

Superstructure of the centromeric complex of TubZRC plasmid partitioning systems

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Edited by J. Richard McIntosh, University of Colorado, Boulder, CO, and approved August 30, 2012 (received for review June 26, 2012)

Bacterial plasmid partitioning systems segregate plasmids into each daughter cell. In the well-understood ParMRC plasmid partitioning system, adapter protein ParR binds to centromere *parC*, forming a helix around which the DNA is externally wrapped. This complex stabilizes the growth of a filament of actin-like ParM protein, which pushes the plasmids to the poles. The TubZRC plasmid partitioning system consists of two proteins, tubulin-like TubZ and TubR, and a DNA centromere, *tubC*, which perform analogous roles to those in ParMRC, despite being unrelated in sequence and structure. We have dissected in detail the binding sites that comprise *Bacillus thuringiensis tubC*, visualized the TubRC complex by electron microscopy, and determined a crystal structure of TubR bound to the *tubC* repeat. We show that the TubRC complex takes the form of a flexible DNA–protein filament, formed by lateral coating along the plasmid from *tubC*, the full length of which is required for the successful in vitro stabilization of TubZ filaments. We also show that TubR from *Bacillus megaterium* forms a helical superstructure resembling that of ParR. We suggest that the TubRC DNA–protein filament may bind to, and stabilize, the TubZ filament by forming such a ring-like structure around it. The helical superstructure of this TubRC may indicate convergent evolution between the actin-containing ParMRC and tubulin-containing TubZRC systems.

FtsZ | X-ray crystallography | DNA segregation

Low copy number plasmids often encode their own segregation machinery, ensuring that copies are partitioned into each daughter cell. These plasmid-partitioning systems organize replicated plasmids and actively separate them. The known plasmid-partitioning systems are minimalist and elegant. They consist of just three components: a DNA centromere, an adapter protein that binds the centromere, forming the centromeric complex, and a nucleotide triphosphate-dependent filament-forming protein, which produces the force to move the plasmids (1, 2).

Plasmid partitioning systems have been divided into types I–III, based upon the homology of their filament-forming proteins to known protein families (2, 3). Type I plasmid partitioning systems (4, 5) are based on deviant Walker A ATPases (6, 7), type II (8) on actin-like proteins (9), and type III (10–12) on tubulin/FtsZ-like proteins (13, 14). Although the structure of the filament implicated in segregation has been determined for all three systems (9, 14–16), only examples of type I and II centromeric complexes have so far been resolved (17–19).

The centromeric complex of the type II (actin-like) ParMRC plasmid partitioning system is formed by the binding of adapter protein ParR to *parC*, a series of parallel (direct), 11-base pair (bp) repeats (20). The superstructure this complex forms is helical, right-handed, and places the *parC* DNA on the exterior of the helix and ParR on the interior (18, 19). This arrangement clusters the interaction surfaces between ParM and ParR, which are located at the tip of the ParM filament and within the last few residues of ParR (18, 19). This clustering probably enables processive filament tip tracking by promoting binding between the filament end and the RC helix, ensuring that only *parC*-bound ParR is able to seed and grow, or at least stabilize filaments.

The centromeric complex of the type III (tubulin-like) TubZRC plasmid partitioning system is also composed of an adapter, TubR, and a DNA centromere, *tubC*. The rough disposition of four of the repeats making up *Bacillus thuringiensis* serovar *israelensis* pBtoxis

(*Bt tubC*, but not the TubR binding site or register, has been assigned by Tang and colleagues who demonstrated binding by *Bt TubR* to this region of pBtoxis (10, 21). Furthermore, the structure of the TubR adapter protein has been determined in the absence of DNA (13), revealing that it forms a recognition-helix dimer with a potential DNA binding surface and demonstrating that it is unrelated to type II ParR (18, 19). Finally, it has been shown that the TubRC complex is capable of recruiting TubZ (13, 21). A summary of previous work on the TubZRC system is provided in Table S1.

