Unravelling post-transcriptional PrmC-dependent regulatory mechanisms in

*Pseudomonas aeruginosa*

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Running Title:

Post-transcriptional regulation in *P. aeruginosa*
**Originality-Significance Statement**

By applying SILAC technology in combination with mRNA-sequencing we provide evidence that the regulatory control of translation termination via the methyltransferase PrmC influences target gene expression in the opportunistic pathogen *Pseudomonas aeruginosa* and has a major impact on the bacterial phenotype. The finding that a bias in stop codon usage determines the *P. aeruginosa* phenotype is unexpected and adds complexity even to well-studied regulatory circuits. Modulation of PrmC activity allows the bacterial cell to cross-regulate targets independently of transcriptional signals. It represents a post-transcriptional regulatory mechanism with an underestimated but major impact on the bacterial phenotype.

**Summary**

Transcriptional regulation has a central role in cellular adaptation processes and is well investigated. In contrast, the importance of the post-transcriptional regulation on these processes is less well defined. The technological advancements have been critical to precisely quantify protein and mRNA level changes and hold promise to provide more insights into how post-transcriptional regulation determines phenotypes. In *Pseudomonas aeruginosa* the methyltransferase PrmC methylates peptide chain release factors to facilitate translation termination. Loss of PrmC activity abolishes anaerobic growth and leads to reduced production of quorum sensing-associated virulence factors. Here, by applying SILAC technology in combination with mRNA-sequencing, we provide evidence that the *P. aeruginosa* phenotype can be attributed to a change in protein to mRNA ratios of selected protein groups. The UAG-dependent translation termination was more dependent on PrmC activity than the UAA- and UGA-dependent translation termination. Additionally, we found a bias towards UAG stop codons in global transcriptional regulators. The finding that this bias in stop codon usage determines the *P. aeruginosa* phenotype is unexpected and adds complexity to regulatory circuits. Via modulation of PrmC activity the bacterial cell can cross-regulate targets independently of transcriptional signals, a process with an underestimated impact on the bacterial phenotype.
The ability of bacteria to adapt to various environmental conditions is one reason for their evolutionary success. A prime example for effective adaptability is the Gram-negative bacterium *Pseudomonas aeruginosa*, which is a ubiquitous environmental bacterium and an opportunistic human pathogen. It plays a dominant role as the causative agent of often devastating hospital-acquired infections. Furthermore, *P. aeruginosa* is frequently recovered from biofilm-associated chronic persistent infections (Bodey et al., 1983; Costerton et al., 1999; Foxman, 2010). Today, *P. aeruginosa* is the most prevalent pulmonary pathogen in cystic fibrosis (CF) patients (Folkesson et al., 2012). *P. aeruginosa* persists in the lungs of these patients within biofilms, which provide protection against antimicrobial therapy and the host immune response. Chronic respiratory tract infections in CF are associated with ongoing inflammation and changes in the structure and function of the affected organ and thus largely determine morbidity and mortality. The increasing development of multi-drug resistant *P. aeruginosa* isolates aggravates the management of acute and chronic infections and further restricts therapy options (Breidenstein et al., 2011).

*P. aeruginosa* is well-known for its remarkable adaptability to various and changing environmental conditions. Survival in altering habitats is largely reflected by changes in the transcriptional profile and a complex network of transcriptional regulators governs e.g. the switch of bacterial lifestyles from a unicellular planktonic to a multicellular biofilm state as well as the production of virulence factors (Jimenez et al., 2012). Although transcriptional regulation is a key step in the control of bacterial gene expression, post-transcriptional regulation seems to play a larger role than previously expected. Examples of post-transcriptional regulators are small regulatory RNAs (sRNAs) or the CsrA/RsmA system which both are involved in highly sophisticated regulatory circuits (Wassarman, 2002; Ventre et al., 2006; Brencic and Lory, 2009; Romeo et al., 2013). Also the modification of the translation machinery can affect RNA processing and thus represents a post-transcriptional regulatory mechanism (Pustelny et al., 2013; Shi et al., 2015; Little et al., 2016). The methyltransferase PrmC methylates class one peptide chain release factors (PrfA, PrfB) on a
glutamine residue of the GGQ motif in various bacteria including *Escherichia coli* and *P. aeruginosa* (Heurgue-Hamard et al., 2002; Nakahigashi et al., 2002; Scarlett et al., 2003; Pustelny et al., 2013). These release factors are responsible for the termination of translation at the stop codons UGA, UAA and UAG (Mora et al., 2007). Deletion of PrmC results in inefficient translation termination and thus downstream sequences are translated which may also include frameshifts (Nakahigashi et al., 2002; Mora et al., 2007). However, rather than generally determining bacterial fitness and growth as a result of inefficient translation termination, loss of PrmC seems to affect specifically bacterial pathogenicity and environmental adaptability. For example a VagH (PrmC-orthologue) deficient *Yersinia pseudotuberculosis* mutant was demonstrated to exhibit an avirulent phenotype similar to a type 3 secretion system (T3SS) negative mutant, and thus it was suggested that absence of VagH affects expression of the T3SS (Garbom et al., 2007). In addition, *prmC* gene expression was shown to be increased in *Porphyromonas gingivalis* and *E. coli* under *in vivo* conditions (Park et al., 2004; John et al., 2005), indicating a potential regulatory function of PrmC that drives pathogenicity. In *P. aeruginosa*, PrmC deficiency resulted in a quorum sensing (QS)-independent reduction of virulence factor synthesis, e.g. pyocyanine, rhamnolipids and a lower production of secreted toxins (Pustelny et al., 2013). However, how PrmC selectively affects pathogenicity in *P. aeruginosa* remained unknown.

