate at UAGA | 1.30   | 0.18   | 0.15    | 0.14    |
| Termination rate at UAGU | 1.42   | 0.16   | 0.06    | 0.12    |
| Relative quantity of RF1 | 100    | 80     | 85      | 85      |

Data are obtained by quantification of Western blots such as those shown in Fig. 3. Termination rate at UAGA and UAGU stop signals is expressed relative to +1 frameshifting in a frameshift window derived from the RF2 frameshift site. The quantity of RF1 in mutant strains, measured by reaction with anti-RF1 antibodies, is expressed as a percentage of that found in the parental strain MC1061.
proteins in the prfA mutant strains somewhat inhibited cell growth, which accounts for the reduced amount of fusion protein in these strains; for reasons that are unclear the inhibitory effect was more pronounced with the UAGA-containing fusion. The values found for the termination rate in parental strain MC1061 are close to those reported by Poole et al. (1995) but lower than those of Mora et al. (2007) in other E. coli K12 strains. It should be recalled that strain MC1061 is a streptomycin-dependent strain, grown in the presence of streptomycin, and that this is likely to influence the kinetic parameters for translation and translation termination. As seen in Table 2 (rows 2 and 3), termination rates for all three mutants at both UAGA and UAGU signals were strongly reduced with respect to the wild type by an average factor of about 10.

Two mechanisms might explain this result: the mutations may affect the folding or stability of RF1, resulting in a strongly reduced amount of RF1 in the cell. Alternatively, the activity of the protein may be reduced compared with that of the normal factor. To resolve this question, the amount of RF1 present in the prfA mutant strains was compared with the parental strain by quantitative Western blotting using anti-RF1 antibodies. The results (Fig. 3C, Table 2, row 4) showed that the reduction in the amount of RF1 in the mutant strains was quite modest (15–20%), insufficient to explain the 10-fold drop in termination rate in the cells; the major part in this drop must therefore be due to reduced activity of the mutant factors.

The small decrease in the amount of RF1 found in the mutant strains suggested that the mutant factors were not strikingly less stable than the wild-type factor. This conclusion might however be invalid if an autocontrol mechanism were present in cells, with the consequence that a lowered peptide release activity were to lead to a higher rate of synthesis of the RF. Such an autocontrol mechanism does exist in the case of RF2 in E. coli and in about 70% of microorganisms for which genome sequences are known (Baranov et al., 2002), based on the competition between termination and frameshifting that we have used to assay termination efficiency. The prfA gene is expressed from the promoter of the upstream gene, hemA. It has been suggested that E. coli and Salmonella may possess a mechanism for the autocontrol of RF1 synthesis, dependent upon readthrough of the weak UAGC tetranucleotide stop signal of the hemA gene, coupled with a suboptimal ribosome binding site for prfA expression (Elliott, 1989). The hypothesis suggests that RF1 synthesis may result to a significant extent from ribosomes that read through the hemA stop signal, translate the intergenic region, terminate just short of the prfA initiation codon and resume translation without dissociating from mRNA. To test this hypothesis, we constructed translational fusions with lacZ extending from the promoter region of hemA to codon 13 of prfA, analogous to those described by Dahlgren and Ryden-Aulin (2004), and studied the effect on β-galactosidase synthesis of mutating the hemA stop signal or introducing mutations in RF1. The results showed that even in the case of the RF1 mutants described in this article, little RF1 is in fact synthesized in E. coli by the proposed readthrough mechanism (L. Mora, unpubl. data). These results further confirm that the detected levels of the RF1 mutants reflect a minor effect on stability and therefore that the isolated mutations do not affect substantially the stability of RF1.

