

Pseudomonas aeruginosa partitioning protein ParB acts as a nucleoid-associated protein binding to multiple copies of a *parS*-related motif

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ABSTRACT

ParA and ParB homologs are involved in accurate chromosome segregation in bacteria. ParBs participate in the separation of *ori* domains by binding to *parS* palindromes, mainly localized close to *oriC*. In *Pseudomonas aeruginosa* neither ParB deficiency nor modification of all 10 *parS*s is lethal. However, such mutants show not only defects in chromosome segregation but also growth retardation and motility dysfunctions. Moreover, a lack of *parB* alters expression of over 1000 genes, suggesting that ParB could interact with the chromosome outside its canonical *parS* targets. Here, we show that indeed ParB binds specifically to hundreds of sites in the genome. CHIP-seq analysis revealed 420 ParB-associated regions in wild-type strain and around 1000 in a ParB-overproducing strain and in various *parS* mutants. The vast majority of the ParB-enriched loci contained a heptanucleotide motif corresponding to one arm of the *parS* palindrome. All previously postulated *parS*s, except *parS5*, interacted with ParB *in vivo*. Whereas the ParB binding to the four *parS* sites closest to *oriC*, *parS1-4*, is involved in chromosome segregation, its genome-wide interactions with hundreds of *parS* half-sites could affect chromosome topology, compaction and gene expression, thus allowing *P. aeruginosa* ParB to be classified as a nucleoid-associated protein.

INTRODUCTION

The segregation of the bacterial genomes is a precise, complex and still only partially understood process. Most what is known about chromosome segregation is based on studies on low-copy-number plasmids in which partitioning systems have been identified (1,2). Despite their diversity, all

these systems contain two protein components: A, an NT-Pase securing energy supply, and B, a DNA binding protein, and the third vital element, a specific DNA motif designated centromere-like sequence (*parC/parS*), recognized and bound by the B component (2–4). The discovery of a *parA-parB* operon located close to origin of replication (*oriC*) and encoding homologs of plasmid partitioning proteins of class IA in the majority of the bacterial chromosomes sequenced has suggested its potential role in chromosome segregation (5–8).

The chromosomal ParAs are Walker-type adenosine triphosphatases lacking the specific DNA binding domain present in the N-terminal part of the plasmid ParAs (9,10). The chromosomal ParBs, similarly to their plasmidic counterparts, contain an extended helix-turn-helix (HTH) motif, a C-terminal dimerization domain and an N-terminal polymerization domain (4,11–17) and have an ability to spread around the *parS* sequences (17–20) and silence nearby genes, at least in test plasmids (14). The chromosomal Par homologs, despite their high conservation, exhibit various features in different species. Deletion of *parA-parB* is lethal in some species, e.g. *Caulobacter crescentus* and *Mycobacterium xanthus* (21,22), whereas in other species *par* mutants are viable but show various severity defects of chromosome segregation in the vegetative and/or sporulation phase of growth, e.g. in *Bacillus subtilis*, *Streptomyces coelicolor*, *Mycobacterium smegmatis*, *Vibrio cholerae*, *Streptococcus pneumoniae*, *Corynebacterium glutamicum* and *Pseudomonas aeruginosa* (23–37).

Chromosomal ParBs recognize multiple (from 2 to 20) *parS* sequences, highly conserved across the bacterial kingdom and localized mainly in the chromosomal *ori* domains (defined as 20% around *oriC*) (8,11,38). They form large nucleoprotein complexes organizing newly replicated *ori* domains and, with the help of ParAs, separate these domains and hold them in defined locations near the poles until the end of replication and cytokinesis (24,30,39–45). It has been shown that in *B. subtilis* and *S. pneumoniae* ParB-bound

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parS sites act as loading platforms for structural maintenance of chromosomes (SMC) complexes facilitating condensation of ori domains and subsequently of all newly replicated DNA (33,46–50). Two major hypotheses of how the ParBs of Class IA build the large nucleoprotein complexes around the *parS* sequences have been put forward. Both are based on the data showing that ParBs bind to the palindromic *parS* sequences as dimers but can also spread laterally along DNA due to dimer–dimer interactions and weak interactions with non-specific DNA (20,51–53). In the looping and bridging model, ParB dimers bound to DNA interact at a distance, which leads to further DNA compaction (20,54). In the nucleation and caging model, ParB bound to *parS* acts as a nucleation center in which protein–protein and protein–nsDNA interactions spatially entrap additional ParB molecules thus forming a large nucleoprotein complex (52).

Besides their involvement in genome segregation, chromosomal ParAs and ParBs have also been shown to influence bacterial growth and cell cycle by affecting DNA replication and cell division, competence or motility (26,27,32,35,36,39,40,55).

In addition to their main function in plasmid partitioning, plasmidic ParBs of class IA may act as global transcriptional regulators by binding to their operators, spreading and non-discriminately silencing nearby promoters, e.g. in the IncP-1 plasmids (56,57) or IncU plasmids (58). In contrast, oligomers of chromosomal ParBs spread by up to 20 kb from the *parS*s (18–20,54) under native conditions but seem to affect gene expression very selectively. A comparison of the transcriptomes of *B. subtilis* wild-type (WT) strain and *spo0J*_{null} mutant (Spo0J is a homolog of ParB) did not reveal significant changes in the expression of genes located around 10 *parS* sites (19). Transcription of only two out of twenty genes close to *oriC* and a few outside a cluster of three *parS* sites was affected in a Δ *parB1* mutant of *V. cholerae* (59). A similar limited regulation was also observed in *S. pneumoniae* (55) for the competence operon *com*, one of the operons localized in proximity of a *parS* cluster.

Pseudomonas aeruginosa is exceptional in this context, since deletion of the *parB* gene leads to global transcriptional changes affecting more than 1000 genes (60). It remains elusive how ParB of *P. aeruginosa* affects the expression of so many genes. Ten potential 16-bp palindromic *parS* sequences have been identified *in silico* in the genome of PAO1 strain of *P. aeruginosa*, numbered clockwise, with four of them (*parS1*–*parS4*) clustering next to *oriC* (14). *In vitro* studies have demonstrated that ParB binds to all ten *parS*s but with different affinity (61). The highest ParB binding affinity was found toward the perfect palindromic sequences *parS2* and *parS3*, slightly lower toward *parS1* and *parS4* with one mismatch, and even lower toward the remaining six *parS*s with two mismatches. *parS5* was the poorest binder of ParB *in vitro*. An analysis of our collection of single and multiple *parS* mutants has demonstrated that the presence of at least one high affinity site from the *parS1*–*parS4* cluster is necessary and sufficient for proper chromosome segregation (61). The remaining six sites were dispensable for chromosome segregation (61,62).

Here, we used chromatin (nucleoprotein) immunoprecipitation followed by next generation sequencing (ChIP-seq)

to analyze the ParB–DNA interactions in *P. aeruginosa* WT strain, a ParB-overproducing strain and several *parS* mutants: *parS*_{null} with all 10 *parS* sites modified, *parS1-4* mutant with the four *parS*s of the highest affinity for ParB inactivated and *parS2*⁺ with only one high affinity site left intact in the background of the *parS*_{null} strain. This allowed us to identify a large number of additional, specific ParB-bound sequences. Consequently, we propose that ParB, having hundreds of defined interaction sites in the genome, represents a new subfamily of nucleoid-associated proteins (NAPs).

MATERIALS AND METHODS

Bacterial strains, growth conditions and plasmid manipulations

Pseudomonas aeruginosa PAO1161 (*leu*[−] *r*[−]), a derivative of PAO1, was provided by Dr B.M. Holloway (Monash University, Clayton, Victoria, Australia). This strain lacks the inversion between the *rrn* operons present in the annotated PAO1 genome (63,64). *Pseudomonas aeruginosa* PAO1161 Rif^R (WT) strain (35), PAO1161 Rif^R *parB*_{null} (36), PAO1161 Rif^R *parS*mut15 with four *parS* sequences modified (here referred to as *parS1-4* mutant), PAO1161 Rif^R *parS*mut28 (here referred to as *parS2*⁺ mutant) with nine *parS* sequences modified but with unchanged *parS2*, and PAO1161 Rif^R *parS*_{null} with all 10 *parS*s modified (61) were used in the analysis. A ParB-overproducing strain (designated ParB⁺⁺⁺) was obtained by transformation of WT strain with pKGB9 (*araC-araBADp-parB*). Chloramphenicol was added to 75 μ g ml^{−1} (liquid cultures) or 150 μ g ml^{−1} (solid media) to maintain the plasmid and 0.02% arabinose was added to induce the expression of *parB* (65).

DNA pull-down assay

DNA pull-down analysis was carried out as previously (66) with modifications described in Supplementary Materials and Methods. Biotinylated DNA fragments containing *parS1* to *parS10* were obtained by polymerase chain reaction (PCR) using appropriate pairs of primers (Supplementary Table S1). ParB binding to the DNA fragments was assessed by detecting ParB in the eluate using western blotting with anti-ParB antibodies (36) and mass spectrometry analysis (Mass Spectrometry Laboratory, IBB PAS).