In this study we examine in detail the superstructures formed by TubRC complexes. We show that *Bt tubC* in fact contains seven high-affinity binding sites, the full region being required for proper in vitro activity. Furthermore we reveal that TubR coats *tubC*: the centromeric complex of TubZRC plasmid partitioning systems therefore takes the form of a DNA–protein filament. We present a 7.0-Å crystal structure revealing the superstructure of the flexible *Bt TubRC* filament and provide electron microscopy and a 3.5-Å crystal structure that imply a preferred conformation for the TubRC filament of *Bacillus megaterium* (*Bm*), allowing us to compare the superstructures of type III (tubulin-like) centromeric complexes to those of type II (actin-like) plasmid partitioning systems.

Results

***Bt tubC* Dissected by Microarray.** We initially set out to precisely define *B. thuringiensis* (*Bt tubC*). The structure of the centromere is defined by the binding of its protein partner; therefore, we dissected the binding of *Bt TubR* to each independent site within this part of pBtoxis. We designed a microarray in which each experimental measurement consisted of a 24-bp double-stranded DNA sequence (Fig. 1*B*), formed by a hairpin stabilized through a tetraloop (22). This 24-base “window” was scanned through the plasmid in successive measurements; a 4-bp shift per measurement was used to conduct a coarse scan of a large area of pBtoxis, and single base pair shifts were then used for a high-resolution scan of the *Bt tubC* region (Fig. 1*B* and *C*). *Bt TubR* bound the array in only a few places, and binding was sustained reproducibly with the movement of the experimental window. The *Bt tubC* region was readily identified as the sole site of strong, sustained binding.

***Bt TubR* Binds to *tubC* Cooperatively.** Peaks of binding by *Bt TubR* were clearly identifiable (Fig. 1*C*). Fourier analysis of this signal with respect to sequence confirmed the repeat length to be between 12 and 13 bp (Fig. S1) (10). We aligned the sequences producing peak signal allowing for a 12-bp repeat and produced a weighted consensus sequence of the repeats found in *Bt tubC* (Figs. S1 and S2). Two directly repeated sequences were required to produce each peak, troughs in binding occurring

Author contributions: C.H.S.A. and J.L. designed research; C.H.S.A. performed research; C.H.S.A. and J.L. analyzed data; and C.H.S.A. and J.L. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org (PDB ID codes 4ASN, 4ASO, and 4ASS).

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This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1210899109/-DCSupplemental.

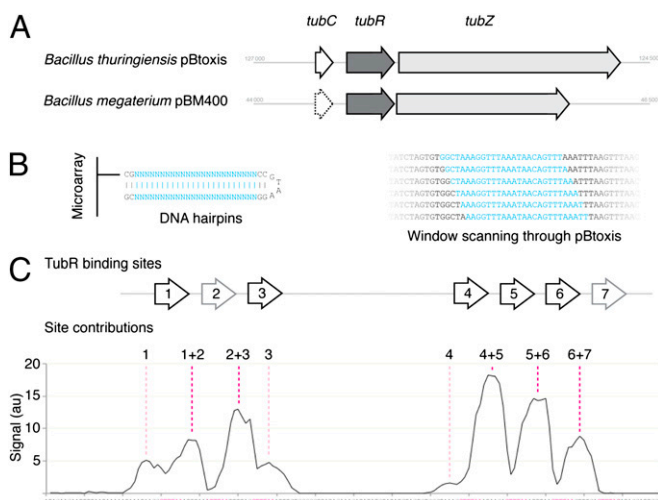


Fig. 1. *Bt tubC* is composed of seven repeats, which bind TubR in a cooperative fashion. (A) Schematic comparing the *tubZRC* loci of *Bt* pBtoxis and *Bm* pBM400. (B) Illustration of the DNA hairpins produced on a microarray to sample the sequence of pBtoxis, and a schematic indicating how this window was scanned through a region of the plasmid sequence by successive single base pair movements. The variable window is shown in cyan, and all other base pairs in black. (C) Plot of the recorded signal for each microarray spot in a 1-bp scan over the region of *Bt tubC* (bp 126688 to 126496). Each point is plotted over the 12th bp of the 24-bp hairpin, with the sequence shown below. The assigned binding sites for *Bt TubR* are shown above, and the corresponding sequences are colored magenta below. The site(s) resulting in each peak have been annotated above the graph.

on centering of a single binding site (Fig. 1C and Fig. S2), implying that *Bt TubR* binding is highly cooperative, dependent on lateral stabilization. This mechanism is consistent with the known activity of TubR as a repressor (12), providing a mechanism by which affinity may be tuned. Stronger sites (such as sites 1, 3, and 4) were capable of supporting binding on adjacent DNA, indicated by low signal peaks in the absence of adjacent cognate sites, implying that spreading of *Bt TubR* along the plasmid is possible (Fig. 1C). We confirmed that such lateral spreading occurs *in vitro* by electrophoretic mobility shift assay (Fig. S3).