prfA mutation leads to hypersensitivity to Kid and RelE toxins

Since RF1 was shown to protect from RNA cleavage by the RelE toxin in vitro (Pedersen et al., 2003), we hypothesized that the mutant RF1 strains should be particularly responsive to the activity of RelE in vivo. For this analysis we selected the prfA mutant KR19 as this mutant showed the lowest overall termination efficiency in the assays (see Table 2). We produced the RelE protein in this mutated prfA strain and compared the levels of cytotoxicity with those achieved in a wild-type E. coli background. To gain insight into the mechanism of RF1-dependent protection, we performed a parallel control experiment in which the Kid toxin was produced in place of RelE. Figure 4 shows the cytotoxic effects of both Kid and RelE proteins over the wild type, MC1061, and the mutant strain KR19 bearing the plasmids pNDM220 (vector), pELI06 (kid) or pELI07 (relE). Figure 4A and D shows that the growth of the strains in the absence of induction of the toxins is unaffected. The partial induction of toxins Kid or RelE with a less-than-saturating concentration of IPTG (100 μM) inhibited cell growth in prfA mutant strains but not in the wild-type strain MC1061 (Fig. 4B, rows 4 and 6 compared with rows 3 and 5). This suggested that KR19 was hypersensitive to both toxins. When this strain was transformed with pELI08 (prfAwt), normal sensitivity to toxins was restored (Fig. 4C, rows 3–6), suggesting a specific effect of the mutations in prfA on the hypersensitivity to the toxins. The growth inhibition of the prfA mutants by the Kid and RelE toxins was neutralized when their respective antitoxin genes kis and relB were coexpressed (Fig. 4E, rows 3–6). Altogether these experiments brought to light a novel and specific functional interplay between RF1 and the Kid and RelE toxins.

Both RelE and Kid toxins are inhibitors of protein synthesis. We therefore evaluated the effect of these toxins on the de novo protein synthesis in the prfA mutant and wild-type strains at the suboptimal induction conditions mentioned above. The results (Fig. 5) indicated that indeed the RelE and Kid toxins inhibited protein synthesis in the prfA mutant but not in the wild-type strain. The enhanced inhibition of protein synthesis by Kid and RelE was prevented when their cognate Ks and RelB antitox-
ins were expressed in the cells, respectively (Fig. 5), indicating as in the previous assay that the inhibitory effect on protein synthesis is specific to the activity of the toxins. Protection by the RF1wt protein in this assay was less efficient, which is likely a reflection of the more specific interaction of a toxin with its cognate antitoxin (data not shown). Altogether, the increased efficiency in protein synthesis inhibition was consistent with the enhanced cytotoxic effect displayed by the Kid and RelE proteins in the mutant \( prfA \) background.

To test whether the extra-sensitivity to the toxins of the \( prfA \) mutants was an indirect consequence of a general stress on the translation apparatus we first compared the sensitivity of the wild type and mutant strain to antibiotics acting on translation, such as chloramphenicol or tetracycline, inhibitors of the peptidyl transferase activity or of aminoacyl tRNA binding respectively (Chopra and Roberts, 2001; Hansen et al., 2003). The sensitivity to different aminoglycosides (kanamycin, gentamicin and paromomycin) was also tested. These family of antibiotics interact with a region that is overlapping with the domain of interaction of RF1 (see Discussion).

As an additional control to test possible unspecific effects of the \( prfA \) mutation on other cellular processes, we used Nalidixic acid (an inhibitor of DNA Gyrase and an inducer of the SOS response) or Rifampycin (an inhibitor of RNA polymerase and therefore of transcription). No

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**Fig. 4.** Hypersensitivity of the \( prfA \) mutant strain KR19 to the Kid and RelE toxins and neutralization by cognate antitoxins and by \( prfA \)wt overexpression. Parental MC1061 and \( prfA \) mutant KR19 strains bearing the pNDM220 plasmid vector (lanes 1 and 2) or its derivatives encoding Kid (lanes 3 and 4) and RelE (lanes 5 and 6) toxins were grown exponentially to an \( A_{600} \) of 0.4 in LB medium. Serial dilutions (10\(^{-1}\) steps from left to right) of the different cultures were spotted in LB solidified medium. In (B) and (C), IPTG (100 \( \mu M \)) was included in the plates to induce the expression of the Kid and RelE toxins from the plasmids. In (A) and (D), the same experiment was performed but IPTG was omitted. Specificity controls were carried out by transforming the cells with plasmid pELI08 (\( prfA \)wt) (C and D) or with the plasmids encoding the respective antitoxins to Kid and RelE, namely plasmids pELI05 (\( kis \)) and pELI09 (\( relB \)) (E). The plates were incubated at 30°C and photographs were taken after 18 h incubation.