In order to address this question, we used stable isotope labeling by amino acids in cell culture (SiLAC) in combination with shotgun proteomics and data dependent mass spectrometry (MS) analysis to determine the impact of PrmC on global *P. aeruginosa* protein biosynthesis. SiLAC is a metabolic labeling strategy and well-established for quantitative proteome analysis in eukaryotic cells (Ong et al., 2002), whereas its application for the analysis of bacterial systems has remained very limited (Soufi et al., 2010; Ping et al., 2013; Jung et al., 2014; Boysen et al., 2015; Chua et al., 2016). In this study we adapted the SiLAC technique to quantify even small magnitude effects in the *P. aeruginosa* proteome and characterized global effects of PrmC deficiency. This comparative proteome analysis was furthermore combined with RNA-seq data in order to precisely decipher
transcriptional and post-transcriptional PrmC-dependent changes and thus provide a detailed view on the mode of action of PrmC.

**Results**

Previous studies revealed a strikingly attenuated phenotype of the PrmC deficient *P. aeruginosa* PA14 mutant in the *Galleria mellonella* infection model (Pustelny et al., 2013). This might be explained by a reduced production of many QS-dependent virulence factors despite normal levels of the QS signaling molecules and/or the inability of the mutant to grow under anaerobic conditions (Pustelny et al., 2013). In order to gain more insights into the molecular mechanisms underlying the attenuation of the PrmC deficient *P. aeruginosa* PA14 mutant, we adapted the metabolic labeling strategy SILAC in combination with shotgun proteomics in the *P. aeruginosa* PA14 strain.

**Adaption of SILAC based protein quantification in *P. aeruginosa***

Sufficient and efficient labeling of peptides by stable isotopic labeled amino acids is an absolute prerequisite for global protein quantification using SILAC. Auxotrophy for the amino acids lysine and arginine ensures high incorporation rates of exogenously added stable isotopic labeled equivalents in eukaryotes (Ong et al., 2002; Dreisbach et al., 2008). Therefore, the first *P. aeruginosa* SILAC experiments in this study were performed in a PA14 ΔlysAΔargB double mutant, which is auxotroph for both amino acids. However, when $^{13}$C$_6$N$_2$-L-lysine (lys8) and $^{13}$C$_6$N$_4$-L-arginine (arg10) were exogenously added, the identification rate of proteins decreased in this auxotrophic mutant over time (Figure S1). Since a successful application of lys8 protein labeling has previously been described for prototroph yeast and *E. coli* (Frohlich et al., 2013), we next fed the PA14 wild-type strain with lys8 only. However, again no stable identification rates could be obtained (Figure S1). To avoid lysine degradation and reincorporation of heavy stable isotopes into amino acids other than lysine, the PA14 tnldcC transposon mutant from the Harvard PA14 mutant library was selected (Liberati et al., 2006), which lacks a functional LdcC protein involved in degradation of lysine (Chou et al., 2010). The number of identified proteins remained constant following lys8 protein labeling in this mutant and
the stability of stable isotopic labeled lysine was ensured (Figure S1). By extending the exposure time of the bacteria to the labeled amino acid a SILAC incorporation efficiency of 98% was reached that nearly reflected the isotopic purity of $^{15}$N and $^{13}$C in lys8. In conclusion, lys8 was used for protein labeling and combined with LysC digestion (Figure S2) for quantitative proteome analysis in *P. aeruginosa*.