***Bt tubC* Is Seven TubR Binding Sites in Two Clusters.** *Bt TubR* binding was limited almost exclusively to the region of DNA 3' from *tubR*. Whereas four direct repeats have previously been identified (21) we found that there are in fact two clusters of *Bt TubR* binding sites, the first comprising three adjacent sites (sites 1–3) and the second four (sites 4–7), which together we identify as the native *Bt tubC* (Fig. 1C). Each block binds *Bt TubR* independently *in vitro* (Fig. S3). Because the two blocks are separated by 54 bp, spreading of *Bt TubR* cannot coat all intervening DNA; however, it is well established in the case of ParR from *Escherichia coli* plasmid R1 that separate blocks of sites are linked by looping of the intervening DNA, so a single complex may be formed (18, 23). Evolution has selected for the production of an array of *Bt tubC*-bound TubR dimers; these must presumably therefore be required to perform the function of the TubRC centromeric complex.

***Bt TubRC* Forms a DNA–Protein Filament *In Vitro*.** To understand why *Bt tubC* had evolved an array of parallel repeats, we examined the complex formed by *Bt TubR* on its centromere using electron microscopy. We produced full-length, untagged *Bt TubR* and purified it to homogeneity (Fig. 2A); it was then bound to a PCR product encompassing the full *Bt tubC* region. DNA–protein filaments composed of blocky subunits were clearly visible in samples of the *Bt TubRC* complex (Fig. 2B). *Bt TubRC* filaments had an average width measuring 5–6 nm, consistent with the

greatest width of a single TubR dimer, whereas the separation of the subunits (from Fourier transform) was 4–5 nm, suggesting that a single fiber of *Bt tubC* DNA was bound by a single row of TubR dimers. The filaments were typically extended, although some exhibited a higher degree of curvature, indicating that these complexes are flexible.

Structure of *Bt TubRC* at 7 Å. Unfortunately the *Bt TubRC* DNA–protein filaments were not sufficiently ordered for either helical or single-particle reconstruction; therefore, we obtained a higher resolution model by cocrystallizing *Bt TubR* with oligonucleotides from *tubC*. Cocrystals were obtained with *Bt TubR* and a 24-bp double-stranded oligonucleotide (*tubC*24). Diffraction was recorded to 7 Å. The structure shown in Fig. 2 C–E was solved by molecular replacement using PDB ID code 3M9A and B-form DNA (13) and verified by phasing selenium anomalous differences to confirm that the difference density obtained identified the selenomethionine sites (Fig. S4 and Table S2). *Bt TubRC*24 crystals were of space group C2, each asymmetric unit containing eight *Bt TubR* dimers and four *tubC*24 oligonucleotides. Electron density was surprisingly good (Fig. S4) and the positions after rigid body refinement resulted in close contacts but few clashes, implying a reasonable model; however, determination of the register of the DNA sequence lay beyond the resolution of our data.

***Bt TubRC* Crystals Contain a DNA–Protein Filament.** *Bt TubRC*24 forms an extended DNA–protein filament in our crystals. The DNA component, *tubC*24, describes a continuous linear fiber, curving in a slight right-handed superhelix, which forms the core of the complex. The outside of the filament is decorated with TubR molecules, forming an 8/1 helix with a rise of 38.4 Å, corresponding exactly to the 12-bp cognate repeat found in *Bt tubC* (Fig. 2 C and D). The coating of the outside of the DNA helix by *Bt TubR* enables the majority of crystal contacts to be formed between protein side chains rather than by DNA, as is expected because of the highly charged nature of the phosphate backbone. The *Bt TubRC*24 filament has very similar dimensions to the extended filaments observed by electron microscopy, and we conclude that this structure is representative of the predominant state of *Bt TubRC* in solution.

