Crystal structure and centromere binding of the plasmid segregation protein ParB from pCXC100

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ABSTRACT
Plasmid pCXC100 from the Gram-positive bacterium Leifsonia xyli subsp. cynodontis uses a type Ib partition system that includes a centromere region, a Walker-type ATPase ParA and a centromere-binding protein ParB for stable segregation. However, ParB shows no detectable sequence homology to any DNA-binding motif. Here, we study the ParB centromere interaction by structural and biochemical approaches. The crystal structure of the C-terminal DNA-binding domain of ParB at 1.4 Å resolution reveals a dimeric ribbon–helix–helix (RHH) motif, supporting the prevalence of RHH motif in centromere binding. Using hydroxyl radical footprinting and quantitative binding assays, we show that the centromere core comprises nine uninterrupted 9-nt direct repeats that can be successively bound by ParB dimers in a cooperative manner. However, the interaction of ParB with a single subsite requires 18 base pairs covering one immediate repeat as well as two halves of flanking repeats. Through mutagenesis, sequence specificity was determined for each position of an 18-bp subsite. These data suggest an unique centromere recognition mechanism by which the repeat sequence is jointly specified by adjacent ParB dimers bound to an overlapped region.

INTRODUCTION
Stable transmission of duplicated genetic material to daughter cells is essential for all life forms. The active partitioning of low-copy-number plasmids has served as a simple model system for DNA segregation studies (1–5). Plasmid partition systems are composed of only three elements: a cis-acting centromere-like site called parS; a filament-forming NTPase, ParA; and a centromere-binding protein (CBP), ParB. These elements are normally arranged as a par operon, with parB located downstream of parA. The centromere region typically consists of multiple direct or inverted repeats near the par operon. Multiple ParB proteins assemble on the centromere repeats to form a high-order nucleoprotein complex, the partition complex. The partition complex recruits ParA to form the segrosome. ParA can polymerize into filaments in an NTP dependent manner. The dynamics of partition filaments is important to separate the attached plasmids into the opposite poles of dividing cells, but the underlying molecular mechanism remains unclear (1–5).

There are two main plasmid partitioning systems, distinguished from each other on the basis of whether the NTPase is a Walker-type (type I) or actin-type (type II) protein (6). The type I partition system is subdivided into types Ia and Ib according to different structure of ParA and ParB, and the relative position of the centromere (6). The centromere region is located downstream of the par loci in the type Ia system but is located upstream in the type Ib and II systems. Transcription of the par operon is under autoregulation by the Par proteins. Type Ia ParA is larger than type Ib ParA because it has an additional N-terminal helix–turn–helix (HTH) domain that binds to the promoter region of the par locus to repress transcription. In type Ib and II systems, ParB associates with the centromere sequence located upstream of the par operon to repress transcription. Type III partition systems were recently characterized that are composed of a tubulin-like GTPase TubZ and a CBP TubR with a dimeric winged HTH motif (7–11).

Type Ia CBPs have a complex multidomain organization, but they share a conserved HTH DNA-binding motif (12–14). In contrast, type Ib and II CBPs are extremely divergent in their primary sequences, obscuring recognition of their DNA binding domain. The structures of two type Ib CBPs—omega and ParG—have been determined (15–17). Each contains a dimeric ribbon–helix–helix (RHH) DNA-binding domain and a flexible N-terminal
domain. The RHH motif was also found in the structures of type II CBPs pB171 ParR and pSK41 ParR (18,19). Although type Ib and II CBPs specifically interact with Walker- and actin-type NTPase, respectively, they appear to share a common DNA-binding structure. CBPs need to function in the context of the partition complex. A type Ib omega–centromere complex shows an extended structure (17), whereas a type II pSK41 ParR–centromere complex forms a superhelical structure (18). The different higher order structures of the partition complex may impact the recruitment of specific ParA (18).

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In this study, we applied X-ray crystallography to determine the structure of the C-terminal DNA-binding domain of pCXC100 ParB, hereafter referred to as ParB, and revealed that it forms a dimeric RHH fold. We also extensively characterized the interaction of ParB with the centromere by hydroxyl radical footprinting, electrophoretic mobility shift assays (EMSA) and isothermal titration calorimetry (ITC). We have mapped the binding sites of ParB on the centromere and divided the structure of the centromere into nine direct repeats. We also characterized in detail how ParB recognizes single subsites and binds cooperatively to tandem subsites. Interestingly, a ParB dimer binds to an extended 18-bp region of single subsites and recognizes the sequence of the immediate repeat as well as flanking repeats.

### MATERIALS AND METHODS

#### Gene expression and protein purification

The ParB gene (residues 1–139) was PCR-amplified from plasmid YB411 (21) with primers WTF1 (5'-ATATACCA TGCGTGTGACACGGTGG-3'); the restriction site is underlined) and WTR139 (5'-CCGCAACGCTGCTTCC CAGTGCGCGCCCGG-3') and cloned into pET28a using NcoI and HindIII sites. The full-length ParB protein contains a non-cleavable six-His-tag at the C-terminus encoded in the plasmid. ParB fragments spanning residues 60–139, 65–139 and 65–135 were generated with primers PF60 (5'-GGCGGAGCTTCTGGAGCGC-3') and PR139 (5'-CCGGGACTTCTATGGCGGCG-3').