Nucleoprotein immunoprecipitation (ChIP)

ChIP was performed as described previously (67) with modifications indicated in Supplementary Materials and Methods. Exponentially growing cultures in L broth at 37°C (OD₆₀₀ ~0.5) were used. For each biological replicate, two independent 50-ml cultures of each strain were pooled together. The immunoprecipitation was performed with affinity purified rabbit polyclonal anti-ParB antibodies (14).

ChIP-qPCR analysis

The immunoprecipitated DNA was used as a template in quantitative PCR (qPCR) performed with Hot FIREPol EvaGreen qPCR Mix Plus (Solis Biodyne) and primers

listed in Supplementary Table S1 in a Roche LightCycler 480. The efficiency of amplification for all primer pairs was between 1.95 and 2.05. The ChIP-qPCR results are presented as ChIP recovery relative to the input DNA used in the procedure, calculated using the formula $100 \cdot 2^{-(Ct_{\text{input}} - 9.965 - Ct_{\text{ChIP}})}$, where Ct are threshold values calculated using the second derivative method with LightCycler 480 software ver 1.5.1.62. Before analysis the input samples were diluted 1000 \times . Each target sequence was analyzed in samples from at least three biological replicates, each with three technical replicates. Primers amplifying a fragment of *proC* gene were used to assess non-specific (background) DNA recovery in the ChIP samples.

ChIP-seq analysis

Library preparation and sequencing on Illumina HiSeq 4000 were performed at Genomed S.A. (Poland). Adapter sequences were removed using Cutadapt (68). Subsequent bioinformatic analyses were performed using the Galaxy platform (<http://usegalaxy.org/>) (69) and R (<https://www.R-project.org/>). Reads were mapped to the *P. aeruginosa* PAO1 (NC_002516.2) genome using Bowtie2 (70). ParB peaks were identified by comparing combined replicates of IP samples (treatment) for each strain with combined negative control (*parB*_{null}) samples using the callpeak function of MACS2 ver 2.1.1.20160309 (71) with default options for paired-end binary alignment map (BAM) files, and 0.05 as false discovery rate (FDR) cut off. Visualization of data was performed using DeepTools (72), Integrative Genomics Viewer ver 2.3.91 (73) and Sushi (74). Coverage files, normalized to 1 \times sequencing depth (RPGC), were generated without binning and smoothing using the bamCoverage tool (72). For simplicity of presentation, the coverage values for individual nucleotides in biological replicates were averaged. ComputeMatrix and plotHeatmap components of deepTools (72) were used to prepare the heatmaps. Identification of the DNA motifs enriched in the ParB peaks was performed using MEME-ChIP (75). Differential binding analysis was performed using Diffbind (76). The ChIP sequencing data have been deposited at the Gene Expression Omnibus (GEO) under accession number GSE110158.

RESULTS

ParB binding to *parS* sequences *in vitro* and *in vivo*

An earlier bioinformatics analysis using the consensus *parS* sequence of *B. subtilis* (11) with allowance of two mismatches has led to the identification of ten putative ParB binding sites in the *P. aeruginosa* PAO1 genome (14). Our previous *in vitro* studies using electrophoretic mobility shift assay (EMSA) confirmed ParB binding to those sequences and demonstrated the highest affinity of purified His₆-ParB to the *parS1-parS4* sites, and the lowest to *parS5* (61). Since the EMSA was performed with short oligonucleotides and tagged ParB, here we used the DNA pull-down method to analyze the binding of unmodified ParB to different *parS* sequences in their natural context. Biotinylated DNA fragments (350–450 bp) encompassing the *parS* sequences were attached to streptavidin-coupled magnetic beads, incubated with PAO1161 WT cell extracts, and washed. Proteins

bound to the DNA were eluted and subjected to western blot analysis with anti-ParB antibodies as well as to mass spectrometry analysis. ParB was detected in the elution fractions in all samples containing DNA with any *parS* sequence with the exception of *parS5* and *parS8* (Figure 1A). These data indicate that in addition to the *parS1-parS4* sequences, required for accurate DNA segregation (61), ParB protein also binds the predicted genomic *parS* sites: *parS6*, *parS7*, *parS9* and *parS10*.

To verify the ParB binding to these *parS* sequences *in vivo*, ChIP with anti-ParB antibodies followed by quantitative PCR analysis was applied. Since a lack of ParB in *P. aeruginosa* PAO1161 *parB*_{null} mutant produced clear-cut effects under conditions of exponential growth in rich medium at 37°C, but not as severe as during growth in minimal medium (36,77), such milder conditions were used for these experiments. The ChIP procedure was performed with PAO1161 (WT) strain, PAO1161 *parB*_{null} mutant (negative control) and PAO1161/pKGB9 (*araC-araBADp-parB*) strain (ParB⁺⁺⁺) with a 5-fold increased ParB level which does not retard bacterial growth (65). Additionally, three *parS* mutants were included: *parS*_{null} with all ten *parS* sites mutated, *parS1-4* mutant with the four high affinity ParB sites inactivated, and *parS2*⁺ strain with nine *parS* sequences modified and only *parS2* left intact (61). Despite producing native amounts of ParB, both *parS1-4* and *parS*_{null} are defective in chromosome segregation to a similar extent as the *parB*_{null} mutant (36). The rationale behind using these strains was that it could help to detect ParB binding not only to the known *parSs* but also to other sites, by either increasing the cellular ParB level or by redistribution of ParB from the main binding sites involved in chromosome segregation, as shown previously for a *parSΔ6* mutant of *B. subtilis* (19).

DNA recovered after ChIP as well as corresponding input DNA was used in quantitative real-time PCR with primers amplifying fragments encompassing *parS1* to *parS10* sequences and a fragment of *proC* gene as a background control (Supplementary Table S1). In WT samples strong ParB binding to *parS1* to *parS4* was confirmed as manifested by high ChIP recovery (Figure 1B). Additionally, a statistically significant enrichment of sequences encompassing *parS6*, *parS7*, *parS8*, *parS9* and *parS10*, but not *parS5*, was found (Figure 1B, inset), indicating that under the growth conditions tested ParB did bind *in vivo* to all the *parS* sequences with the exception of *parS5*. Also in the ParB⁺⁺⁺ strain ParB was bound to all the *parSs* but *parS5* (Figure 1C). Interestingly, while the regions encompassing *parS1* to *parS4* were recovered with a similar efficiency as in the WT, the remaining *parS* sites were significantly more enriched than in WT (Figure 1B and C). As expected, in the *parS2*⁺ strain, the region encompassing *parS2* showed the highest enrichment (Figure 1D), but also the DNA region around *parS1*, adjacent to *parS2*, was significantly enriched, suggesting that ParB bound to *parS2* spread to adjacent DNA sequences. In the *parS1-4* mutant the fragments containing *parS1* to *parS4* were not significantly enriched (Figure 1E) but those encompassing *parS6* to *parS10* were recovered with a higher efficiency than for the WT strain. Finally, no ParB binding to the altered *parS* sequences was found in the *parS*_{null} strain (Figure 1F), and for the *parB*_{null}