**Robustness of SILAC to quantify even small magnitude effects in the *P. aeruginosa* proteome**

We next investigated the robustness of SILAC based protein quantification in *P. aeruginosa* and determined the overall degree of variation of the quantified proteins between biological replicates. We therefore grew two PA14 tnlC cultures under identical conditions and supplemented one with unlabeled (lys0) and the other with labeled (lys8) lysine. We extracted the proteins from lys0 and lys8 labeled bacteria, mixed them in a 1:1 ratio and subjected them to shotgun proteomics to determine labeled/unlabeled protein ratios. We calculated the threshold for reliable fold-changes in protein abundance to be a ratio of 1.2 which corresponds to a 95% confidence interval of all protein ratios (Figure S3).

**Quantitative, PrmC dependent proteome analysis**

We used the adapted SILAC technique to evaluate whether production of *P. aeruginosa* proteins and particular those that drive pathogenicity were affected by expression of the release factor methyltransferase PrmC. We therefore performed comparative proteomics of the PA14 tnpC strain harboring the empty vector as control (tnprmc) against the PA14 tnpC mutant complemented with the *prmC* gene *in trans* (tnprmc::prmC) in order to ensure measurement of PrmC-dependent effects in the identical strain background. The lys8 labeled proteins extracted from the PA14 tnlC mutant (tnlC) were used as “spike in” standard (Geiger et al., 2011). Therefore the tnpC and tnpC::prmC samples were mixed with the heavy labeled reference and analyzed individually by liquid chromatography (LC)-MS/MS analysis. Ratios between the heavy labeled “spike
in” proteins and the unlabeled proteins from the samples were generated and compared to determine the final ratios for the relative protein abundance of the tnprmC versus tnprmC::prmC samples (Fig. 1). Replicates were plotted against each other to evaluate the biological reproducibility of protein changes. All replicates showed high correlation (Squared Pearson’s correlation coefficient of 0.87-0.93, Figure S4).

Among the overall 2722 detected and quantified proteins, 1699 proteins were found in all three replicates of the tnprmC and tnprmC::prmC strains (listed in Table S1). Contrary to the 2D-based proteome analysis, which revealed only 26 quantified proteins (Pustelny et al., 2013), this extensive data set enables global similarity analysis of termination sequences. In the SILAC based analysis we observed 94 proteins with a significant higher abundance and 296 proteins with a significant lower abundance in the tnprmC mutant (p-value ≤ 0.05; Fig. 2). This highlights that in general PrmC deletion negatively affects the protein translation in P. aeruginosa. In accordance with the previous observation of a lower virulence factor production in tnprmC (Pustelny et al., 2013), the mutant produced lower levels of e.g. proteins involved in phenazine biosynthesis (PhzM, PhzB1, PhzB2, PhzS, PhzE2, PhzD2; Table S1) and of enzymes for rhamnolipid synthesis (RhlB, RmlC; Table S1).

**Proteome-transcriptome correlation**

Changes on protein level in the tnprmC mutant may be the direct result of a translational termination deficiency. However, if protein levels of e.g. transcriptional regulators were affected, those might transcriptionally affect downstream targets and thus may lead to lower corresponding protein levels as a secondary effect of inefficient translation termination. We therefore also acquired RNA-seq data in order to examine the correlation between protein and mRNA abundance (Table S2).

We first compared the mRNA data of the tnprmC::prmC strain with its global protein quantification data (Table S3). A correlation analysis revealed a squared Pearson’s correlation coefficient of 0.20 of protein to mRNA abundance (Figure S5). Nevertheless, in the group of genes that exhibited a high protein to mRNA ratio (upper 25% of the genes) the functional PseudoCAP categories “amino acid
were enriched. Vice versa in the group of genes that exhibited a low protein to mRNA ratio (lower 25% of the genes) the functional categories “secreted factors (toxins, enzymes, alginate)”, “transcriptional regulators” and “two component regulatory system” were enriched (Figure S5).