#### Limited proteolysis

The limited digestion reactions were set up with 24 μg of the full-length ParB protein and 0, 0.004, 0.016 or 0.064 μg trypsin (Sigma) in a 20 μl volume containing 5 mM HEPES-K, pH 7.6, 5 mM MgCl2 and 100 mM KCl. For the DNA protection experiment, ParB was preassembled with 10 μg R4–18 DNA for 30 min before digestion. The digestion reactions proceeded at 20°C for 24 h. Half of each sample was separated on a 4–20% gradient SDS denaturing gel and stained with Coomassie blue. Half of each of the DNA-bound sample was also analyzed on a 5% native polyacrylamide gel and stained with ethidium bromide. For mass spectroscopy identification, the band corresponding to the major cleavage product was excised, in-gel digested by trypsin and analyzed with Q-Star (ABI).

### Crystallization and structure determination

The crystal of the free ParB RHH domain was obtained during crystallization of ParB–DNA complexes. ParB 65–135 was assembled with DNA R4–18 in a 1:1 molar ratio and concentrated to 10 mg/ml in 5 mM HEPES-K (pH 7.6), 5 mM MgCl2 and 100 mM KCl. The DNA complex was incubated with 0.1% (w/w) trypsin on ice for 2 h prior to crystallization to trim disorder regions (22). Crystals were grown by vapor diffusion at 20°C in a hanging drop containing 1 μl of the ParB–DNA complex and 1 μl...
of mother solution containing 0.01 M MgCl₂, 2.4 M (NH₄)₂SO₄, 0.05 M MES, pH 5.6. The crystal was cryoprotected with 30% glycerol in the mother solution and flash-frozen in liquid nitrogen. To make a gold derivative, the crystal was soaked in the mother liquor supplemented with 10 mM KAu(CN)₂ for 12 h. This structure was initially solved using the single isomorphous replacement with anomalous scattering method (SIRAS) based on a 1.7 Å native dataset and a 3.0 Å derivative dataset collected at an in-house generator (Rigaku), and revealed only the free protein. We later obtained a better crystal of free protein during the screen of ParB 65–139 and DNA complex. This crystal grew from 1.8 M Li₂SO₄, 0.01 M MgSO₄, 0.05 M cacodylate-Na and pH 6.0, diffracted to 1.4 Å at Shanghai Synchrotron Radiation Facility (SSRF) beamline BL17U1, and provided a dataset for final refinement.

Diffraction data were reduced using Denzo and Scalepack in-house or HKL2000 at the synchrotron (23). Heavy atoms were detected by SHELXD (24). The phases were calculated, refined and solvent modified in SHARP (25). ARP/wARP was used for automated model building (26). The model was built in COOT (27) and refined in Refmac (28). The current model contains two ParB chains containing residues 69–128 and 70–128, respectively, 10 sulfate ions and 162 water molecules. The Ramachandran plot indicates that 100% of the residues are in a favored region.

Hydroxyl radical footprinting assay

Hydroxyl radical footprinting was performed as previously described (29). For 3’-labeling of the forward strand, a 170-bp centromere DNA sequence was amplified by PCR from a plasmid containing the whole par operon (20,21) with primers ParS3141-F (5’-CTAGAGCAAAGGACCC ACCGACCG-3’) and ParS3310EcoRI-R (5’-GGATTCC TCTGGAATTCCGACCCCGC-3’). The PCR product was cleaved by EcoRI at one end. The resultant sticky end was filled in with a Klenow fragment in the presence of [α-32P]dATP (Furui Biotech, Beijing) and other unlabeled dNTPs, resulting in 3’-end labeling of the forward strand. The sequencing ladders were prepared using the Sanger dideoxy method with the 5’-labeled primer ParS3310-5R (5’-TCTGAGGAAATTTCCAG CCCGCCG-3’). The sequence reads have been converted to denote complementary nucleotides. The reverse strand of the centromere DNA was similarly 3’ labeled using primers ParS3141EcoRI-F (5’-GGATTCCGAAAGGA CCCGCCGACCCCG-3’) and ParS3310-R (5’-TCTGGAAT TTCCAGCCCGCCG-3’). The corresponding sequencing ladders were prepared using the 5’-labeled primer ParS3141-5F (5’-CTAGAGAAAGGACCCCAGACC G-3’).

Labeled DNA probes (20 nM) were incubated in a buffer containing 5 mM sodium cacodylate, pH 7.0, 5 mM MgCl₂, 100 mM KCl, 0.02% NP-40, 0.05 mg/ml poly(dI–dC) with ParB 60–139 or full-length ParB in a final volume of 10 μl at 25°C for 10 min. The concentrations of ParB 60–139 and full-length ParB were both 0.5 and 1 μM in the forward-strand reaction. In the reverse-strand reaction, ParB 60–139 was used at 0.25, 0.5 and 1 μM, and full-length ParB was used at 0.1, 0.2, 0.25, 0.33, 0.5, 0.6, 1 and 2 μM. Three 1 μl drops of 10 mM sodium ascorbate, 0.6% H₂O₂ and 1 mM FeCl₃/2 mM EDTA were added separately on the wall of tube, mixed on the wall by pipetting and spun down to the sample. The reactions were incubated for 4 min and quenched by adding 1 μl of 100 mM thiourea. The reactions were then diluted with 300 mM sodium acetate, pH 5.2, to 200 μl and extracted with phenol–chloroform. The upper phase was collected and 1 μl of GlycoBlue (15 mg/ml; Ambion), and 500 μl of ethanol were added. DNA was precipitated in liquid nitrogen for 2 min and harvested by centrifugation. The pellets were washed with 70% ethanol, briefly dried, resuspended in 8 μl of loading buffer containing 95% (v/v) deionized formamide, 0.5 mM EDTA, 0.025% (w/v) SDS, 0.05% (w/v) xylene cyanol, 0.05% (w/v) bromphenol blue, and separated on an 8% polyacrylamide/8 M urea sequencing gel. The gel was visualized by autoradiography. Software SÅFA (semi-automated footprinting analysis) was used to extract the volumes of resolved bands (30). The volume data were imported into OriginPro 8 (OriginLab) and fit to a sine function:

\[ y = y_0 + A \sin \left( \frac{2\pi(x - x_0)}{w} \right), \]

where \( y \) is the band volume, \( x \) is the nucleotide number, \( w \) is the period, \( y_0 \) is the baseline, \( x_0 \) is the phase and \( A \) is the amplitude of protection.