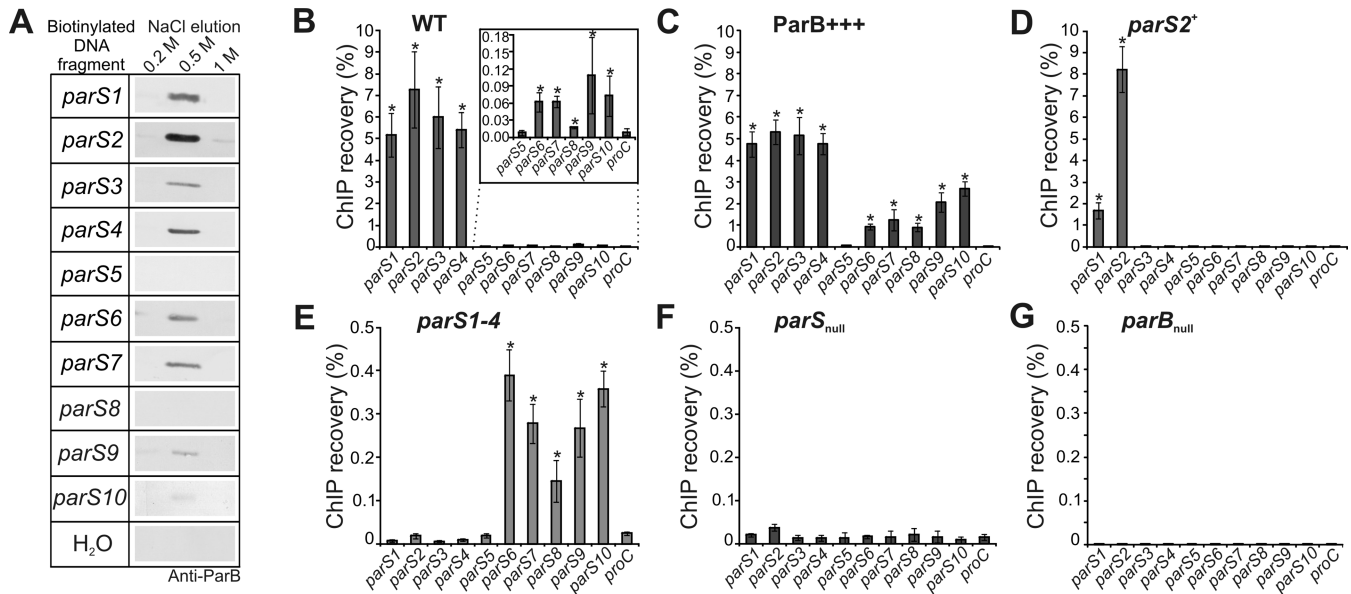


Figure 1. ParB binding to *parS* sites in *Pseudomonas aeruginosa* genome. (A) Western blot analysis of elution fractions from DNA pull-down assays performed with DNA fragments containing *parS1*–*parS10* sequences. Biotinylated DNA was coupled with magnetic beads, incubated with extracts of PAO1161 cells and washed. Proteins bound to the DNA were eluted with increasing concentration of NaCl (0.2–1 M). The negative control contained water instead of biotinylated DNA. Western blotting was performed using polyclonal anti-ParB antibodies. (B–G) qPCR analysis of DNA obtained by nucleoprotein immunoprecipitation using anti-ParB antibodies and extracts from exponentially growing cells of *P. aeruginosa* strains (B) WT, (C) ParB⁺⁺⁺, (D) *parS2*⁺, (E) *parS1-4*, (F) *parS*_{null} and (G) *parB*_{null} (negative control). qPCR was performed using primers flanking the *parS* sequences and for the *proC* gene (background control). Data represent percentage of DNA recovered by ChIP relative to corresponding input samples and are shown as mean \pm SD for at least three biological replicates analyzed in three technical replicates. The significance of differences in recovery of *parS*s relative to the background control (*proC*) was evaluated by two-sided Student's *t*-test assuming equal variance, with $P < 0.01$ considered as significant (marked by asterisk).

strain (negative IP) the recovery was less than 0.01% for all the sequences analyzed (Figure 1G).

Taken together, these results confirm that the *parS1* to *parS4* sequences are the major ParB binding sites in *P. aeruginosa* cells and that they are saturated by ParB in the WT strain. The fact that the lower-affinity sites *parS6* to *parS10* were also enriched, and that this enrichment was higher in ParB-overexpressing cells and also in cells lacking functional *parS1*–*parS4* sequences, suggests that when a ParB pool not bound to its major targets appears, it may specifically associate with other sites of the bacterial chromosome.

Genome-wide identification of ParB-enriched regions

To gain a global insight into the ParB interactions with the *P. aeruginosa* chromosome, chromatin immunoprecipitation was combined with next-generation sequencing. The anti-ParB immunoprecipitated DNA from two independent biological samples of each: WT, ParB⁺⁺⁺, *parS2*⁺, *parS1-4*, *parS*_{null} and *parB*_{null} cells was sequenced and the reads were mapped to the *P. aeruginosa* PAO1 (NC_002516.2) genome. PlotFingerprint analysis indicated the strongest local ChIP enrichment (a large part of the reads mapped to a few genomic bins) for the WT strain and a more dispersed distribution of the ParB-bound DNA regions in strains lacking functional *parS1* to *parS4* sequences and in the strain overproducing ParB (Supplementary Figure S1A). Some enrichment in specific regions of the genome was observed for the negative control *parB*_{null} strain, which

was likely due to cross-reactions of the polyclonal antibodies used.

A principal component analysis (PCA) on the mapped reads obtained for the six strains produced three clusters (Figure 2A). WT, ParB⁺⁺⁺ and *parS2*⁺ formed one group, *parS1-4* and *parS*_{null} formed a second group, and the *parB*_{null} (negative control) strain was clearly separated from those two. Similar grouping was obtained by hierarchical clustering analysis based on correlation coefficients between different samples (Supplementary Figure S1B). Additionally, inspection of the distribution of the sequence reads along the reference genome showed that the reads from the five strains expressing ParB mapped roughly to the same regions which were different from the regions overrepresented in the *parB*_{null} strain (Supplementary Figure S1C). Overall, these analyses confirmed that the obtained reads were enriched in sequences specifically bound by ParB.

To identify the sequences corresponding to the ParB binding sites we performed peak calling on merged data for the two biological replicates what was justified by the high correlation of coverage data within each pair for a given strain (Supplementary Figure S1B and C). Using the FDR cut off value of 0.05, 420 specific peaks were found for the WT strain, 968 peaks for ParB⁺⁺⁺, 823 for *parS2*⁺, 1043 for *parS1-4* and 1077 peaks for *parS*_{null} (Supplementary Tables S2–6). The peaks identified in the WT displayed fold enrichment (FE) between 1.15 and 67.5, with a median of 2.9 (Figure 2B). A similar median FE (3–3.1) was observed for all the other strains analyzed (Figure 2B). Remarkably, not only the number of peaks but also the number of peaks with

FE > 10 was substantially higher in the ParB+++ strain as well as in the strains with the high-affinity *parS* sites impaired (Figure 2C), indicating feasibility of the approach used for the identification of additional ParB binding sites.

The lengths of the sequences encompassed by the peaks varied greatly between 0.2 kb and about 50 kb in WT, ParB+++ and *parS2+* samples (Figure 2D). In *parS1-4* and *parSnull* the broadest peaks reached only 8 and 5 kb, respectively. Nevertheless, the median values of the peak width did not differ substantially among the strains (Figure 2D).

The peaks obtained for the five strains were matched based on the proximity of the peak summits (Supplementary Table S7). Approximately 90% of the corresponding peaks in various strains had their summits <20 bp apart. Overall, the analysis yielded 1305 groups of peaks representing regions with increased ParB occupancy (Supplementary Table S7, hereafter referred to as ParB-bound regions). Of those, 324 were common to all five strains, representing a high-confidence set of ParB-bound regions in the *P. aeruginosa* genome (Figure 2E and Supplementary Table S7). Three other most numerous pools consisted of: 375 sites identified in four strains (ParB+++ and the three *parS* mutants), 153 in three strains (ParB+++ , *parS1-4* and *parSnull*), and 112 sites common to *parS1-4* and *parSnull*. Less than 10% of the peaks were unique to any single strain.

The ParB-bound regions generally displayed an increased read coverage in their central parts (Figure 2F and G). The 420 regions identified in WT strain (Figure 2F, top) were also occupied by ParB in all the other strains, with the exception of those harboring *parS* sequences that were inactivated in the respective *parS* mutants. Importantly, no increased coverage was visible for *parBnull* or WT input. A similar inspection of ChIP-seq data for the remaining 885 ParB-bound regions detected only in ParB+++ , *parS2+* , *parS1-4* or *parSnull* revealed that most of these regions also displayed a slight increase of coverage in WT (Figure 2F, bottom and G), suggesting that ParB could also bind to these sites in WT cells. Overall, this data indicated the existence of more than 1000 specific ParB-bound regions in the *P. aeruginosa* genome. The high reproducibility of these regions in different *P. aeruginosa* strains implies robust and specific binding of ParB to many more sites beyond the canonical *parS* sites.

Inspection of ParB ChIP-seq peaks in various strains

The distribution of the ParB ChIP-seq peaks in various strains was visualized on the *P. aeruginosa* PAO1 genome (Figure 3A). For clarity, only those with FE > 3 were considered. The ParB peaks were present across the entire genome, but the peaks with the highest FE (marked with dots) were almost exclusively found in the genome half containing *oriC*. Notably, the broadest peaks (>5 kb), almost exclusively found in only three strains, WT, ParB+++ and *parS2+*, were tightly clustered in less than ca. 8% of the genome around *oriC* (triangles in Figure 3A). In the *parS1-4* mutant only two peaks broader than 5 kb were detected. Interestingly, these peaks encompass *parS6* and *parS10*.

The 20 peaks with FE > 10 identified in the WT (Figure 2C) were analyzed in detail in the four other strains (Table 1). The cluster of four high affinity *parS* sites (*parS1-*

parS4) in the WT strain produced an extended peak of 53 kb with the maximum FE of 67 (Figure 3B). The corresponding 50-kb peak in ParB+++ strain was enriched only 20-fold. In the *parS2+* mutant a peak with the summit at *parS2* encompassed 21 kb and had an FE of 33. No such broad peaks were detected in the *parS1-4* or *parSnull* strains but the absence of ParB binding to the *parS1-parS4* cluster in these mutants unmasked an additional 15 or 16 peaks, respectively, one of them upstream of *dnaA* with an FE of ca. 19. The role of the ParB binding in the expression of *dnaA* operon requires further studies.