**Fig. 5.** Inhibition of protein synthesis in vivo by the Kid and RelE toxins. Strains MC1061 (top) or KR19 (bottom) were transformed with plasmids as indicated: lane (1) plasmid vector pNDM220, lane (2) pSS100 (\( kid \)), lane (3) pELI07 (\( relE \)), lane (4) pSS100 (\( kid \)) and pELI05 (\( kis \)), lane (5) pELI07 (\( relE \)) and pELI09 (\( relB \)). When relevant, expression of the toxin and antitoxin genes was triggered by adding IPTG (100 \( \mu M \)) and arabinose (0.5%) to the medium respectively. The proteins synthesized at time 0 or 60 min after the induction of the toxin and antitoxin genes were fractionated by SDS-PAGE (12.5%) and identified by autoradiography as described in Experimental procedures.
The differences between wild type and prfA mutants to the sensitivity to these inhibitors were found (Fig. 6). This further supports the specificity of the increased sensitivity to the RelE and Kid toxin observed in the mutants.

**Discussion**

In this article we report the isolation and characterization of novel mutations in prfA affecting the translation termination factor RF1. As previously described these mutations were selected by means of a novel TA system-based genetic screen leading to the efficient counter-selection of cells with intact translation termination capacities. The three mutant residues isolated in RF1 are grouped quite closely together and are nearby the region of RF1 known to be involved in stop codon recognition (Fig. 7). This includes a tripeptide motif (PxT, 188–190 in RF1, SPF, 205–207 in RF2) (Ito et al., 2000) important in the recognition of the first and second stop codon nucleotides (Laurberg et al., 2008) and amino acid residues at the tip of helix α5 (shown in blue in Fig. 7) in RF1 and RF2, which help discriminate against purines at the constant first position U of stop codons (Petry et al., 2005; Laurberg et al., 2008). One of the mutations, G121, is present at the tip of this helix, and is within about 6 Å of the two first nucleotides, U4 and A5 of the mRNA stop codon (Fig. 8, Table 3). Mutation R303 is close to A1492 and A1493 in 16S rRNA, involved both in recognition of sense codons in the A site by tRNA (Ogle et al., 2001; Ogle and Ramakrishnan, 2005) and in translation termination (Laurberg et al., 2008), as well as to other adjacent nucleotides. Both R303 and G301 are close to ribosomal protein S12 (Fig. 8, Table 3). Mutations in S12 are known to affect the accuracy of stop codon recognition. Thus all three mutations are situated close to the stop codon recognition site, which is consistent with their effect

**Table 3.** Components of mRNA, 16S rRNA and proteins of the 30S ribosomal subunit within about 6 Å of the mutant residues in RF1.

| Mutant residue | mRNA | 16S rRNA | 30S proteins |
|----------------|------|----------|--------------|
| G121           | U4, A5 | A1493, G1494 | S12: T43 |
| G301           |       | A1491, A1492 | S12: T43, P44 |
| R303           |       | A1491, A1492 | S12: T43, P44 |

The nearby components were identified using the structure file of the RF1:70S ribosomal complex (Laurberg et al., 2008). See also Fig. 8.
on the efficiency of translation termination mediated by RF1. It would be of interest to study their effect on the accuracy of stop codon recognition.