EMSA

DNA oligos were synthesized and gel-purified by Invitrogen (Supplementary Table S1). Complementary strands (500 μM) were annealed in 5 mM HEPES-K, pH 7.6, at 95°C for 1 min. DNA duplexes were radiolabeled at the 5’ end with \([γ-^{32}P]ATP\) (5000 Ci/mmol; Furui Biotech, Beijing) by T4 polynucleotide kinase (NEB). The 170-bp centromere DNA was PCR amplified as described above followed by dephosphorylation and phosphorylation. Labeled DNA (~0.2 nM) were incubated in binding buffer (5 mM MgCl₂, 100 mM KCl, 0.01% NP-40, 5 mM HEPES-K and pH 7.6) with various concentrations of ParB 65–139 in a final volume of 10 μl at 4°C for 30 min. An equal volume of loading buffer containing 10% glycerol, 0.05% (w/v) xylene cyanol FF and 0.05% (w/v) bromphenol blue was added to the reactions, which were then resolved on a 5% native polyacrylamide gel run in 1 x Tris-glycine buffer at 20 V/cm at 4°C for ~1 h. Gels were dried, exposed to phosphorimagery screens and read by a Typhoon 9400 scanner (GE Healthcare). The volumes of the DNA bands were integrated in Quantity One (BioRad). For stoichiometric assembly, 5 μM DNA was mixed with 0, 2.5, 5, 7.5, 10, 12.5, 15 and 20 μM ParB 65–139 in a 10 μl volume containing the same binding buffer indicated above. The gel was stained by ethidium bromide.
Analysis of two-ParB binding data

We consider the case of ParB binding to a DNA probe with two identical binding sites. The fraction of DNA bound to $i$ ParB molecules $\Theta_i$ ($i = 0, 1, 2$) is given by (31):

$$\Theta_0 = 1/Z, \quad \Theta_1 = 2KL/Z, \quad \Theta_2 = \omega K^2 L^2/Z,$$

where $K$ is the intrinsic microscopic binding constant of ligand to a single binding site, $\omega$ is the cooperativity factor between adjacent interacting ligands, $L$ is the concentration of free ligand, and $Z$ is the binding polynomial:

$$Z = 1 + 2KL + \omega K^2 L^2.$$  

Parameters $K$ and $\omega$ were determined by non-linear global fit of experimentally measured values $\Theta_i$ ($i = 0, 1, 2$) to Equations (2)-(5) as a function of $L$. Because the labeled DNA has a concentration of $\sim 0.2 \text{nM}$ in our experimental conditions and the amount of bound ligand is negligible, the concentration of free ligand $L$ is approximated by the total concentration of ligand. The global non-linear fit analysis of the 2R3-27 and 2R6-27 data was carried out in MATLAB 6.1 with home-written scripts. The intrinsic dissociation constant $K_{d, \text{int}}$ is equal to $1/K$.

ITC

The titrations were performed at 25°C using an ITC-200 microcalorimeter (MicroCal Inc.). DNA solutions (5 or 10 $\mu$M) were prepared by diluting concentrated stocks into the binding buffer containing 5$mM$ HEPES-K, pH 7.6, 5 $mM$ MgCl$_2$ and 100 $mM$ KCl. ParB 65–139 was prepared in the same binding buffer with a concentration of 50 or 100 $\mu$M. Solutions were degassed for 2–5 min before loading. The sample cell was filled with 200 $\mu$l of DNA. ParB 65–139 was injected in a volume of 0.4 $\mu$l for the first injection and 2 $\mu$l for the next 19 injections using a computer-controlled 40-$\mu$l microsyringe with an injection interval of 150 s. Titration of protein into the binding buffer or titration of the binding buffer into the DNA solution produced negligible heat. Integrated heat data were analyzed using a one-set-of-sites model in MicroCal Origin following the manufacturer’s instructions. The first data point was excluded in analysis. The binding parameters $\Delta H$ (reaction enthalpy change in cal mol$^{-1}$), $K$ (binding constant in M$^{-1}$) and $n$ (bound ParB dimer per DNA) were floating in the fit. The binding free energy $\Delta G$ and reaction entropy $\Delta S$ were calculated using the relationships $\Delta G = -RT \ln K$ ($K = 1.9872 \text{cal mol}^{-1} \text{K}^{-1}$, $T = 298 \text{K}$) and $\Delta G = \Delta H - T \Delta S$. The dissociation constant $K_d$ was calculated as $1/K$. The change of the ParB binding free energy of a subsite relative to R3–18 was calculated using the equation $\Delta \Delta G = \Delta G_{\text{subsite}} - \Delta G_{R3-18}$.

RESULTS

The C-terminal half of ParB forms a DNA-binding domain resistant to proteolysis

Sequence analysis of pCXC100 ParB did not reveal any known domains. Searching for ParB homologs by PHI-BLAST returned $\sim 10$ significant hits, but none of them has been biochemically characterized. To experimentally probe the domain organization of ParB, full-length ParB protein was subjected to limited proteolysis by increasing amounts of trypsin and was analyzed by SDS–PAGE (Figure 1A). Full-length ParB protein was digested into a major species with a molecular weight of $\sim 6 \text{kDa}$. Mass spectroscopy analysis showed that the species includes residues 67–129, suggesting that this region folds into a structural core of ParB.