The remaining 19 highest peaks detected in WT strain were also present in the four other strains and had the FE > 10. Interestingly, in the three *parS* mutants the accumulation of ParB at these 19 loci was much higher than in WT or ParB+++ (Table 1). Notably, none of these 19 ParB-associated peaks corresponded to known *parS* sequences. Nevertheless, the *parS6* to *parS10* sites (but not *parS5*) were ParB-bound in three strains, with 3- to 4-fold higher ParB occupancy in ParB+++ and *parS1-4* strains compared to WT (Table 1 and Figure 3C).

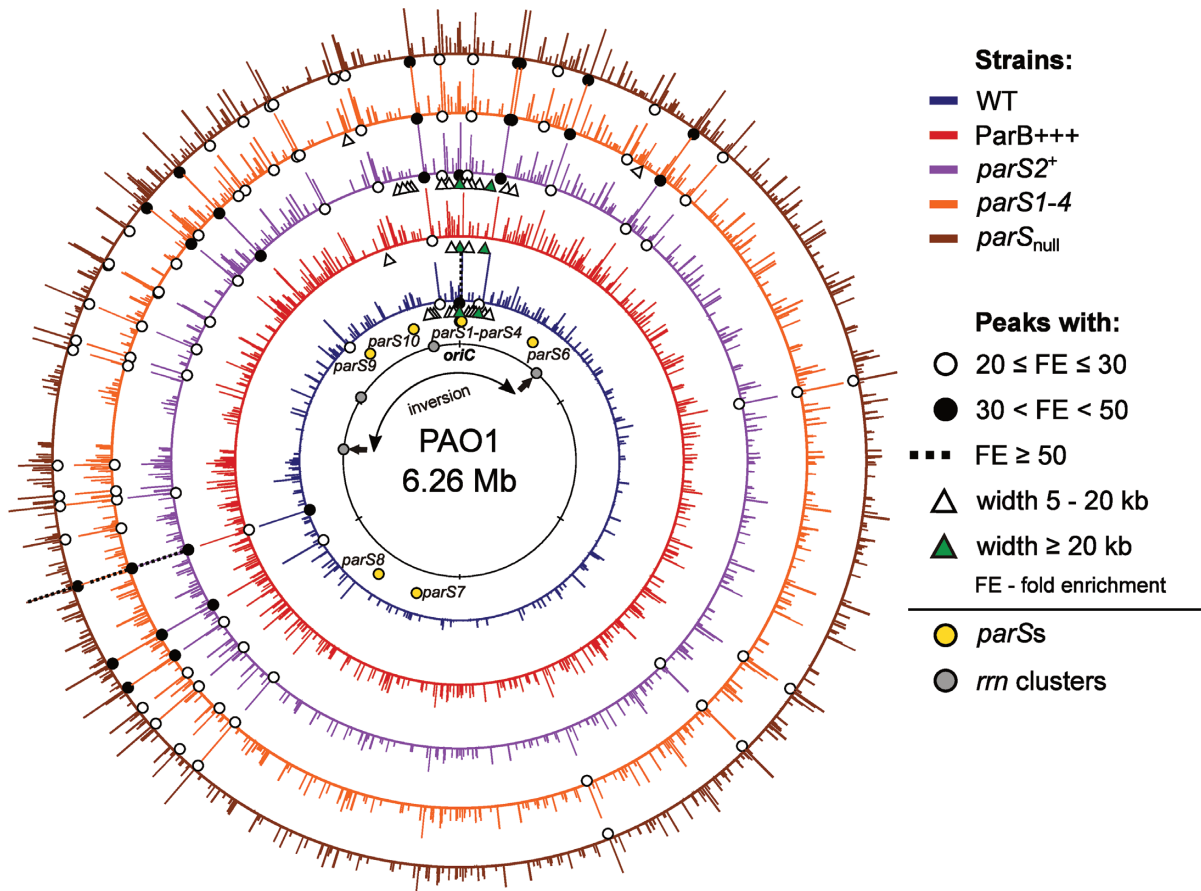
The width of the *parS*-encompassing peaks varied for different *parS*s and also depended on the relative availability of ParB. The *parS6*- and *parS10*-containing peaks encompassed ca. 2 kb each in WT, but were ca. 6 kb wide when an excess of ParB was available, i.e. in ParB+++ and *parS1-4*. ParB bound to *parS7*, *parS8* and *parS9* occupied ~400 bp in WT and under conditions of ParB overabundance the ParB-bound region extended significantly (~3 kb) only for *parS7*. These data suggest that ParB bound to a *parS* site can spread for variable distances depending on the individual *parS* sequence and possibly its context, and the availability of ParB.

Overall, the ChIP-seq analysis demonstrates that *in vivo* ParB binds not only to the *parS* sequences identified previously, with the exception of *parS5* (Figure 3B and C; Table 1), but also to additional regions across the entire genome creating a specific, highly reproducible pattern of distribution (Figure 3A).

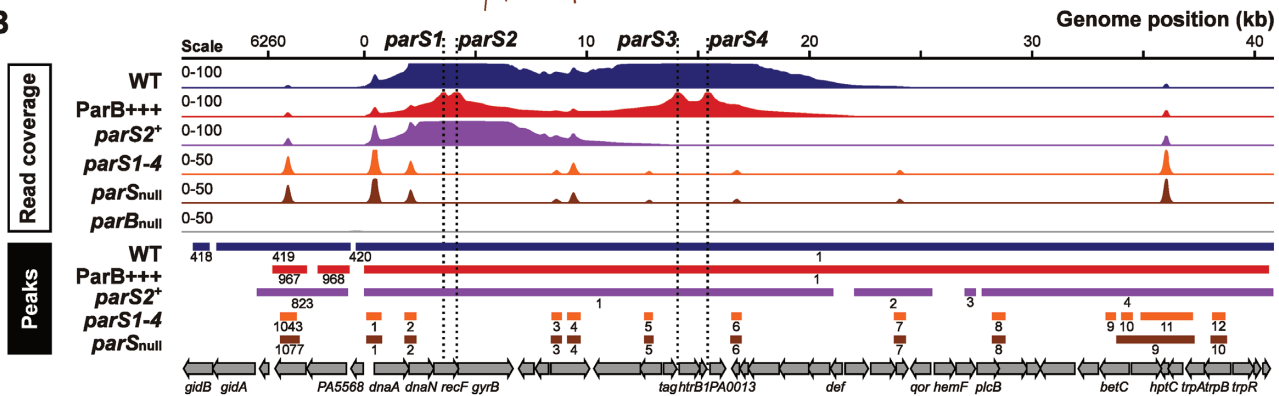
DNA motif other than palindromic *parS* is bound by ParB in *P. aeruginosa* genome

To check if the genome-wide ParB binding involved a defined DNA motif, we performed a search of DNA sequences flanking the summits (± 75 bp) of the peaks using MEME ChIP (75). Remarkably, a similar 7-bp DNA sequence logo was identified in the data from all five strains (Figure 4A). The GtTcCac motif showed strict conservation of positions 1, 3, 5 and 6 whereas at the three other positions some freedom (T→C/A at position 2, C→T at positions 4 and 7) was allowed (Figure 4A). Notably, this motif resembles the GTTCCAC sequence, which is a conserved part of the 8-bp inverted arms of the four major *parS*s critical for chromosome segregation, *parS1-parS4* (Figure 4B), and is also present in one arm of the remaining *parS* sequences, with the exception of *parS5*, the only *parS* to which ParB binding was not found (Figure 1). In *parS6-parS10* the other arm carries substitutions at positions 1, 3 and/or 6, the positions strictly conserved in the deduced motif (Figure 4B),

A



B



C

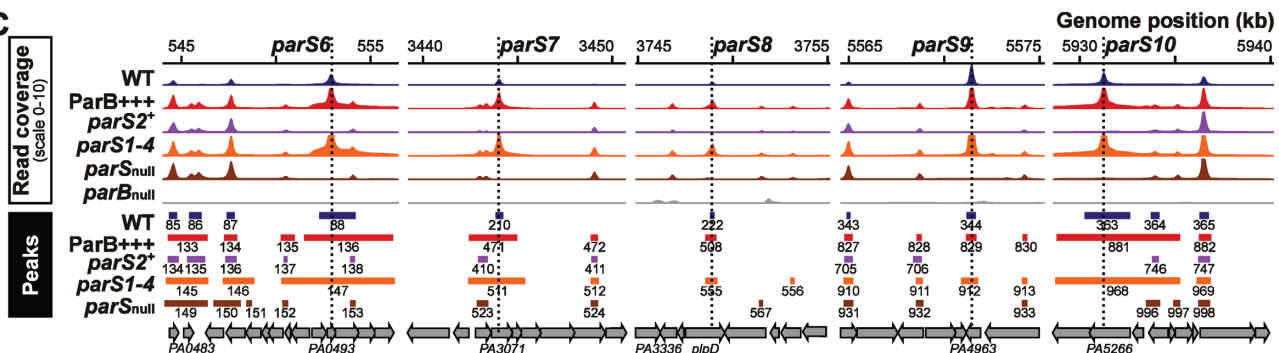


Figure 3. Localization and characteristics of ParB ChIP-seq peaks. (A) Distribution of peaks detected in five analyzed strains. Bar height corresponds to peak FE. Only peaks with FE > 3 are shown and all peaks with FE ≥ 30 are shown at the same height. Genomic coordinates are according to the PAO1 genome. The *rrnA*–*rrnD* gene clusters are marked with gray dots, arrows indicate *rrnA* and *rrnB* loci where inversion occurred in PAO1 (63,64) but not in PAO1161 strain. (B) Coverage of the genome regions encompassing *parS1*–*parS4* cluster and (C) *parS6*–*parS10* sequences in each strain. The histograms show normalized read counts (averaged for two biological replicates) for corresponding positions of the PAO1 genome. Ranges of peaks, their numbers as in Supplementary Tables S2–6 and annotated genomic features are indicated below.