DNA Binding by *Bt TubR*. *Bt TubR* bound predominantly within the major groove of *tubC*. The N termini of the paired “recognition” helices together protrude deep into the major groove where they are able to make contact with 5 bp. This is the only major groove contact made, and it is therefore likely to correspond to the sequence-specific component of *Bt tubC*. The consensus binding sequence GTTTAA is not a simple twofold however, suggesting that DNA conformation must also play a role. On either side of the major groove binding site, the phosphate backbone descends into a cleft between the short helix and loop N-terminal to the recognition helix and the β -hairpin of *Bt TubR*. The negative charge of the phosphates will be complemented by lysines 43, 54, and 79, and arginine 74, which form the sides and base of this cleft. Finally, the tip of the β -hairpin protrudes toward the minor groove where arginine 77, which clashes with the phosphate backbone after rigid body refinement, must intrude into the groove itself (Fig. 2E).

Currently, only one other structure of a recognition helix dimerized form of the winged helix–turn–helix fold bound to DNA is available, that of FadR from *E. coli* (24). The DNA binding mode is conserved between the two structures, although the minor groove binding β -hairpin is positioned slightly differently in FadR. The conformation of the DNA in the two structures deviates substantially, however. Both proteins extend the DNA helix relative to its conformation in standard B-DNA; however in the case of FadR, the DNA is bound closely within the minor grooves, bending the course of the fiber toward the surface of the protein, whereas *Bt TubR* binds less closely in the minor groove, the DNA bending slightly away from the protein. The minor groove is also elongated, possibly due to the interaction with the β -hairpin. The full length of DNA contacted by a single *Bt TubR* dimer is 16 bp,