We also digested ParB in the presence of an 18-bp DNA (R4–18) derived from the centromere. The DNA-bound ParB degraded into a stable species of a size similar to that of the free ParB. Native gel analysis showed that the stable species remained bound by DNA (Figure 1B). In the absence of DNA, the $\sim 6 \text{kDa}$ species could be completely digested within 24 h at the highest concentration of trypsin tested whereas the DNA-bound species remained highly resistant to digestion, indicating that DNA interaction stabilizes the structure core. Hence, these results
suggest that the C-terminal half of ParB forms the DNA-binding domain.

Crystallization and structure determination

Given the distinct sequence of ParB, determination of its three-dimensional structure would be essential for resolving its DNA-binding motif. We obtained the crystal of the ParB DNA-binding domain in the free-state during crystallization of ParB–DNA complexes. The structure was solved by the SIRAS method based on a gold derivative. Although the derivative dataset has a high Rmerge value of 0.183, the experimental electron density map is of excellent quality (Supplementary Figure S1). The structure revealed only the DNA-binding domain of ParB but no bound DNA in the crystal. The asymmetric unit of the crystal contains two chains composed of residues 69–128 and 70–128 that fold into an intertwined dimeric structure. The structure has been refined to 1.4 Å resolution with an R factor of 0.198 and a free R-value of 0.212 (Table 1).

Dimeric RHH structure of the ParB DNA-binding domain

The DNA-binding domain of ParB adopts a dimeric RHH structure (Figure 2A). Each subunit consists of a short β-strand, β1 (residues 72–74), followed by two helices, α1 (residues 77–94) and α2 (residues 99–118). Two monomer subunits are tightly intertwined to form a symmetric homodimer. The N-terminal β1 strands pair with each other, forming an antiparallel β-ribbon. The symmetry-related α2 helices cross over by 107° and occupy the center of the structure. The α1 helices are located at the peripheral region with a bend in the last helical turn. The dimerization interface is extensive and buries a total of 1808 Å² of solvent accessible surface area. Residues V73, V75, L83, A86, Y87, L89, F99, F102, I103, L107, V111, L114 and F124 from both subunits constitute the hydrophobic core of the structure, and most of them are conserved among homologs of ParB (Figure 3).

Table 1. Data collection and refinement statistics

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*The values for the data in the highest resolution shell are shown in parentheses.

**Rmerge = \sum |I_i - \langle I_i \rangle| / \sum I_i$, where $I_i$ is the intensity of the measured reflection and $\langle I_i \rangle$ is the mean intensity of all symmetry-related reflections.

$Rwork = \sum |F_o - F_c| / \sum |F_o|$, where $F_o$ and $F_c$ are the observed and calculated structure factor amplitudes of reflection $hkl$.

$Rfree$ is the same as $Rwork$, but calculated on 5% reflections not used in refinement.

Figure 2. Crystal structure of ParB DNA-binding domain. (A) Ribbon representation of the dimeric RHH structure in two orthogonal views. The two subunits are colored blue and green. Secondary-structure elements are labeled. The image on the bottom is viewed along the dyad axis of the dimer. (B) Structural superimposition of ParB (green) with other RHH CBPs: TP288 ParG (blue, 1P94), pSM19035 omega (orange, 2BNW), pSK41 ParR (magenta, 2Q2K) and pB171 ParR (red, 2JD3). These structures were aligned by their two α2 helices in PyMOL. The type and plasmid of the CBP are indicated.
b\textsuperscript{1} are likely responsible for sequence-specific recognition of the centromere. In addition, the conserved residue R84 in helix a\textsubscript{1} may contact with the phosphate–sugar backbone of DNA.

Through structure determination, the RHH motif has been identified in type Ib CBPs TP288 ParG (16) and pSM19035 omega (15,17), and type II CBPs pSK41 ParR (18) and pB171 ParR (19). Structural alignment of monomer subunit to ParB yielded an rmsd of 1.4 Å over 39 C\textalpha pairs for ParG, an rmsd of 1.9 Å over 36 C\textalpha pairs for omega, an rmsd of 2.4 Å over 30 C\textalpha pairs for pSK41 ParR and an rmsd of 3.2 Å over 24 C\textalpha pairs for pB171 ParR. Hence, ParB is structurally more similar to type Ib CBP than to type II CBP. Among the five RHH CBPs including ParB, the two a\textsubscript{2} helices display the most conserved position with an inter-helical angle of 95–113\degree. Alignment of the five structures by their a\textsubscript{2} helices is shown in Figure 2B. The structural conservation of helices a\textsubscript{2} underlies that the N-termini of a\textsubscript{2} helices need to mediate highly conserved contacts with two phosphate groups across the DNA major groove (17,18,33–38). In the alignment, equivalent b\textsubscript{1} strands occupy a similar position, but the conformation of helix z\textsubscript{1} and its connecting loops to helix z\textsubscript{2} and strand b\textsubscript{1} are highly divergent. These regions often mediate variable interactions between adjacent RHH dimers bound to tandem repeats.

In addition, the ParB RHH domain contains a unique C-terminal loop (residues 119–128) following helix a\textsubscript{2} that makes contact with helix a\textsubscript{1} of its dyad mate, contributing to the dimerization interface. Use of extra structural elements for dimerization was also found in the structure of pB171 ParR, which possesses an extensive C-terminal helical domain involved in dimerization (19).