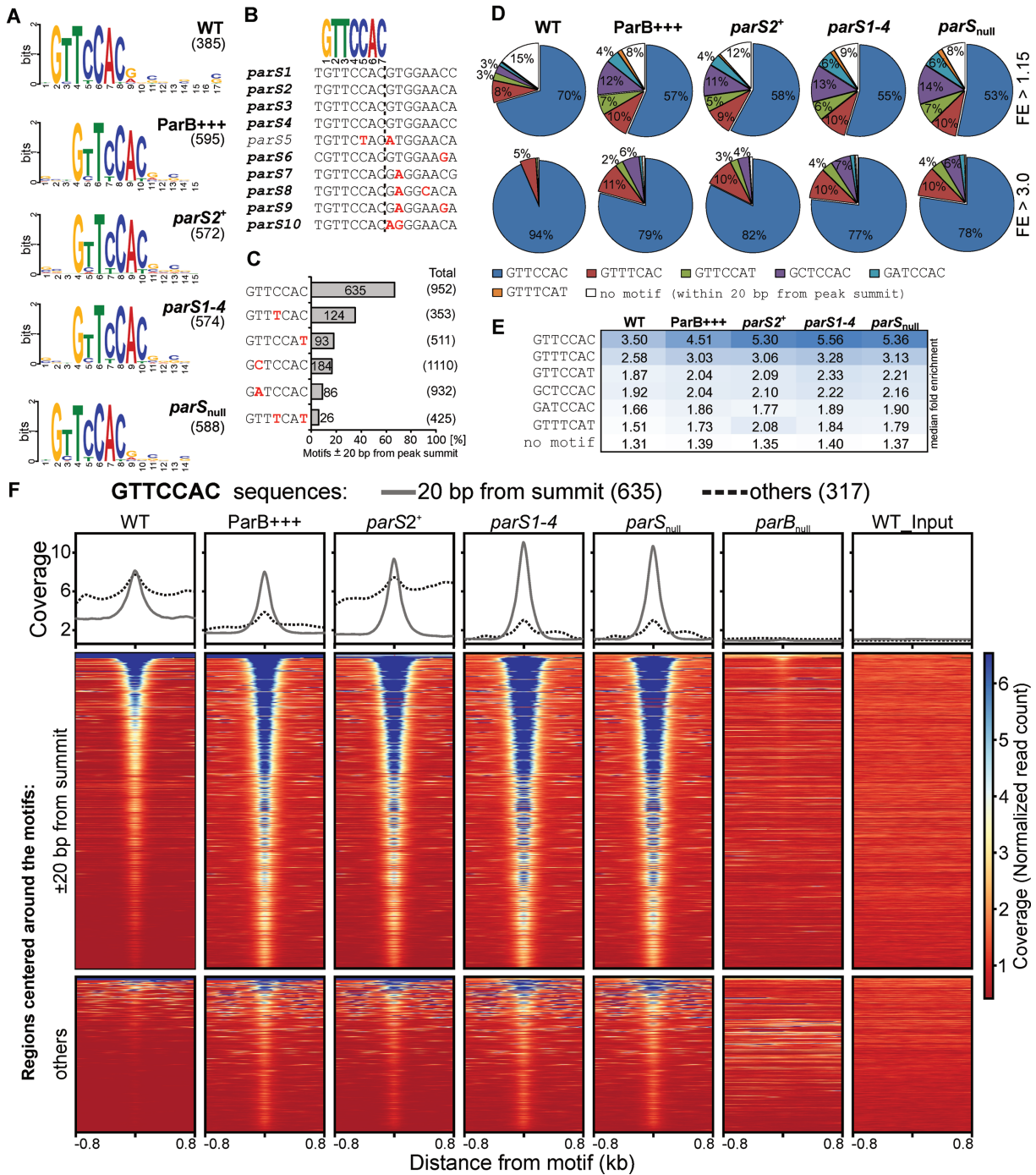


Figure 4. Recurrent sequence motifs in ParB-bound regions of *Pseudomonas aeruginosa* genome. (A) Sequence logos of DNA motifs enriched in genome sequences ± 75 bp from summits of ParB ChIP-seq peaks identified in indicated strains. Motif search was performed using MEME-ChIP. The number on the right indicates the total amount of sequences used to build the logo. (B) Alignment of *parS* sequences from the *P. aeruginosa* PAO1 genome. Logo represents fully conserved sequence in both arms of *parS1*–*parS4* palindromes and in one arm of the remaining *parS* sequences, with the exception of *parS5*. Mismatches relative to GTTCCAC are indicated in red. (C) GTTCCAC sequence variants enriched around summits of ParB ChIP-seq peaks. The percentage of the sequences found within ± 20 bp of any peak summit relative to the total number of copies in the PAO1 genome is shown. Only motifs enriched more than four times are presented. Data for remaining variants are included in Supplementary Table S8. Mismatches relative to GTTCCAC are indicated in red. (D) Percentage of peaks containing the indicated heptanucleotide motifs as closest to the summit. The analysis was performed for all peaks (upper panel) and for peaks with FE > 3 (bottom panel). Category ‘no motif’ includes peaks with none of the analyzed variants present ± 20 bp from summit. (E) Median FE of ParB ChIP-seq peaks carrying indicated variant of the heptanucleotide motif as closest to the summit in different strains. (F) Read coverage around GTTCCAC motifs in the PAO1 genome in ParB ChIP samples from different strains. Each line in the heatmap represents normalized read counts for each nucleotide of region ± 800 bp around a GTTCCAC motif sorted in descending order of mean coverage value, averaged for two biological replicates and colored according to scale. Upper part corresponds to fragments with GTTCCAC motif identified within 20 bp from ParB peaks summit, bottom part to fragments with remaining GTTCCAC motifs. The same regions analyzed in *parBnull* and WT input are included as controls. Plots above the heatmaps indicate mean coverage score separately for the two groups of GTTCCAC motifs.

Table 1. Selected ParB-bound regions in *P. aeruginosa*

Peak Number (WT)	Coordinates in PAO1 genome				Peak fold enrichment (FE)					Locus tag closest	Additional information/ name
	Start	End	Summit	Peak width	WT	ParB+++	<i>parS2</i> ⁺	<i>parS1-4</i>	<i>parS</i> _{null}		
1	14	53 134	14 136	53121	67.5	19.7	33.5	nd	nd	PA0011 (ig)	<i>parS1-4</i>
12	123 814	158 647	147 478	34834	29.8	19.1	40.7	42.7	41.6	PA0129 (ig)	d
69	325 608	326 332	326 024	725	13.8	18.0	28.1	33.2	31.9	PA0290	
94	620 609	621 160	620 843	552	12.5	19.5	29.7	35.8	35.0	PA0564	*1
103	713 611	714 166	713 888	556	10.4	14.3	22.7	26.5	26.6	PA0660	
133	1 366 827	1 367 323	1 367 079	497	10.0	13.1	21.7	26.4	24.3	PA1259	
166	2 354 358	2 354 890	2 354 602	533	10.7	11.9	22.3	26.6	24.0	PA2138	h
225	3 839 047	3 839 623	3 839 369	577	12.9	13.9	24.3	28.7	28.0	PA3430	*3
232	3 987 269	3 987 755	3 987 519	487	11.0	15.5	24.2	29.7	28.0	PA3559	
238	4 099 347	4 099 809	4 099 572	463	11.1	15.0	27.5	32.2	31.1	PA3659	*2
245	4 170 680	4 171 338	4 171 009	659	26.4	19.8	44.4	46.4	45.4	PA3725	g
262	4 381 238	4 381 931	4 381 579	694	31.7	21.8	50.8	52.9	50.8	PA3911	f
279	4 586 068	4 587 049	4 586 456	982	15.7	13.4	27.3	29.1	28.6	PA4102	e
310	5 098 227	5 098 844	5 098 560	618	13.4	13.0	24.4	26.4	26.5	PA4550	a
324	5 378 372	5 378 982	5 378 701	611	12.7	16.2	26.9	31.8	30.8	PA4790	
337	5 496 504	5 497 129	5 496 819	626	12.5	17.0	nd [#]	nd [#]	nd [#]	PA4899	
338	5 498 127	5 499 905	5 498 743	1779	20.9	19.2	38.0	41.5	40.7	PA4901	b
359	5 775 771	5 776 395	5 776 084	625	11.7	15.4	23.1	27.6	27.9	PA5126	
398	6 140 926	6 143 944	6 142 506	3019	23.5	22.6	42.4	46.8	46.6	PA5452	c
412	6 203 326	6 217 050	6 215 975	13725	11.6	16.1	23.8	29.0	27.6	PA5523	
88	552 324	554 217	552 921	1894	5.8	11.1	nd	15.5	nd	PA0493	<i>parS6</i>
210	3 443 861	3 444 275	3 444 056	415	3.6	7.5	nd	11.1	nd	PA3071	<i>parS7</i>
222	3 748 815	3 749 078	3 748 950	264	2.2	4.6	nd	6.2	nd	PA3339	<i>parS8</i>
344	5 571 227	5 571 683	5 571 466	457	7.8	10.9	nd	20.9	nd	PA4963	<i>parS9</i>
363	5 930 304	5 932 676	5 931 318	2373	6.0	11.0	nd	15.1	nd	PA5266	<i>parS10</i>

nd: not detected

nd[#]: not detected as a separate peak but included in the region around PA4901 locus