We also recorded the transcriptional profile of tnprmC mutant and compared it to the RNA-seq data of tnprmC::prmC strain. We observed 90 genes with a significantly higher expression and 183 genes with a significantly lower expression in the tnprmC mutant (adjusted p-value ≤ 0.05, (Anders and Huber, 2010), genes are listed in Table S2). Among the latter were genes involved in phenazine biosynthesis and other QS-regulated genes. This indicates that the lower levels of QS-dependent virulence factors in the prmC mutant are due to lower mRNA abundances and not necessarily due to an inefficient translation termination of the respective mRNAs.

In order to identify those proteins that are directly post-transcriptionally influenced by PrmC, the protein to mRNA correlation ratios were calculated for tnprmC and tnprmC::prmC. Therefore, all quantified proteins (1699) were assigned to the corresponding mRNA data. The high correlation coefficient ($r^2=0.92$) indicates that most of the ratios were independent of the presence of PrmC. However, some proteins diverged and exhibited PrmC-dependent higher or lower protein abundance in relation to the level of the corresponding mRNA (Fig. 3).

One would expect a lower protein to mRNA ratio if mRNA was not sufficiently translated due to inefficient translation termination in the tnprmC mutant. Those 100 genes that exhibited the lowest PrmC-dependent ratio between protein and mRNA (top100 low) are listed in Table S4. The 100 genes with the highest ratio between protein and mRNA are listed in Table S5 (top100 high). Interestingly, genes of the functional PseudoCAP category “Chaperones & heat shock proteins” were highly enriched in the top100 high group, whereas genes of the functional category “Transport of small molecules” were enriched in the top100 low group (Figure S6).
Among the top100 low group of proteins we did not identify a transcriptional regulator of one of the P. aeruginosa QS-systems that might directly explain the lower production of virulence factors in the tnpmC mutant. However, the global regulator DNR was identified at lower protein levels in the tnpmC mutant. DNR controls the expression of several denitrification genes (Giardina et al., 2011).

Under anaerobic conditions, nitrate or nitrite are used as the terminal electron acceptor in P. aeruginosa. Since the tnpmC mutant has previously been shown to be severely affected under anaerobic conditions, we complemented the tnpmC mutant with the DNR gene in trans (tnpmC::dnr) and determined growth under anaerobic conditions. As shown in Fig. 4A the tnpmC mutant could be indeed partially rescued by dnr complementation under anaerobic growth conditions in glucose containing medium. Incomplete complementation might be explained by inefficient dnr translation even if the transcript level is increased. However, restoration of anaerobic growth via dnr expression in trans was not sufficient to compensate the lower virulence of the tnpmC mutant in the Galleria mellonella infection model (Fig. 4B). Interestingly the infection assay revealed that pyocyanine production is not required for full pathogenicity.

**tnpmC is differentially regulated under diverse environmental conditions**

If the activity of PrmC impacts on post-transcriptional regulatory processes, it is reasonable to assume that tnpmC itself might be subject to regulation by environmental cues. We therefore re-analyzed previously recorded data on mRNA expression levels in PA14 grown under 14 different conditions (Dotsch et al., 2015). Indeed, tnpmC was significantly differentially expressed under following conditions as compared to exponential growth phase (in LB medium): stationary growth phase, attachment experiments in M9 minimal medium (Godeke et al., 2012), heat shock at 42 °C or 50 °C, in vivo growth within a mouse tumor and iron deficiency (Figure S7). The changes in tnpmC transcript level could be caused by a differential in transcription initiation or in mRNA stability. We also quantified tnpmC mRNA levels in 369 clinical P. aeruginosa isolates. There was a high coefficient of variation (CV) for tnpmC of 0.36 across all isolates. This CV value was significantly higher than that of six housekeeping genes (proC, rpoD, nuoD, ppsA, nadB, mutL) (Figure S8). Remarkably, heat shock
proteins were enriched in the top100 high group of PrmC-dependent protein to mRNA ratios. Thus, as prmC expression is affected by heat shock, this regulatory mechanism might allow for an optimized translation efficiency of heat shock proteins after heat shock treatment.