ParB homologs

About 10 sequences share 48% similarity and 29% identity on average with ParB at the RHH motif region (Figure 3). Such a degree of conservation suggests that they all similarly adopt a dimeric RHH fold and likely function in binding specific DNA sequences. These sequences are encoded in seven Gram-positive actinobacteria with most in plasmid DNA and two in the genome. The function of these proteins has not been experimentally studied. Notably, Yag1E\textsubscript{pAG1} in plasmid pAG1 of Corynebacterium glutamicum, Yag1E in the chromosome of Propionibacterium acnes SK137; NP_052577 (Yag1E\textsubscript{pAG1} in plasmid pAG1 of Corynebacterium glutamicum); YP_829452 (Arth_4230 in plasmid 3 of Arthrobacter sp. FB24); YP_829155 (Arth_4403 in plasmid 1 of Arthrobacter sp. FB24); YP_950238 (Aaur_pTC20074 in plasmid TC2 of A. aurescens TC1); YP_002478563 (Achl_4636 in plasmid pACHL02 of A. chlorophenolicus A6); YP_002477960 (Achl_4183 in plasmid pACHL01 of A. chlorophenolicus A6); YP_002784447 (ROP_pKNR-00030 in plasmid pKNR of R. opacus B4). Sequence Achl_4183 is incomplete at its N-terminal region in GenBank and has been corrected here. The numbers in parentheses indicate extra C-terminal residues not shown. Residues with 100 and 80% conservation are shaded in black and gray, respectively.
ParB binding site on the centromere

We previously showed that the binding of ParB protected a ~90-bp centromere region upstream of the par locus from DNase I cleavage (21). To precisely locate ParB binding sites within the centromere and define the boundaries and organization of the centromere, we applied the hydroxyl radical footprinting assay to a 170-bp DNA probe covering the whole centromere region. The hydroxyl radical primarily attacks the C5' atom of deoxyribose, resulting in the breakage of the phosphate-ribose backbone independent of the DNA atom of deoxyribose, resulting in the breakage of the a/C2 complex contains two ParB dimers. ParB apparently differences between the C1 and C2 complex suggest that the dimer to DNA. This observation and the migration differences between the fourth and fifth nucleotides of a segment and each segment roughly corresponds to a binding site of ParB RHH domain. Sequence alignment of segments 1–9 reveals modest conservation with a consensus sequence ‘AGNTGGAAA’ (Figure 4F), indicating that the centromere is composed of nine unspaced direct 9-bp repeats. Repeat 10 was not included in the alignment for it does not bind ParB by itself, as shown below. The centromere is less likely to contain an inverted repeat, as none of inverted repeats matches the consensus sequence better than their direct counterparts (Supplementary Figure S2).

A ParB dimer contacts an 18-bp region around its binding center

Association of a ParB dimer to a single subsite is the first step in the assembly of a full partition complex. We ask how ParB recognizes a single subsite. To find a suitable length of subsite required for ParB binding, we applied ITC to compare binding of the ParB RHH domain to four 14–20-bp probes centered around repeat R4 (Figure 5A). A 14-bp probe that fully covers repeat R4 failed to bind ParB. Elongation of the probe from 16- to 18-bp reduced $K_d$ value from 530 to 60 nM, while further elongation to 20-bp caused no change in $K_d$ value (68 nM). Repeat R3 also requires an 18-bp region around its binding center to bind ParB (Supplementary Table S2). We conclude that a length of 18-bp around the binding center is required for ParB to efficiently bind a single subsite.

Binding stoichiometry of ParB with one- and two-site probes

These 18-bp one-site probes formed only a single species of complex in EMSA (Figure 6), which we interpreted as one ParB dimer-bound complex. However, the complex might actually contain a ParB dimer-of-dimers, which is the minimal functional unit of most RHH domains (32). To directly assess the binding stoichiometry, we preformed EMSA with different molar ratios of ParB 65–139 and unlabeled DNA probes (Figure 5B). We compared the assembly of an 18-bp DNA containing repeat R3 and its flanking sequences (R3–18), an 18-bp DNA containing repeats R2 and R3 (R2R3–18) and a 27-bp DNA containing repeats R2 and R3, and their flanking sequences (R2R3–27). One-site probe R3–18 was assembled almost completely into a C1 complex with equal molar amount of ParB dimer, indicating that the C1 complex contains one ParB dimer. In contrast, two-site probes R2R3–18 and R2R3–27 formed predominately a C2 complex with a ~1.5-fold molar ratio of ParB dimer to DNA. This observation and the migration differences between the C1 and C2 complex suggest that the C2 complex contains two ParB dimers. ParB apparently
Figure 4. The ParB-centromere interaction probed by hydroxyl radical footprinting. (A) Footprinting of the forward strand of ParB-bound centromere. A 170-bp centromere DNA was labeled at the 3'-end of the forward strand, assembled with 0.5 and 1 μM of ParB RHH domain (residues 60–139) (lanes 2 and 3), and 0.5 and 1 μM of full-length ParB (lanes 4 and 5) and subjected to hydroxyl radical cleavage. Lane 1 contains no protein. Sequencing ladders (A, T, G, C) are indicated. The most protected nucleotides derived from curve fitting are shown on the right. (B) Footprinting on the reverse strand. The ParB 60–139 concentrations were 0.25, 0.5 and 1 μM in lanes 2–4, and the ParB 1–139 concentrations were 0.1, 0.2, 0.25, 0.33, 0.5, 0.66, 1 and 2 μM in lanes 5–12. (C) The band volumes of the cleavage products are plotted against nucleotide numbers. Lane 3 (boxed) of the forward strand and lane 4 of the reverse strand were analyzed. Smaller volume corresponds to greater protection at the respective site. The lines represent the best fit of a sine function to the protection data with a period of 9.07 ± 0.08 for the forward strand (top panel) and 8.96 ± 0.09 for the reverse strand (bottom panel). (D) Structural model of the ParB RHH domain in complex with DNA. A standard B-form DNA was positioned based on the aligned pSK41 ParR-DNA complex structure (2Q2K). Red circles refer to the DNA backbone regions contacted by the N-terminus of helix α2. (E) The structure of the centromere. The forward strand is numbered from 5' to 3', and nucleotides in the reverse strand are numbered as their pairing nucleotides in the forward strand and are denoted by prime. Nucleotides with maximal protection in hydroxyl radical footprinting are marked by red circles and their nucleotide numbers. Ellipses denote binding centers of ParB dimer. Hollow ellipses are extrapolated from incomplete protection data. Alternative repeats are shaded in green and gray. Repeats R0, R10 and R11 do not bind ParB by themselves and are denoted in italics. (F) Alignment of repeats R1–R9 and the consensus sequence. The sequence logo is indicated at the top. A loose criterion of 50% conservation was used to define the consensus sequence.
associates with the two-site probes in a sequential manner as the one-ParB bound C1 complex was transitively populated.