Shown are parameters of ParB ChIP-seq peaks with FE higher than 10 as determined by MACS2 analysis of ChIP-seq data for *P. aeruginosa* PAO1161 (WT) strain using *parB*_{null} strain as a control. Corresponding peaks in other strains were identified by peak matching (Supplementary Table S7). Data for peaks containing *parS6*–*parS10* sequences are also included. Last column provides names of *parS* sequences and designations of the ParB binding sites identified before (62).

suggesting that these parts of *parS6*–*parS10* might not be involved in initial ParB binding.

To identify the sequence variants which could be recognized by ParB, a search of DNA motifs in the PAO1 genome was performed using the GTTCCAC sequence with the allowance of two mismatches. The total pool of such variants was then limited to the 183 ones detected at least once within 20 bp from any peak summit (Supplementary Table S8). Overall, the sequences in proximity of peak summits analyzed here represent 1.3% of the genome. Remarkably, six variants: GTTCCAC, GTTTCAC, GTTCCAT, GCTCCAC, GATCCAC and GTTTCAT showed at least a 4-fold enrichment in sequences ± 20 bp of peak summits relative to the total genome (Figure 4C and Supplementary Table S8). Among them, GTTCCAC was the most over-represented, since 66.7% of the 952 sequences present in the PAO1 genome were found close to peak summits. The second most enriched motif in proximity of summits was GTTTCAC (35.1%).

An analysis of the peaks in all five strains revealed that $\geq 85\%$ contained at least one GTTCCAC, GTTTCAC, GTTCCAT, GCTCCAC, GATCCAC or GTTTCAT motif within 20 bp from the summit (Figure 4D), with approximately 70% of peaks in each strain containing the motif < 5 bp from the summit (Supplementary Figure S2). GTTCCAC was the most frequently identified variant lo-

calized closest to the summit (Figure 4D, upper panel), followed by GTTTCAC. Remarkably, when the analysis was limited to the peaks with FE > 3 these two variants were identified in 94 and 5% of WT peaks, respectively (Figure 4D, bottom panel). A similar trend was observed for other strains. Strikingly, peaks without any of the six analyzed motifs ± 20 bp from summit represented $< 1\%$ of the peaks with FE > 3 (Figure 4D, bottom panel).

A complementary analysis of the FE of the peaks containing indicated variant as the closest one to the summit gave the highest median FE for peaks with GTTCCAC, followed by those with GTTTCAC (Figure 4E). The median FE for the peaks without motif ± 20 bp from summit was < 1.4 suggesting that these peaks might represent low affinity binding sites (Figure 4E).

To further confirm that the genome fragments carrying the six indicated sequences were enriched by precipitation with anti-ParB antibodies we analyzed normalized read coverage around each motif in the genome, separately for those found ± 20 bp from the summits and for the remaining ones. As expected, the mean normalized read count around GTTCCAC present ± 20 bp from a peak summit revealed a strong central increase in all ParB-expressing strains, but not for *parB*_{null} (negative control) or WT_input DNA (Figure 4F, upper part of the heatmap). A similar analysis for the remaining GTTCCAC motifs also revealed an increase

in read coverage for sequences in proximity of most of these motifs in all five strains (Figure 4F, bottom part of the heatmap). The observations were confirmed by quantification of the coverage data for both groups (Figure 4F, plots). Similar results were obtained for GTTCCAC, GCTCCAC, GTTCCAT, GATCCAC and GTTTCAT motifs (Supplementary Figure S3). Overall, the high enrichment of six sequence variants related to one arm of *parS* in proximity of ChIP-seq peak summits strongly suggests that these are true targets for ParB binding.

To check whether ParB binding to the 7-bp motif is followed by ParB spreading (as observed after binding to palindromic *parS*s) we analyzed the width of peaks containing the motifs. Positive correlation between the peak width and the number of motifs encompassed by the peak (Supplementary Tables S2–6), but not between FE and the number of motifs was observed, revealing that appearance of broad peaks results from the presence of multiple closely located ParB binding sites (Supplementary Figure S4A and B). Concomitantly, analysis of the widths of peaks containing a single *parS* half-site revealed that the vast majority of such peaks spanned <600 bp (Supplementary Figure S4C), twice the average size of the DNA fragments used for ChIP, indicating no obvious ParB spreading around the *parS* half-sites.

ParB bound to *parS1-4* cluster provides a platform for extended ParB–DNA interactions

Most ChIP-seq peaks contained one of the six variants of the GTTCCAC motif <20 bp from the summits (Supplementary Figure S2), but a fraction of the peaks lacked such motifs. We asked therefore how consistent was presence of motif-less peaks among different strains, how strongly enriched regions they represented and whether they occurred in specific locations. Combined ChIP-seq data from the five analyzed strains led to the identification of 1305 ParB-bound regions that were then grouped according to the number of strains they were found in (Figure 2E and Supplementary Table S7). The ParB-bound 324 regions detected in all five strains were considered to be the most reliable ones, whereas those found in the ChIP-seq data of any single strain (221 in total) were considered the least reliable. Presence of any of six variants of the 7-bp motif showed positive correlation with the number of strains in which the region was identified (Figure 5A). Out of the 324 ParB-associated regions common to all strains, 321 (99.1%) contained the motif close to the summit in comparison with only 41% (92 out of 221) found in any single strain. Overall, the motif was identified ± 20 bp from the peak summit in 1097 of the 1305 ParB-bound regions (Figure 5B), and in 59 ParB-bound regions the motifs were also found but more than 20 bp away from the summit (Figure 5B). Only 149 regions lacked the analyzed motifs. Of those, 70% were detected in a single strain only (Supplementary Table S7) and showed low enrichment (median FE 1.26), suggesting that they may represent non-specific ParB binding. Notably, most of the motif-less peaks were detected in the WT, ParB+++ and/or *parS2*⁺ strains, i.e. the strains carrying at least one *parS* from the *parS1-4* cluster (Figure 5B) in which a large nucleoprotein complex formed at *oriC* (Fig-

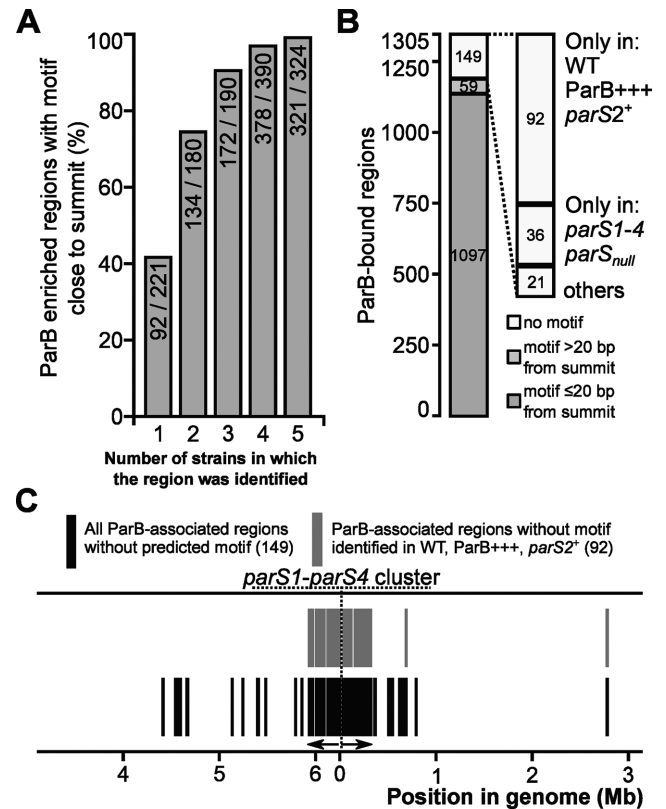


Figure 5. Heptanucleotide motif-independent ParB binding to DNA. (A) Fraction of ParB-associated regions carrying six analyzed variants of heptanucleotide motif ± 20 bp from peak summit as a function of number of strains in which the region was detected. Numbers in columns relate to ParB-bound regions with the motif close to the summit versus summed up regions detected in any single strain (221), any two strains (180), any three (190), any four (390) and all five strains (324). (B) Distribution of heptanucleotide motifs (as above) in ParB-bound regions (Supplementary Table S7). The regions are divided into three categories: with a motif ± 20 bp from summit, with a motif further than ± 20 bp from summit but within the enriched region and with no motif in the enriched region. The motif-less regions were also categorized according to the strains they were detected in. (C) Localization of ParB-bound regions lacking a heptanucleotide motif in the *Pseudomonas aeruginosa* PAO1 chromosome.

ure 3B). Indeed, almost all these motif-less peaks were in the vicinity of *oriC* (Figure 5C) suggesting that the ParB binding to a high affinity site(s) resulted in DNA regions fairly distant from the *parS1-4* cluster and lacking the 7-bp motif being recovered in the anti-ParB immunoprecipitate. Hence the large complex formation at *oriC* involves specific ParB binding to high affinity *parS1-parS4* sites and multiple lower affinity *parS* half-sites (e.g. 39 such motifs are present in the *oriC* encompassing peak in WT strain) as well as non-specific DNA interactions.