The UAG stop codon is enriched in genes with low PrmC-dependent protein to mRNA ratio

Since PrmC is involved in the activation of peptide chain release factors that are important to terminate translation at the UAG, UAA, or UGA stop codons, we explored whether there is a difference in the frequency of the three stop codons in the top100 low versus the top100 high group of genes. Indeed, the UAG codon was enriched in the top100 low group. Thus, overall efficient UAG-dependent translational termination of mRNAs seemed to be more dependent on the methylation of peptide chain release factors than UAA- and UGA-dependent translation termination (Fig. 5). Next, distinct functional PseudoCAP gene categories were analyzed for their stop codon preference. Interestingly, an enrichment of the UAG stop codon was found in the category “transcriptional regulators” (Figure S9). This enrichment is particularly noteworthy, since changes in the abundance of transcription factors have major effects on the cellular protein profile. Concordant with the PseudoCAP enrichment of the category “chaperones & heat shock proteins” in the top100 high group, the analysis revealed a low UAG but a high UAA stop codon frequency, indicating that the UAA stop codon is less associated with PrmC deficiency.

PrmC deletion results in higher readthrough rates of translation associated proteins

The MS-raw data were reanalyzed and scanned for C-terminal successional peptides to determine whether global readthrough events were more frequent in PA14 tnpmC. In total, 273 proteins with readthrough events were detected; 63 readthrough peptides were found in frame, 97 with a +1 frameshift and 153 with a +2 frameshift. All these peptides were identified as specific LysC peptides and had an average Andromeda peptide ion score of 109 indicating highly specific detection (Cox et al., 2011). The intensities of readthrough peptides of the tnpmC strain were compared with that of tnpmC::prmC to identify whether they are PrmC-dependent. Despite the role of PrmC in the
activation of the peptide chain release factor, most of these readthrough events were detected independently of $prmC$ expression. Nevertheless, 23 readthrough events could be detected exclusively in the PA14 $tnprmC$ strain in at least two of three replicates (in frame: 5 proteins; +1 frameshift: 9 proteins; +2 frameshift: 9 proteins; Table S6, Table S7, Table S8), whereas only three readthrough events were detected exclusively in at least two replicates of the $tnprmC::prmC$ strain (in frame: 0, +1 frameshift: 2 and +2 frameshift: 1). Of note, 18 of these PrmC deficiency dependent peptides, were detected in the low molecular weight fraction of the SDS gel and thus were separated from the mature proteins. This indicates that proteins with readthrough events upon PrmC deletion are more likely degraded by endogenous proteases.

Discussion

The capability of $P. aeruginosa$ to strive in various and challenging habitats is facilitated by its versatile metabolic capacity, which is controlled by environment-driven flexible changes in the transcriptional profile. Although transcriptional regulation is the major key in control of bacterial gene expression, post-transcriptional regulation seems to play a much larger role than previously anticipated.

With the aim to shed more light on post-transcriptional regulation in the opportunistic human pathogen $P. aeruginosa$, we adapted SILAC technology to precisely quantify protein abundances in this organism and analyzed the effect of PrmC on protein and corresponding mRNA levels. PrmC methylates class one peptide chain release factors (PrfA, PrfB) which control efficient termination of translation (Heurgue-Hamard et al., 2002; Nakahigashi et al., 2002; Pustelny et al., 2013). Altered PrmC protein levels are therefore expected to result in global proteomic changes. We have previously characterized the $P. aeruginosa$ PA14 $tnprmC$ mutant (Pustelny et al., 2013) which did not exhibit a growth defect under aerobic conditions. However, the PrmC protein was found to be essential for growth under anaerobic conditions and pathogenicity of $P. aeruginosa$ in the Galleria mellonella infection model (Pustelny et al., 2013).
In this study, we detected and quantified as many as 1910 proteins in *P. aeruginosa* PA14 following lys8 labeling in a LC-MS/MS based shotgun proteome analysis. Remarkably, auxotrophy for lysine was not necessary to reach incorporation rates of more than 98%. These results are in accordance with a study of Fröhlich *et al.* demonstrating that stable isotopic labeled lysine can be used as a labeling strategy for protein quantification experiments of prototroph microorganisms (Fröhlich *et al.*, 2013). However, in this study the prevention of *P. aeruginosa* lysine degradation seemed to be important to ensure stable labeling with lys8.

Various studies in bacteria and eukaryotes revealed a rather low correlation of protein and mRNA levels, which might indicate the importance of post-transcriptional regulation (Maier *et al.*, 2009; Wu *et al.*, 2013; Kwon *et al.*, 2014; Borirak *et al.*, 2015). In this study, the application of RNA-seq and SILAC-based protein quantification allowed for the detailed analysis of mRNA to protein ratios in *P. aeruginosa* and uncovered direct and indirect PrmC induced changes in the protein profile. The overall high correlation of the ratio of mRNA to protein levels in the *prmC* mutant as compared to the complemented *P. aeruginosa* strain clearly demonstrated that PrmC-dependent changes on translation efficiencies of mRNAs is not a common effect on all proteins but is rather selective. Our study additionally revealed that *prmC* is differentially regulated under various environmental conditions, e.g. after heat shock treatment. Remarkably, heat shock proteins were enriched in the top100 high group of PrmC-dependent protein to mRNA ratios. Thus, this regulatory mechanism seems to allow for an optimized translation efficiency of heat shock proteins after heat shock treatment.