These novel prfA mutants show increased sensitivity to the RelE and Kid toxins. This phenomenon, detected both in growth and in protein synthesis assays, and the suppression of this sensitivity by the overproduction of prfAwt in vivo revealed an unexpected confluence in the pathway of action of these two toxins. Previous reports showed that the RelE toxin acts as an endoribonuclease in concerted action with the ribosome to cleave at the UAG termination codon at the A site of the ribosome, thus leading to translation inhibition. Consistent with this it was found that RF1 prevents RNA cleavage by this toxin (Pedersen et al., 2003). RelE seems to have a shape similar to the C-terminal region of translation elongation factor EF-G (Takagi et al., 2005), which could explain its access to the UAG codon in the A site of the ribosome. The protection conferred by RF1 against the action of RelE likely reflects a competition for the same substrate. The mutations could affect this competition ultimately favouring the action of the toxin. A similar explanation could account for the extra-sensitivity of the prfA mutant to paromomycin and other aminoglycosides, which interact with the 30S subunit of the ribosome at a region overlapping with the site of interaction of RF1. Paromomycin binding to 30S ribosomes directly displaces A1492 and A1493 from 16S rRNA helix 44 to a bulged-out state characteristic of ribosomes containing cognate tRNA in the A site (Ogle et al., 2001). The modes of binding of kanamycin and gentamicin to the ribosome are similar (Francois et al., 2005). The extrahelical position of A1493 sterically impedes RF1 binding (Laurberg et al., 2008), explaining the competitive inhibition of termination by paromomycin (Youngman et al., 2007). These aminoglycosides therefore introduce perturbations into the same region of the decoding site as the RF1 mutations we describe here, suggesting that the increased sensitivity of the mutant strains to the antibiotics is due to an accumulation of factors interfering with stop codon recognition by RFs.

Unlike RelE, the Kid and ChpAK (MazF) toxins are able to cleave RNA in the absence of ribosomes (Zhang et al., 2003; 2004; Munoz-Gomez et al., 2004; 2005) following a mechanism similar to the one described for RNase A and RNase T1 (Christensen et al., 2003; Pedersen et al., 2003; Kamphuis et al., 2006). Since the structure of the Kid protein is clearly different from that of RelE, similarities in their mechanisms to access the A site of the ribosome seem unlikely. It is noteworthy however that the Kid toxin can cleave the minimal substrate UpA which is part of the termination codon UAG/A (Kamphuis et al., 2006). It remains to be seen if the hypersensitivity to Kid observed in the prfA mutants could be due to interactions between Kid and RF1, which remain to be elucidated. Alternatively, in view of the fact that reduced activity of the RF1 can induce ribosome stalling at the stop codon and subsequent mRNA cleavage (Li et al., 2007), the increased toxicity of the prfA mutants to the Kid toxin could also be due to an enhancement of this stalling. Whatever the mechanism, it seems clear that the increased sensitivity of the prfA mutants to aminoglycosides and to the Kid and RelE toxins is a specific effect of these mutations rather than the unspecific consequence of a general stress on translation induced by them. This result provided the first evidence for the involvement of RF1 in the pathway of Kid toxicity.

The new prfA mutants contributed to showing that a post-transcriptional mechanism is not involved in maintain-
ing the levels of the RF1 in the cells. The mutants could be used to further explore the contribution of new residues close to the RF decoding region in the recognition of the termination codons and/or in the transmission of the signal that activates the transpeptidase of RF1 at the PTC distal site. In addition, the experiments reported provide the first in vivo evidence for a functional interaction between the RelE toxin and the RF1 protein. Furthermore, evidence is provided for a role of translation termination factors in the cytotoxicity pathway of the Kid toxin. The results in turn reveal an unexpected functional confluence between two distinct families of bacterial toxins, namely the RelE and Kid families. Similarities and differences in these pathways remain to be fully established. The hypersensitivity of the prfA mutants to the RelE and Kid toxins also opens the possibility of exploring their effects on other toxins belonging to the kis–kid/ccd and the relBE/parDE superfamilies (Gerdes et al., 2005). These include toxins targeting DNA gyrase such as CcdB and ParE in addition to toxins which acts on RNA as Kid and RelE. In theory it can be anticipated that the prfA mutants should also be extra-sensitive to the RNase-type toxins of the family but not to the toxins targeting the DNA Gyrase. Finally, the results and methods reported here provide a rationale for the design of screening approaches to isolate inhibitors of RF1 activity. Such compounds would likely potentiate the antibacterial activity of TA systems. Most interestingly, some antibiotics have been reported to specifically activate TA systems (Sat et al., 2001). In this respect as well, compound drugs inhibiting RF1 activity would be particularly appealing as they would provide a back-up induction of TA toxicity in antibiotic-resistant strains. Novel therapeutic alternatives against drug-resistant pathogens are urgently needed and the work reported here certainly opens new avenues towards that goal.