To get an insight into the postulated *parS1-parS4* dependent enrichment of certain DNA sequences a differential binding analysis was performed using Diffbind (76). The ChIP-seq data for strains carrying at least one *parS* of the *parS1-parS4* cluster (WT, ParB+++ and *parS2*⁺) was compared pairwise with the data for strains lacking such *parS* site (*parS1-4* and *parS*_{null}). The analysis was not limited to the ChIP-seq peaks, but instead non-overlapping 200-bp windows covering the whole genome were used. The WT

versus *parS1-4* comparison revealed that the majority of bins showing statistically significant difference between the strains displayed decreased coverage in WT (Figure 6A). This was in agreement with the previous observation that the ChIP-seq peaks in the *parS* mutants showed higher FE than the corresponding peaks in the WT (Table 1). Conversely, some bins showed an increased coverage in WT relative to *parS1-4* and these were strictly confined to a 400-kb chromosome region encompassing *oriC* (Figure 6A). The highest differences between the two strains were observed from 10 kb upstream of *parS1* to 20 kb downstream of *parS4* (Figure 6A) with a clear negative correlation between the magnitude of the difference between the strains and the distance from the *parS1-parS4* cluster. This relation was less clear for bins located further away from *parS1-parS4*. Remarkably, multiple bins in the 115 to 150 kb region showed higher coverage difference between WT and *parS1-4* than bins located relatively closer to *parS1-parS4* (Figure 6A and B). Similar results were obtained when WT was compared with *parS_{null}* as well as when *parS2⁺* or ParB+++ were compared with *parS1-4* (Supplementary Figure S5). Significantly increased coverage of sequences from the 115–150 kb region in the WT versus *parS1-4* comparison as well as the presence of a 20-kb gap separating this region from the peak encompassing *parS1-parS4* were also observed when WT_input was used as a background control in Diffbind analysis (Figure 6B), indicating that the observed distribution is not an artefact caused by an elevated number of reads mapping to this part of the genome in *parB_{null}* data, used as a background control in all previous analyses.

Overall, the data indicate that ParB binding to *parS1-parS4* is required for an enrichment of specific 50 kb region around *oriC* but also at quite distant locations (e.g. the 115–150 kb region) from it. The fact that these regions are discontinuous suggests that the enrichment cannot be a sole consequence of lateral spreading of ParB along the DNA adjacent to the *parS1-parS4* cluster but further supports the hypothesis of interactions at a distance involving bridging/looping or caging of DNA (20,52–54,78).

DISCUSSION

Bacterial genomes are highly compact structures whose distribution within cell is tightly controlled spatially and temporally (79,80). Chromosome segregation must be coordinated with DNA replication and cell division to assure delivery of full genome copies to progeny cells. It has been demonstrated that ParA, ParB and *parS* sequences are engaged in genome partitioning in many bacteria (14,21,24–37). Additionally, they have been shown to be involved, in a species-specific manner, in the regulation of DNA replication initiation by DnaA (26,27), in the formation of a loading platform for SMC on newly replicated DNA to assure its proper condensation (33,46,47,50), in the positioning of *oriC* domains at defined cell locations (21,24,36,37,43,44,81), coordination of replication and cell division (30,39,82) and also in the regulation of gene expression (55,59,65,82).

Our earlier studies have confirmed the partitioning role of ParA and ParB proteins in *P. aeruginosa*, identified ten

parS sequences bound by ParB with different affinities and showed that any one of the four *parS* sites closest to *oriC*, *parS1-parS4*, is required and sufficient for accurate chromosome segregation (14,61). Remarkably, expression of over 1000 genes was affected by a lack of ParB (60), suggesting that ParB plays a role in global gene regulation. The plausible explanations of that effect were ParB interactions with sites other than *parS*s and/or an involvement of ParB partners in the regulation. A recent study on ParB in *P. aeruginosa* (62) claimed that *in vivo* ParB bound to the *parS1-parS4* and eleven other regions but not to other *parS*s. In contrast, the results of our *in vitro* (DNA pull-down) and *in vivo* (ChIP) confirmed ParB binding to *parS6-parS10*. To solve this apparent discrepancy regarding the ParB–DNA binding *in vivo* we performed a ChIP-seq analysis using a WT strain, three *parS* mutants as well as a ParB-overproducing strain, with *parB_{null}* serving as a negative control. Unlike in the aforementioned study (62), untagged ParB and polyclonal anti-ParB antibodies were used rather than a ParB-FLAG chimera. During the data analysis we did not apply a strict and, by its nature, arbitrary FE cut off, but used the criterion of statistical significance alone, which allowed us to identify a high number of ChIP-seq peaks. Importantly, more than 90% of the peaks were identified in at least two different strains (Figure 2E) indicating that, despite their often relatively low FE, these peaks represent bona fide ParB-binding regions.

The ChIP-seq analysis showed unequivocally that *in vivo* ParB is bound to all the palindromic *parS* sequences identified previously, except *parS5*, with the highest degree of ParB association in proximity of *oriC* coinciding with the *parS1-parS4* sites (Figure 3B). Mutations in the *parS* palindromes blocked the ParB binding which confirmed that the indicated *parS*s are specifically bound by ParB *in vivo*. Remarkably, our data indicate that ParB occupies 420 regions in WT strain, covering almost 7% of the genome and ca. 1000 regions under conditions of ParB abundance.

Using the stringent enrichment criterion of FE > 10, 20 high-affinity regions were identified in the WT, including the major one encompassing the *parS1-parS4* sites and all 11 ‘secondary ParB binding sites’ detected previously (62) in *P. aeruginosa* strains grown on minimal medium (Table 1). This partial correspondence of the two sets of results indicates a significant degree of specificity and conservation of ParB binding to the genome. However, the drastically different experimental setups and approaches used in data analysis prevent conclusions about growth conditions-dependent ParB binding to DNA.

When the availability of ParB in the cell is limited, as is the case in the native situation, the protein must discriminate between its binding sites. The occupation of individual sites may be determined by the degree of affinity, interactions with cellular partners (e.g. ParA), local ParB availability, on-going DNA transactions (e.g. transcription and replication), presence of other factors interacting with DNA, DNA conformation, physiological state of the cell or growth conditions. An increased ParB availability in the cells enabled the identification of ParB binding to an additional 885 loci not identified in the WT strain. Importantly, most of these loci do show a read coverage slightly above the background value also in WT ChIP-seq data suggesting