Global regulators have recently been suggested to be subject of extensive post-transcriptional regulation in bacteria (Schmidt *et al.*, 2016). Here we found that translation of the mRNA of selected genes was dependent on PrmC function. This bias can at least in part be explained by a differential use of stop codons in affected mRNA. Intriguingly, on a global scale the use of stop codon UAG entailed a lower and the UAA stop codon a trend to a higher PrmC-dependent protein to mRNA ratio. Interestingly, transcriptional regulators exhibit an overall higher content of UAG stop codons. Our
finding that the UAG stop codon is responsible for a differential translational termination efficiency upon PrmC deletion is supported by studies of Mora et al. (Mora et al., 2007). They clearly presented that the readthrough rate in *E. coli* strongly increases upon PrmC deletion if UAG terminates the open reading frame. Despite the general trend, not all of the mRNAs encoding for proteins that were negatively affected by PrmC deficiency harbored UAG as the stop codon and not all mRNAs that were more efficiently translated harbored the UAA stop codon. Other factors play a role in the PrmC-dependent changes on protein to mRNA ratios. The identity of the adjacent codons might very well aggravate or mitigate the effect of PrmC-dependent translation termination efficiency at the various stop codons. Furthermore, C-terminal elongation of proteins due to stop codon readthrough might be neutral, but they might also stabilize or destabilize the protein. Moreover, the readthrough might not be complete and, for an individual protein, a fraction might be correctly terminated (Mora et al., 2007). This could vary from protein to protein and also under changing environmental conditions. In *P. aeruginosa* at least some of the PrmC-dependent changes in protein abundance seem to be due to a destabilization of the protein, since we detected the major part of readthrough peptides upon PrmC deletion in the lower molecular weight fraction. The artificially extended proteins may also be nonfunctional because of misfolding. Notably, Beznoskov et al. and Eswarappa et al. reported programmed readthrough in yeast and mammalian endothelial cells, respectively (Eswarappa et al., 2014; Beznoskova et al., 2015). Their results indicate that certain proteins acquire new or differing functions by C-terminal extension. This could also apply for bacterial proteins and would indicate new ways of post-transcriptional regulatory mechanisms.

In conclusion, by using advanced protein profiling technology we uncovered a mechanism of gene regulation that is based on a bias in stop codon usage in selected functional groups of genes. Modulating the activity of PrmC and thus readthrough events at defined stop codons that impact protein stability adds complexity even to well-studied regulatory circuits. This strategy allows the bacterial cell to cross-regulate targets independently of transcriptional signals, a process with an underappreciated but major impact on the bacterial phenotype.
Materials and methods