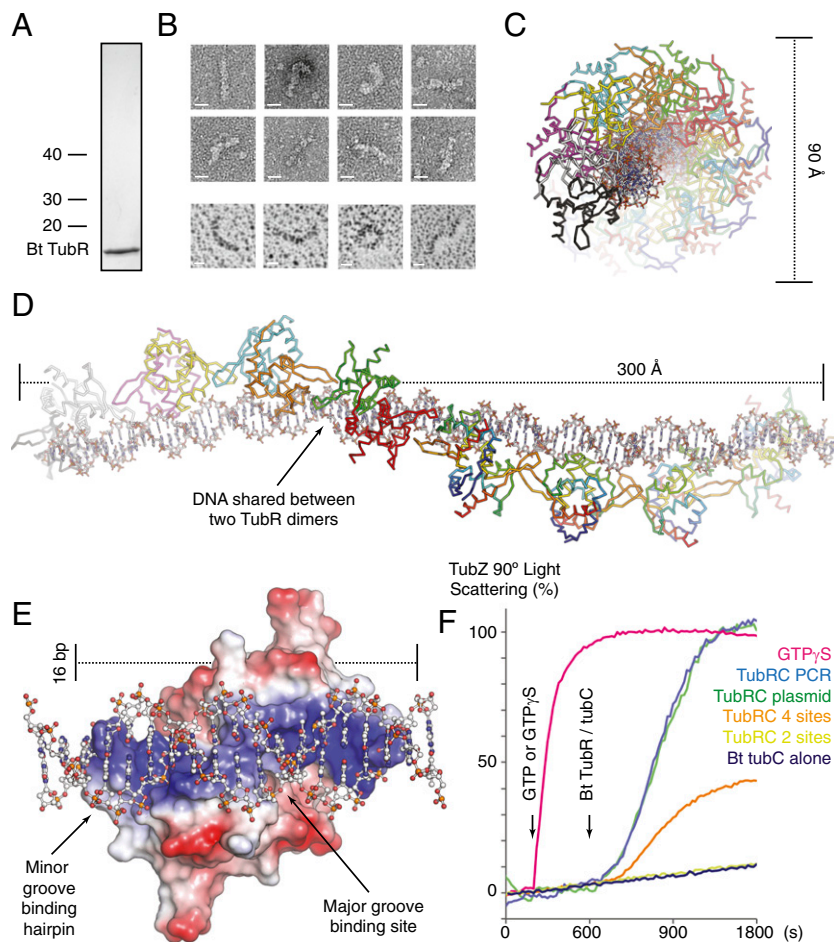


Fig. 2. *Bt* TubR bound to *tubC* forms a DNA-protein filament that effects TubZ polymerization. (A) Coomassie-stained SDS/PAGE of purified *Bt* TubR. (B) Electron micrographs of negatively stained (Left) and rotary shadowed (Right) *Bt* TubR bound to full-length *tubC* showing the morphology of the DNA-protein filaments formed. (Scale bars, 10 nm.) (C and D) Crystal structure of *Bt* TubR (α ribbon representation, the first four dimers are colored by chain, the second four a continuum between blue at the N terminus and red at the C terminus) bound to two repeats of *tubC* (stick representation, C in white/CPK colors). Dimensions in angstroms are shown alongside the structure, whereas the filament has been rotated by 90° between the two plates. (E) Single *Bt* TubR dimer (surface charge representation, red negative, blue positive) from the structure, with B-DNA [ball and stick representation, C in white/Corey, Pauling, Koltun (CPK) coloring] extended from the 12-bp section used in refinement in order to show the interaction with the full 16-bp region covered by the dimer. (F) Effect of *Bt* TubRC on the polymerization of TubZ measured using 90° light scattering. GTP or GTP γ S was added to all reactions at 200 s, and the complexes indicated to the Right of the graph were added at 600 s (color coded). The four- and two-site versions of *Bt tubC* encompassed binding sites 4–7 and 6–7.

although the *tubC* repeat length is 12 bp (13). This requires 4 bp to be shared between adjacent dimers, which occurs at the site of protein-protein contact between the minor groove binding hairpins (Fig. 2 C–E). These β -hairpins from adjacent *Bt* TubR dimers are tightly paired in the minor groove, their opposition favoring the DNA curvature in the *Bt* TubRC24 structure.

TubRC Stabilizes TubZ Polymerization in Vitro. To confirm that the TubRC complex we had discovered was relevant to the activity of the TubZRC plasmid partitioning system, we assayed its effect on TubZ polymerization by 90° light scattering. A recent study by Oliva and colleagues (25) has shown that TubRC increases polymerization of TubZ. We therefore mixed *Bt* TubZ with GTP below the critical point of filament formation measurable by light scattering. *Bt* TubRC complex was then added, which stabilized filament growth. A similar effect, with the same maximum polymerization, can be produced by replacing GTP with GTP γ S, implying that the bar to growth is due to hydrolysis of the nucleotide and disassembly of the filament at a higher rate than growth (Fig. 2F). Given that the *Bt* TubRC complex was present at a substoichiometric level, this suggests that it might prevent depolymerization of TubZ filaments, possibly switching from treadmilling to elongation.

To determine whether or not the whole *Bt* TubRC complex is required for the action of the system, we performed the assay with *Bt tubC* containing plasmid and with linear DNA representing full-length *tubC*, four repeats (sites 4–7) and two repeats (sites 6 and 7). These *Bt* TubRC complexes were compared at a constant concentration of TubR binding sites. *Bt* TubRC complexes bearing only two repeats did not have any measurable effect on TubZ polymerization below stoichiometric levels, whereas both linear DNA and plasmid containing *Bt tubC*

showed a similar substoichiometric effect. The four-repeat long *Bt* TubRC complex affected polymerization; however, this effect was substantially reduced relative to full-length *tubC*, requiring four times as much complex to match polymerization (Fig. 2F).

TubR from a *tubZR* Operon in pBM400. Having defined *tubC* in the *Bt* TubZRC plasmid partitioning system and shown that *Bt* TubRC forms a flexible DNA-protein filament, the full length of which is required to match native in vitro efficacy, we moved on to consider whether or not there might be a preferred structural mode that the TubRC complex might take up, given that the flexible TubRC complex must presumably become ordered upon binding to the TubZ filament, which might explain the requirement for the length of *tubC*.

Fortuitously, in the course of our investigation of TubZRC plasmid partitioning systems we discovered an additional TubR homolog that adopts a more consistent preferred conformation as a DNA-protein complex, shedding light on this question. *Bm* TubR (UniProt Q848W2_BACMQ) lies within an operon containing a TubZ homolog in *B. megaterium* QM B1551 (*Bm*) plasmid pBM400 (Fig. S5). The gene encoding this TubR was synthesized and we overexpressed and purified the full-length protein without any tags or additions (Fig. 3A).