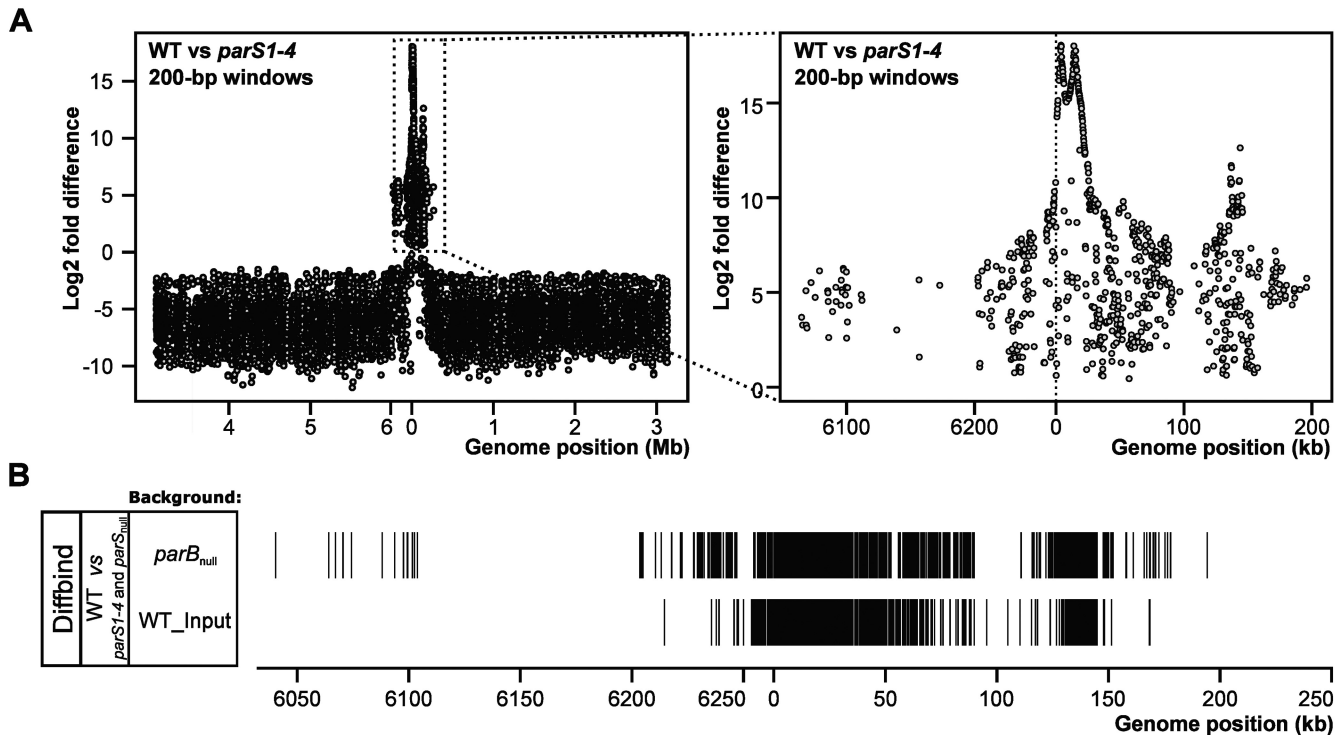


Figure 6. *parS1-parS4* dependent ParB distribution in *Pseudomonas aeruginosa* genome. (A) Comparison of ParB binding to DNA in WT strain versus *parS1-4* strain (lacking functional *parS1-parS4* cluster). Differential binding analysis was performed in non-overlapping 200-bp windows covering the whole genome using Diffbind. Only bins showing statistically significant difference of coverage (FDR of 0.05) between WT and *parS1-4* are shown. On the right, blow up of the *oriC* region with bins showing higher number of reads in WT than in *parS1-4* is shown. (B) Distribution of *parS1-parS4* dependent ParB ChIP enriched sequences in proximity of *oriC* (± 200 kb). Diffbind analysis was performed for WT versus *parS1-4* and *parS_{null}* ChIP-seq data using 200-bp windows. Subsequently the analysis was performed with the 200-bp windows shifted by 100 bp. Data represent windows (or their halves) showing a statistically significantly (FDR ≤ 0.05) higher coverage of reads in WT strain relative to *parS1-4* and *parS_{null}* for both sets of genomic windows using *parB_{null}* or WT_Input data as a background control.

low-occupancy ParB binding in WT cells and not only in the mutants used in this study (Figure 2F). Concomitantly, most of the peaks common to WT and mutant cells showed higher FE in the four strains with increased ParB availability. In a similar study in *B. subtilis* deletion of six out of ten *parS* sites resulted in re-distribution of Spo0J (ParB) toward the low-affinity *parS* sites left in the mutant strain (19), whereas in *P. aeruginosa* the inactivation of the high affinity *parS* sites led to the re-distribution of ParB not only to the remaining *parS*s but also to hundreds of additional ParB binding sites.

The extent of ParB spreading has been analyzed in diverse chromosomal systems. Most data suggested the spreading for up to 10 kb from a *parS* site in natural conditions (19,55,59,82). Recently, ParB from *C. crescentus* has been shown to spread by no more than 2 kb from a single *parS* site, while the extension of the ParB-associated region in the vicinity of *oriC* to 10 kb involved ParB binding to a cluster of four *parS* sequences located in a 5-kb region near *oriC* (83). The *parS1-parS4* cluster located within 15 kb of *oriC* in *P. aeruginosa* seems to facilitate even farther ParB-DNA interactions since the most conspicuous region of ParB spreading encompassed more than 40 kb in the WT strain (Figure 6A). A comparison of the distribution of the ParB-associated DNA in *oriC* region between the WT and *parS1-4* (or *parS_{null}*) revealed a discontinuous

ca. 400 kb stretch around the *oriC* whose occupation was strictly dependent on the presence of *parS1-parS4* (Figure 6). Models explaining how ParB shapes the *ori* domain presumed binding to the strong affinity *parS* site(s), spreading due to the non-specific DNA binding activity and forming a large nucleoprotein complex due to 3D interactions between DNA-bound ParB molecules (20,52–54,78). Showing that except *parS*s, *P. aeruginosa* ParB has several hundreds of other binding sites in the genome we reveal another important mode of ParB action, which allows it not only to shape the *ori* domain but also to influence the global chromosome topology.

Notably, we show that the ParB-bound sequences contain the GtTcCac motif corresponding to seven out of eight nucleotides from a single arm of the canonical *parS* palindrome (Figure 4A and B). A systematic analysis of the motif variants showed that GTTCCAC, GTTTCAC, GCTCCAC, GTTCCAT, GATCCAC and GTTTCAT motifs were found in proximity of ParB ChIP-seq peak summits significantly more often than expected by chance. In any given strain more than 85% of all peaks and 99% of peaks with FE > 3 carried one of the six sequence variants in proximity of the peak summit. Among these six variants, GTTCCAC was not only the most frequently occurring one but was also found closest to the summits of peaks with the highest FE. In the most consistent peaks,

those detected in four or five strains, 97 and 99%, respectively, contained one or more of the six motif variants within 20 bp from the summit (Figure 5A). Remarkably, inspection of the ChIP-seq read coverage around all such motifs in the PAO1 genome showed an increased coverage around them indicating that these motifs were likely targeted by ParB *in vivo*. Our data also demonstrates that the binding to the *parS* half-sites (in contrast to the ParB binding to palindromic *parS*s) does not prompt long distance spreading of ParB.

Recent reports indicate that some ParB proteins do not strictly require a symmetrical *parS* structure for DNA binding. ParB2 of *V. cholerae* was shown to bind a non-centromeric site containing a half-*parS* sequence (82). Additionally, in the multipartite genome bacterium *Burkholderia cenocepacia* ParBc1, ParBc3 and ParBpBC displayed enhanced affinity for the corresponding half-*parS* relative to random-sequence DNA (84). This data show that the ability of ParB to bind to one arm of *parS* is not restricted to *P. aeruginosa*. However, the biological significance of the ParB binding to *parS* half-sites in the genome remains elusive.

We speculate that the half-*parS* sequences scattered across the genome may play a role in concentrating ParB molecules on the nucleoid. The interaction of ParB with multiple sites combined with the interactions between ParB molecules could provide a scaffold for chromosome compaction and affect DNA topology, possibly playing a role in modulation of gene expression. Our earlier transcriptomic analysis of *P. aeruginosa parB_{null}* mutant versus WT indicated altered expression of 1166 genes (60). A large fraction of these genes seemed to be involved in the stress response due to disturbances in chromosome segregation (85). A complementary study with a strain showing 5-fold overproduction of ParB (ParB+++), a ParB excess leading to no visible changes in growth rate or genome segregation, revealed altered expression of 211 loci (65). The present ChIP-seq data suggest that the ParB–DNA binding relies on the presence of specific sequence(s). A comparison of the distribution of the analyzed variants of the *parS* half-site with the transcriptional organization of the genome shows that in PAO1 these sequences are mainly intragenic (GTTCCAC—92%, GTTTCAC—76%, GCTCCAC—89%, GTTCCAT—94%, GATCCAC—93% and GTTTCAT—84%), which suggests their roughly random distribution between coding and non-coding sequences (genes make up 90% of the PAO1 genome). The loci with expression affected by ParB deficiency are putatively expressed from 853 promoters. Only 113 of these promoters contain one of the six indicated sequences. Similarly, the 211 loci affected by ParB overproduction are expressed from 130 promoters only 28 of which contain one of the motifs. Whereas ParB binding to the palindromic *parS3* and *parS4* is responsible for the repression of *PA0011p* and *PA0013p*, respectively (65), the low correlation between the ParB-dependent changes in expression and the presence of half-*parS* sequences in the promoters of other genes suggests that their ParB-dependent regulation does not rely solely on a direct ParB–promoter interaction. The mechanism linking ParB and gene expression regulation in *P. aeruginosa* requires further studies.

Summarizing, the most prominent features of the *P. aeruginosa* partitioning protein ParB are: i) sequence-specific DNA binding to palindromic *parS* sequences, dedicated primarily to the specific task of *oriC* separation and chromosome segregation; ii) specific binding to numerous heptanucleotide sites throughout the genome; iii) acting at a distance, probably by a combination of spreading and bridging/looping/or caging; and iv) widespread influence on gene expression. These features allow ParB to be classified as a NAP involved in DNA compaction, genome topology maintenance and modulation of gene expression (79,86,87).

DATA AVAILABILITY

All sequencing data have been deposited at the Gene Expression Omnibus (GEO) under accession number GSE110158.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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