Bacterial strains, media and growth conditions

Strains, plasmids, and primers that were used in this study are listed in Table S9. All chemicals were provided by Sigma-Aldrich or Carl Roth unless otherwise stated. PA14 strains were cultured in modified M9 medium containing 42.2 mM Na$_2$HPO$_4$ x 2 H$_2$O, 22 mM KH$_2$PO$_4$, 18.7 mM NH$_4$Cl, 8.6 mM NaCl, 1 mM MgSO$_4$, 0.1 mM CaCl$_2$, 20 mM Glucose, 10 µM Fe(II)sulfate and 0.1 mM of each of the 20 canonical amino acids except lysine and arginine unless otherwise stated. Isotopically labeled amino acids or unlabeled counterparts were used at the following concentrations: $^{13}$C$_6^{15}$N$_2$-L-lysine (lys8): 1.368 mM; $^{13}$C$_6^{15}$N$_4$-L-arginine (arg10, both SILANTES): 0.949 mM. For anaerobic growth experiments, strains were precultivated in Luria broth (LB) medium. Washed and concentrated precultures were used to inoculate main cultures in Hungate tubes containing PYG-KNO$_3$ medium (modified from http://www.dsmz.de/microorganisms/medium/pdf/DSMZ_Medium104.pdf: 20 g/l casein hydrolysate, 10 g/l yeast extract, 10 g/l glucose, 500 mg/l cysteine-HCl, 10 mg/l CaCl$_2$ x 2 H$_2$O, 20 mg/l MgSO$_4$ x 7 H$_2$O, 40 mg/l K$_2$HPO$_4$, 40 mg/l KH$_2$PO$_4$, 400 mg/l NaHCO$_3$, 80 mg/l NaCl, 100 mM KNO$_3$, distributed under N$_2$) to a start OD$_{600}$ of 0.01. Growth was monitored by taking samples of 100 µl with a cannula and subsequent OD$_{600}$ measurement. Incorporation efficiency of lys8 into the proteome of the tn$\Delta$dcc mutant was determined by preparation of 10 mL precultures in modified M9 medium either supplemented by lys0 or lys8. Each preculture was used to inoculate a main culture supplemented with lys8 with a start OD$_{600}$ of 0.05. Cells were harvested after 6 and 9.5 hours incubation at 37 °C and 180 rpm. Each sample was prepared in triplicates. To obtain protein extracts and mRNA samples for comparative quantitative LC-MS/MS analysis and RNA-seq, cultures of the strains tn$\Delta$prmC and tn$\Delta$prmC::prmC were harvested at an OD$_{600}$ of 2. To generate heavy labeled protein as internal standard, tn$\Delta$dcc cells were grown in the same defined M9 medium but containing stable isotope labeled lys8 instead of lys0 up to an OD$_{600}$ of 2. Before protein extraction, the tn$\Delta$prmC and the complemented mutant cells were mixed 1:1 with lys8 labeled tn$\Delta$dcc “spike-in”. Carbenicillin was added to a final concentration of 400 µg/ml to strains harboring the pUCP20 plasmid.
Preparation of LC-MS/MS samples, LC-MS/MS analysis and data processing

Bacterial cells were lysed by SDS-lysis buffer (20 mM TRIS-HCl, pH 6.8, 2% SDS, 6.8% glycerol, 20 mM DTT, 1x Protease inhibitor (Roche)) and subsequent sonification. Extracted proteins were fractionized by SDS-PAGE and conducted for in gel digestion using LysC (Wako Chemicals). Measurement of peptide samples was performed as described previously (Schroder et al., 2015) with minor changes using a reversed phase nanoflow ultrahigh pressure liquid chromatography (RSLC) system (Thermo Fisher Scientific) and an LTQ Orbitrap Velos mass spectrometer (Thermo Fisher Scientific). The raw files generated by LC-MS/MS analysis were further processed using the MaxQuant software package (version 1.5.2.8) (Cox and Mann, 2008). Mapping of detected fragments to peptide sequences was performed by the implemented search engine Andromeda (Cox et al., 2011) using a self-made database of combined P. aeruginosa strain PA14 annotated entries of UniProt database and Pseudomonas Genome Database (Winsor et al., 2011; UniProt, 2014). The resulting txt-file containing the identified proteins was further processed by spreadsheet application Excel (Microsoft). For more detailed information see Materials and Methods S1.

Comparative transcriptome analysis

RNA samples were extracted from the same P. aeruginosa cultures as the samples for proteome analysis. RNA was extracted from cell pellets using the RNeasy Mini Kit (Qiagen) in combination with Qiashredder columns (Qiagen) according to the manufacturer’s instruction with some modifications. For removal of ribosomal RNA the Ribo-Zero Bacteria Kit (Illumina) was used and cDNA libraries were generated with the ScriptSeq v2 Kit (Illumina). The samples were sequenced in single end mode on an Illumina HiSeq 2500 device involving 50 cycles. Mapping was performed using stampy (Lunter and Goodson, 2011) and testing for differential expression was performed with the R package DESeq (Anders and Huber, 2010). For transcriptome analysis only two biological replicates for each strain were used. Adjusted p-values for differential gene expression were calculated as described by Anders and Huber (Anders and Huber, 2010).
Analysis of readthrough events in \textit{tnprmC} cells

In order to identify readthrough events upon \textit{prmC} deletion, raw files of the proteome analysis were reanalyzed by MaxQuant. Therefore, multiplicity was set to 1. Instead of the existing PA14 database containing known coding sequences, three new databases were prepared containing the in frame, +1 frameshift and +2 frameshift protein sequences starting from the amino acid after the first lysine downstream of the stop codon to at least 50 amino acids downstream of the stop codon plus the sequence to the next lysine of the translated non-coding region. Only peptide sequences without stop codons and only unlabeled peptides were considered. For comparison of \textit{tnprmC} and \textit{tnprmC::prmC}, only proteins with detected peptides with intensities in at least two of the three \textit{tnprmC} replicates and no intensities in all of the \textit{tnprmC::prmC} replicates were considered.