Experimental procedures

Bacteria strains

*Escherichia coli* K-12 strains used in this work are described in Table 4.

| Strain   | Genotype                                      | Reference                        |
|----------|-----------------------------------------------|----------------------------------|
| BL21 (DE3) | F^-ompT hsdS2 (rB- mB') gal dcm λDE3         | Novagen                          |
| MC1061   | F- araD139 galE15 galK16 rpsL hsdR2 (K- mK+) mcrA mcrB1 | *E. coli* Genetic Stock Center   |
| KR19 (KR2) | MC1061 prfA301                               | This work                        |
| KR4      | MC1061 prfA303                                | This work                        |
| KR17     | MC1061 prfA121                               | This work                        |
| MRA8     | prfA1(ts)                                     | Zhang et al. (1994)              |
| W3110    | F- λ rph-1 INV(mD, mE)                        | Bachmann (1972)                  |

*Table 4. Strains used in this work.*

Plasmid construction

Plasmids used and constructed in this work are listed in Table 1. To construct the pBAD18-derivative plasmids (Table 1), PCR fragments were generated using primers PRL (+) (5'-GACAGCTAGGGCTGGAGTA-3'; NheI site underlined) and PRR (-) (5'-GGTAGCATGCTCCAGGAGATTCTC-3'; SphI site underlined) and genomic DNA originated from the wild-type strain as well as from the different mutants as templates. The resulting fragments were introduced into pBAD18 vector digested with Nhel and Spel, to yield pELI01, pELI02, pELI03 and pELI04 plasmids.

Plasmids pELI05 and pELI06 (Table 1) were constructed by generating PCR fragments encoding the *kis* and *kis74* alleles using primers KISSN (+) (5'-GGCGAGCATATGGAGGG TGAAGAAT-3'; Ndel site underlined) and KIS CTR (-) (5'-GCTGGAATCTCAGAATTTTCCCTGGAAGTTT-3'; BglII site) and genomic DNA originated from the wild-type strain as well as from the different mutants as templates. The resulting fragments were introduced into pFUS2 after being digested with Ndel and BglII, resulting in the pELI05 and pELI06 plasmids.

To construct the pELI07 plasmid, a PCR fragment encoding *reIe* was generated using primers *reIe*B1 and *reIe*E2 described in Gottfredsen et al. (1998) and genomic DNA from W3110 as template. The PCR product containing *reIe* was introduced into pNDM220 after being digested with BamHI and Sall, yielding the pELI07 plasmid.

The pELI08 plasmid was constructed using a fragment containing the wild-type prfA gene generated from the diges-
tion of pELI02 with HindIII and NsiI inserted into the pFUS2 vector cleaved with the same enzymes.

The pELI02 plasmid was constructed by introducing a PCR fragment encoding relB into pFUS2 after a double digestion with Ndel and HindIII. The PCR fragment was generated using W3110 as DNA template and degenerated primers relB1 and relB2 described in Gotfredsen and Gerdes (1998), relB1 (5′-CCGCCCATATGTAAATTACAGATCGTAAGC-3′) and relB2 (5′-CCCCCTCGAGAGCTTACAGGTCCATCCACGCACGTTACTGG-3′).

pKK1120 (p15a, kis74, kid) was constructed by ligating the 1.2 kb FspI–EcoRI fragment of pAB1120 containing the kis74 (amber)-kid mutant operon and the 3.8 kb EcoRI–Scal vector fragment from pACYC184.