The above EMSA experiments were performed with protein and DNA concentrations in the micromolar range. Under these conditions, excess protein was found to lead to larger complexes that are likely caused by non-specific binding. These complexes appeared as smeared bands for R3–18 and R2R3–27. Notably, a discrete C3 complex is evident for R2R3–18, and less so for longer R2R3–27. The C3 complex likely contains 3 ParB dimers as judged from its migration rate, but its assembly mode is unclear. The non-specific complexes were not observed in EMSA using low concentrations (10 nM) of 32P-labeled DNA.

Affinity variation among individual subsites

The 10 centromere repeats are relatively divergent in sequence, raising the question of whether they have a similar affinity toward ParB. To this end, we characterized the binding of ParB RHH domain (residues 65–139) to individual subsites. We used EMSA and ITC to evaluate the binding affinity of 12 18-bp DNA probes that center around each of 10 repeats (R1–R10) as well as the putative repeat R0, which is upstream of R1, and the putative repeat R11, which is downstream of R10 (Figure 6). The ITC data were analyzed with a one-set-of-sites model that assumes one ParB dimer binding to one DNA. Some fit curves show systemic deviation from the data, which is probably caused by additional binding events, as shown in Figure 5B. The $K_d$ values derived from ITC are indicated in Figure 6, and the full sets of binding parameters are listed in Supplementary Table S2.

Subsites around R0, R10 and R11 display virtually no binding in both EMSA and ITC experiments, indicating that the centromere core contains only repeats R1–R9. Although binding of repeat R10 was detected in the footprinting experiment, this result is likely due to the high affinity of subsite R9 and binding cooperativity between adjacent ParB molecules (see below). ITC measurements showed that subsites R9, R3, R8 and R4 have the highest binding affinities with $K_d$ values of 11–87 nM, whereas the rest subsites show $K_d$ values in the submicromolar range (380–950 nM). The two orders of magnitude variation in binding affinity among individual subsites is probably caused by their divergent sequences.

Sequence recognition of ParB

To assess the contribution of individual nucleotide to ParB recognition, we replaced each of 18 base pairs in the one-site probe R3–18 to other three alternatives and measured the ParB binding affinity by ITC (Table 2, Supplementary Table S2, and Figure 7A, Supplementary Figure S3). Sequences in positions 4, 5, 7, 8, 11, 14 and 16 are not recognized by ParB, as corresponding point mutants retain the similar binding affinity. The most significant mutational effect occurs at positions 9 and 10, which are immediately adjacent to the ParB binding center. Mutation of T9 to adenine and guanine reduced the binding affinity by 5- and 12-fold, respectively, while replacement of G10 by cytosine yielded an ITC profile lacking two-state transition (Supplementary Figure S3). Moreover, guanine substitution of A6 or A12 decreased the binding affinity by 7-fold. Since steps 6–12 are within 4-bp of the ParB binding center, these sequences may be recognized by the β-ribbon structure of ParB dimer, according to the known RHH–DNA interaction mode.
Interestingly, ParB also recognizes distal sequences from the preceding and succeeding repeat. Specifically, steps 1–3 favor strongly the ‘GGA’ sequence and mutations at these positions reduced the binding affinity by up to 4–7-fold. At the downstream region, steps 13, 15, 17 and 18 show marginal selection of sequence with corresponding mutants causing <3-fold increase of $K_d$ value.

To understand the affinity variation among different subsites R0–18 to R11–18 (Figure 6), we attempted to estimate the change of the ParB binding free energy ($\Delta \Delta G$) of these subsites relative to R3–18 on the basis of their sequence. To the first-order of approximation, $\Delta \Delta G$ of a subsite sequence can be calculated as a sum of $\Delta \Delta G$ due to single substitution. The predicted $\Delta \Delta G$ values generally agree with the experimentally measured values (Figure 7B). However, subsites R9–18 and R10–18 show some discrepancy between the predicted and measured $\Delta \Delta G$, likely due to coupling interaction among multiple substitutions.

Verification of the structural organization of the centromere

We attempted to verify by a second approach the positions of the ParB binding centers on the centromere that are predicted from the footprinting experiment.