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References


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Figure 1. SILAC “spike in” experiment for global protein quantification in *P. aeruginosa*. For quantitative proteome analysis the cultures of the PA14 tnprmC transposon mutant either complemented with pUCP20::EV (tnprmC) or pUCP20::prmC (tnprmC::prmC) were mixed 1:1 with the lys8 labeled PA14 tnlDC standard (tnlDC). Subsequently cells were lysed, proteins fractionized by SDS-PAGE and in gel digested by LysC. LC-MS/MS analysis was performed and resulting raw data were processed by MaxQuant (Cox and Mann, 2008). To generate tnpC to tnpC::prmC ratios, the individual ratios to the “spike in” standard were combined.
Figure 2. Volcano plot depicting PrmC-dependent changes of protein abundance in PA14. Squares represent proteins which are significantly lower abundant in the PA14 \textit{tnprmC} transposon mutant complemented with pUCP20::EV (\textit{tnprmC}) compared to PA14 \textit{tnprmC} complemented with pUCP20::\textit{prmC} (\textit{tnprmC::prmC}, Student’s t-test \textit{p} \leq 0.05) with a ratio (\textit{tnprmC}/\textit{prmC::prmC}) \leq 0.83. Triangles represent proteins which are significantly higher abundant in the \textit{tnprmC} mutant with a ratio \geq 1.2. Circles represent proteins which have either no differential abundance (ratio <1.2; ratio >0.83) or exhibit no significant change (\textit{p} >0.05) in abundance. \textit{n}=3 for this experiment.
Figure 3. Correlation of PrmC-dependent protein to mRNA ratios. The protein (P) to mRNA (T) ratios of the PA14 tnpC transposon mutant either complemented with pUCP20::EV (tnpC) or pUCP20::prmC (tnpC::prmC) were calculated and plotted against each other. Most of the protein to mRNA ratios did not significantly differ between the two strains, indicating that the ratios were independent of the presence of PrmC. PrmC-dependent protein to mRNA ratios were calculated by dividing protein ratios (tnpC/tnpC::prmC) by mRNA ratios. Squares represent the 100 proteins exhibiting the lowest values for the ratio between protein to mRNA ratios (top100 low, S3 Table). Diamonds represent 100 proteins exhibiting the highest values for the ratio between protein to mRNA ratios (top100 high, S4 Table). R^2 indicates squared Pearson’s correlation coefficient.
Figure 4. Complementation of the \textit{tnprmC} mutant by DNR. (A) For monitoring PrmC and DNR dependent anaerobic growth behavior in Pyg medium, OD$_{600}$ was measured. Anaerobic cultures were inoculated by PA14 wt or \textit{tnprmC} strains complemented either with pUCP20::EV or pUCP20::\textit{dnr}. For negative control the wt strain was cultivated without KNO$_3$; n=3 for the experiment. Data points are presented as mean ±SD. The * indicates significantly altered values between \textit{tnprmC::EV} and \textit{tnprmC::dnr} with a p-value ≤ 0.05 as determined by Student’s t-test. (B) To test the PrmC and DNR dependent pathogenicity of PA14 the \textit{Galleria mellonella} virulence assay (Jander et al., 2000) was applied with minor changes. Larvae were infected with 50 cells of PA14 wt, \textit{tnpqsa} or \textit{tnprmC} complemented either with pUCP20::EV or pUCP20::\textit{dnr} or PBS as negative control. Survival rates of 30 larvae per strain (10 per replicate, n=3) were determined by video monitoring. Mortality time point was asserted by melanization of the cuticle and lack of movement for at least 20 min. Significance was determined by Kaplan Meier estimation (Kaplan and Meier, 1958).
Figure 5. Stop codon usage is responsible for PrmC-dependent changes in the translation efficiency.

An enrichment analysis for the three stop codons UGA, UAA and UAG was performed in the top100 low and the top100 high groups (as defined in Fig. 3) in comparison to all genes quantified on the protein level as the control. The * indicates significantly altered values with a p-value \( \leq 0.05 \) as determined by hypergeometric distribution.