Western blotting for readthrough assay

Cultures growing overnight in LB media were diluted 1:100 and induced with 1% arabinose when the OD600 reached 0.1. At OD600 0.3, aliquots of 2 ml were centrifuged and the pellet was re-suspended in 60 μl of Laemmli buffer (Laemmli, 1970) and boiled for 10 min in order to obtain the cell lysates. Sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) was performed with a Hoefer SE 260 apparatus in the presence of running buffer [Tris 0.025 M, glycine 0.192 M, SDS 0.1% (w/v) pH 8.5]. The separating gel contained 15% (w/v) of acrylamide. Following electrophoresis, proteins were transferred onto polyvinylidene difluoride (PVDF) membrane (Sequi-Blot Bio-Rad Laboratories) using a Trans-Blot SD Semi-Dry Transfer Cell apparatus (Bio-Rad Laboratories) at a constant current of 12 V for 90 min at room temperature. PVDF membrane was blocked overnight using a buffer containing KCl 3 mM, NaCl 68 mM, KH2PO4 9 mM and re-suspended in 10 ml of lysis buffer (50 mM Tris-HCl, pH 6.8, 100 mM DTT, 2% SDS, 0.1% bromphenol blue, 10% glycerol). Aliquots of 20 μl in the case of the parental strain MC1061 and volumes containing equivalent amounts of total protein in the case of the mutant strains were applied to 10% polyacrylamide gels for separation as described by Laemmli (1970). Transfer to nitrocellulose membranes, Western blotting with antibodies and quantification were performed as described above. Western blots were performed with rabbit anti-RF1 diluted 1:5000 commercially prepared from pure factor.

Complementation assays

Cultures of strain MRA8 (prfA1ts) bearing pBAD-derivative plasmids (prfA wild type and mutants) were grown overnight at 30°C prior to being diluted 1:100 into two different aliquots of fresh medium (LB medium; ampicillin, 25 μg ml−1), with or without arabinose (0.2% v/v). The cultures were transferred to 42°C, and the OD600 of both cultures was monitored every 20 min over a period of 160 min.

Toxicity assays

MC1061 wild-type and prfA mutant strains bearing pNDM220-derivative plasmids containing or not pELI08 were grown at 30°C to mid-exponential phase in LB broth plus ampicillin (50 μg ml−1) and kanamycin (as needed, 50 μg ml−1). The same strains containing the pFUS2 derivatives containing the antitoxin genes were grown in a similar way. Aliquots of the different cultures and serial dilutions (10−1 steps) were spotted onto plates of the same solidified medium containing or not IPTG 100 μM as inducer of the toxins and were incubated at 30°C for 18 h. Antitoxins were induced using 0.5% arabinose. A semi-quantitative assessment of the relative levels of toxicity was derived from the differential dilution factor obtained for two spots presenting an equivalent number of colonies. Similar procedures were used to test the differential sensitivity of KR19 and MC1061 strains to the different antibiotics.

In vivo de novo synthesis of 35S-labelled proteins

One millilitre of cultures of wild-type and mutant strains KR19 bearing pSS100 (kid) plus or not pELI08 (prfAwt) or pELI05 (kis), or KR19 bearing pELI07 (relE) plus or not pELI08 (prfAwt) or pELI09 (relB) were labelled with 14.3 μCi of [35S]Met/Cys (80:20, Amersham Biosciences) for 2 min at 37°C, then 1 mg of unlabelled methionine was added and incubation was continued for 8 min at 37°C. Cells were centrifuged at 20 000 g for 2 min at 4°C, the pellet was washed with a buffer containing KCl 3 mM, NaCl 68 mM, KH2PO4 1.5 mM and NaH2PO4 9 mM and re-suspended in 10 μl of...
protein loading buffer 2× [0.05 M Tris-HCl pH 6.8, 10% (w/v) SDS, 0.01 M EDTA 25% (w/v), Bromophenol blue 0.5 g l⁻¹ and β-mercaptoethanol 5% (v/v)]. Proteins were separated by 12.5% SDS-PAGE and the radiolabelled products were identified by autoradiography.

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