Figure 6. Binding affinity of individual sites. Twelve 18-bp probes covering repeats R0-R11 were assessed for binding of ParB RHH domain (residues 65–139) by EMSA (top panel) and ITC (bottom panel). The concentrations of ParB dimer in EMSA were 0, 2, 8, 16, 32, 64, 128 and 512 nM. The curves in the ITC panels are the best fit to a one-set-of-sites model that dose not account for possible additional binding events. Fit of the R7-18 data was unsuccessful.
We examined the binding of the ParB RHH domain with an array of eight 18-bp probes that are consecutively shifted by one base pair to cover repeat R4 (Figure 8A). The R4–18 probe that has its midpoint at the predicted binding center of repeat R4 should display the highest affinity. If the midpoint of a probe is off the binding center, we would expect to see reduction of the binding, as one side of the binding center may become too short for optimal contact. The EMSA results showed that probes R4–18−1 and R4–18+1, with a midpoint 1-bp off the binding center of R4, display a binding affinity similar to that of R4–18, whereas probe R4–18+2, which has a midpoint 2-bp downstream of the binding center of R4, shows significantly reduced binding. These observations are consistent with the predicted ParB binding center on repeat R4.

Interestingly, when the probe midpoint was moved farther upstream (R4–18−C0/2, R4–18−C0/3) or downstream (R4–18+3, R4–18+4), we observed a second slow-migrating DNA–protein complex C2, which apparently corresponds to two-ParB bound complex. In these cases, the probes contain part of the R3 or R5 repeat. Despite being incomplete, the adjacent R3 and R5 sites show significant binding to a second ParB molecule. In contrast, efficient binding to a single-site probe requires a length of 9-bp at either side of a binding center. These results suggest that the ParB molecule bound at repeat R4 strongly attracts a second ParB molecule to adjacent weak sites. Comparison of probe R4–18−C0/2 versus R4–18+2 and R4–18−C0/3 versus R4–18+3 shows that even in their incomplete repeat forms, repeat R3 is better than R5 in cooperating R4 binding of ParB, which is consistent with the former having an 24/25-fold higher intrinsic binding ability (Figure 6). These results are fully consistent with the proposed structural model of the centromere and also indicate strong cooperativity for ParB in binding tandem repeats.

### Binding cooperativity

To quantify the binding cooperativity of ParB, we analyzed ParB binding to probes containing two subsites. Different affinities associated with the natural centromere binding sites would complicate measurement of cooperativity. To simplify the analysis, we engineered probes that contained two identical subsites over an 18-bp region surrounding each binding center. To construct such probes, we duplicated the sequence of a 9-bp repeat four times and selected a 27-bp segment between the two outermost binding centers. Two such probes—2R3−27 and 2R6−27—were derived from repeats R3 and R6, which represent a strong and a weak binding site, respectively (Figure 9A and B). The corresponding one-site 18-bp probes for R3 and R6 (R3−18 and R6−18) were also created.

As expected, the one-site probes form only a single complex with ParB, whereas the two-site probes form two types of complexes corresponding to one- and two-ParB bound complexes (Figure 9C–F). For two-site probes, the two-ParB complex is more favorably formed over the one-ParB complex even at low protein concentrations. The cooperativity effect was the most dramatic for the weak repeat, R6. Global analysis of the binding isotherms yielded a cooperativity of 16 for tandem R3 subsites and 112 for tandem R6 subsites (Figure 9G and H).

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**Table 2.** Dissociation constants $K_d$ (nM) of R3–18 point mutants and ParB 65–139 measured by ITC

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<th>No.</th>
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**Figure 7.** Sequence recognition of ParB. (A) Binding constants of R3–18 and its point mutants plotted against the position of the replaced nucleotide. For the purpose of illustration, mutant G10C that shows weak ParB binding is set to have a $K_d$ value of 300 nM. (B) The predicted and measured change in the binding free energy relative to R3–18 for 18-bp subsites. Repeats are shown as arrows and ParB binding centers as vertical lines. Each nucleotide is also labeled by its position in a repeat and distance to the central ParB binding center. N.B.: No binding; N.F. Not fit.
Figure 8. Binding of ParB to frame-shifted R4 probes. (A) Eight frame-shifted R4 probes. The R4 repeat is shown in black and its flanking sequences from repeats R3 and R5 are shown in gray. Vertical bars indicate the positions of the binding centers of R3, R4 and R5. (B) EMSA of the ParB RHH domain (65–139) and R4 probes. Lane 1 in each panel has no ParB, and the subsequent lanes have increasing amounts of protein in concentrations of 2, 8, 16, 32, 64, 128, 512 and 5120 nM. C1 and C2 denote complexes formed by one and two ParB dimers, respectively.

Figure 9. The cooperativity of ParB binding to tandem identical repeats. (A and B) Sequence of two- and one-site probes designed for repeats R3 (A) and R6 (B). (C–F) EMSA of ParB 65–139 with one-site probe R3–18 (C) and R6–18 (D), and two-site probes 2R3–27 (E) and 2R6–27 (F). C1 and C2 denote complexes formed by one and two ParB dimers, respectively. (G and H) Global analysis of the binding isotherms for 2R3–27 (G) and 2R6–27 (H). The lines are the best fit to Equations (2–5), with intrinsic dissociation constant $K_{d_{int}} = 112$ nM and cooperativity $\alpha = 16$ for 2R3–27; $K_{d_{int}} = 877$ nM and $\alpha = 110$ for 2R6–27. (I) EMSA of ParB 65–139 with a $^{32}$P-labeled 170-bp DNA covering the whole centromere region.
The association of ParB to a 170-bp DNA covering the whole centromere region is governed by an even stronger cooperativity (Figure 9I). Majority of the DNA was transformed into a major partition complex in the presence of 8 nM ParB 65–139, while at the same concentration the two-site probes remain not fully assembled (Figure 9E–F).

DISCUSSION

The active partition system for low-copy-number plasmids is a simple and autonomous mechanism for separating duplicated DNA into daughter cells. Studies on a few model plasmids have begun to reveal the molecular mechanisms of many aspects of plasmid partitioning (5). The first step of plasmid partition is assembly of CBP and centromere repeats into the partition complex, which then recruits filament-forming ParA protein. In this study, we structurally and biochemically characterized the CBP ParB from a type Ib partition system recently reported in plasmid pCXC100 (20,21). The obtained results indicate that the pCXC100 partition system is generally similar to other type Ib systems in terms of ParB structure, centromere organization and cooperative ParB-centromere interaction, and also reveal a distinct overlapped assembly mode of ParB to the centromere repeat.

ParB structure

We show that the DNA-binding domain of ParB is located at its C-terminal half and folds into a dimeric RHH structure. The RHH motif of ParB could not be predicted based on sequence homology, as is the case with many other RHH domains. Our structure adds to the list of four known RHH domains of type Ib (omega and ParG) and type II (pSK41 ParR and pB171 ParR) CBPs (15,16,18,19), and further supports the prevalence of the RHH domain in centromere binding. In addition to its C-terminal RHH domain, ParB contains an N-terminal domain (residues 1–65). This region is likely flexible as it is highly susceptible to trypsin digestion in both the absence and the presence of DNA (Figure 1) and is predicted to lack secondary structure (40). The presence of a flexible tail upstream the RHH domain appears to be a common feature of type Ib RHH CBPs. A shorter N-terminal flexible tail is also present in ParG (~32 residues) and omega (~20 residues) (15,16). The sequences of the N-terminal domains are not similar among ParB, omega and ParG, nor are they similar among ParB homologues (Figure 3). Nevertheless, the N-terminal domain may be functionally important, as the corresponding region of ParG has been shown to be involved in the assembly of the ParG-DNA higher structure for transcription repression and in the activation of polymerization and ATP-hydrolysis activity of the ParG partner ParF (41,42). The N-terminal region of omega was also critical for stimulation of ATPase activity and polymerization of its partner delta (43).

Structure of the centromere

We previously relied on sequence analysis to find a weak 9-bp periodicity within the centromere region and divided the centromere into nine 9-bp repeats (21). The phase of the repeat was arbitrarily set due to lack of strong conservation in repeat sequences. Now, we have experimentally mapped ParB binding sites on the centromere together with high-resolution hydroxyl radical footprinting on both strands of the centromere (Figure 4). Based on the known RHH-DNA interaction mode, we were able to deduce the structure of the centromere and the precise location of ParB binding centers. We found that the centromere is composed of nine unspaced 9-bp direct repeats. The repeat is non-palindromic and the binding center of ParB dimer is likely located between the fourth and fifth base pairs of each repeat. The current repeats have the same 9-bp cycle as the previously defined ones, but they differ in phase and boundary. The previously defined repeats are shifted 4-bp downstream compared to the current ones and cover a region corresponding to current repeats R2–R10.

Cooperativity in ParB–centromere interaction

The cooperativity of ParB in binding multiple repeats is similar to that of other RHH domains. Indeed, cooperativity is a crucial feature characterizing RHH-DNA interaction (32). The majority of RHH domains have extremely weak or no binding toward a single subsite and bind efficiently only to tandem subsites as a dimer-of-dimers. In this regard, it is notable that ParB is capable of binding 18-bp single subsites with affinities up to 11 nM. Arc repressor of bacteriophage P22 is one RHH protein sharing this property. Although Arc binds as a dimer-of-dimers to a 21-bp operator that contains two subsites, it can also bind as a dimer to DNA fragments containing single subsite with nanomolar affinities (44). Arc dimer has a strong cooperativity (\( \alpha = 5900 \)) towards full operator that literally prevents the detection of one-dimer bound intermediate complex (44). In contrast, ParB exhibits a moderate cooperativity (\( \alpha = 16–110 \)) and one-dimer bound complex remains observable for two-site probes (Figures 5B, 8 and 9).

The binding cooperativity would allow the association of ParB with intrinsically weak sites within the centromere and spread out of the centromere core. The latter phenomenon is reflected by the detection of binding of repeat R10 in the context of the whole centromere in the footprinting experiment (Figure 4) and the observation of binding of repeat R0 in the presence of repeat R1 by EMSA (data not shown). This spreading effect was previously observed for the interaction of omega with its cognate DNA (45).

Sequence specificity of ParB interaction

A ParB dimer requires a region of 18-bp covering one central repeat as well as two surrounding half repeats for binding single subsites (Figure 5A). This is unprecedented given that an RHH domain normally contacts only ~8–10 bp of DNA. Such an interaction mode implicates that ParB dimers bound adjacent to the centromere...
would contact an overlapped region. Consequently, the sequence in a repeat could be jointly specified by adjacent ParB dimers. The mutagenesis data support that ParB dimer recognizes the sequences of flanking repeats, particularly the fifth to seventh nucleotide of the preceding repeat (Table 2 and Figure 7). For instance, the sixth nucleotide of repeat, conserved as guanine, appears to be specified by the succeeding ParB rather than the immediately bound molecule. Determination of the ParB–DNA complex structure would elucidate the molecular details about their unusual interaction mode.

**PDB ACCESSION CODE**

The atomic coordinates and structure factors have been deposited in the Protein Data Bank under accession code 3NO7.

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

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