***Bm* TubR Forms Ring-Like DNA-Protein Filaments.** To determine whether or not the *Bm* and *Bt* TubR-DNA complexes were comparable, we imaged the *Bm* TubR-DNA complex by electron microscopy. Whereas *Bm* TubR binds its intergenic DNA 5' to *tubZR* with a slight preference over *Bt tubC* (Fig. S3), we were unable to identify the exact *Bm* TubR binding sites by microarray (its structure suggests an explanation for this). Electron microscopy of *Bm* TubR bound to DNA revealed DNA-protein

conformation from that in *Bt* TubR; however, this region has higher B factors in both cases, and given that it is also slightly separated from the body of the protein, it may be flexible in vivo.

***Bm* TubR Helices Externalize the DNA Binding Site.** Importantly, the dimers within the crystal structure of *Bm* TubR form helical filaments (Fig. 3 *E* and *F*). The dimensions of these helices are similar to the *Bm* TubR–DNA rings visualized by electron microscopy, with a filament width of 48 Å and a maximum helical width of 155 Å. The helices place *Bm* TubR dimers back to back, allowing the wedge-shaped proteins to make significant contacts with one another. The mean buried interface between pairs of dimers is 399.5 Å² in our structure of *Bt* TubRC, but this increases to 723.8 Å² in the case of our structure of *Bm* TubR. The extended β -hairpins of adjacent dimers are paired with a small gap between each dimer, extending the groove between recognition helices around the entire helix. Significantly this superstructure displays the positively charged DNA binding surface on the exterior of these helices (Fig. 3*H*).

Discussion

***Bm* TubR Helices Are in Principle Compatible with DNA Wrapping.** The length of DNA used to produce the ring-shaped *Bm* TubR–DNA filaments resolved by electron microscopy is consistent with DNA wrapping on the external surface. Ring diameter averaged 17.5 nm, implying a circumference of 55 nm, whereas the expected length of the PCR product would be 50 nm. The internal circumference would therefore be too short, whereas the external diameter would roughly match the expected length. Given the similarity in the dimensions of the *Bm* TubR–DNA rings, and the crystallographic helix, the most parsimonious explanation is that these structures are similar. If we assume that the DNA binding mode of *Bm* TubR is identical to that of *Bt* TubR, given their structural similarity, then superimposing the available structures of this class of DNA binding proteins (*Bt* TubRC/FadR–DNA) onto the structure of the *Bm* TubR helix will describe the path of the DNA. When this is carried out, it is clear that, although the surface is not immediately compatible with a continuous strand of DNA, only a relatively small distance (6.1 Å, *Bt* TubRC/3.4 Å, FadR–DNA) would separate the DNA backbones from adjacent dimers. FadR–DNA better matches the curve of the *Bm* TubR helix as the DNA from this structure curves in the same direction. Closer pairing of the minor groove binding β -hairpins would be required to bind DNA, which might be achieved through closure of the notch in the surface of *Bm* TubR or a slight rotation of the helix (Fig. 3 *G–I* and Fig. S7).

TubRC Is a DNA–Protein Filament. Our results conclusively show that the centromeric complex of TubZRC plasmid partitioning systems forms a DNA–protein filament. The disposition of *tubC* has evolved to facilitate its formation, ensuring coverage of a minimum length of plasmid DNA; furthermore, the binding mechanism of TubR is highly cooperative, allowing coating, further favoring its formation. We have shown that such a filament is required for the activity of TubR upon TubZ polymerization in vitro, activity falling to nil as the length of the centromeric complex decreases, even though the total concentration of TubR–DNA binding pairs was held constant. It is clear, therefore, that a TubRC filament is essential for the activity of the centromeric complex. Why must this be the case?

Interaction with the centromeric complex must stabilize the assembly of its cognate filament, whereas the free adapter protein must be incapable of this activity. The formation of the complex must therefore either produce a novel binding site between its components with substantially higher affinity or cluster the adapter protein in space to achieve binding to the filament through higher avidity. In both the ParMRC and TubZRC plasmid partitioning systems, it would seem that the second mechanism appears to have won out over the first. This may well occur because this mechanism can take advantage of both the clustering of the adapter protein on an array of DNA binding

sites and clustering of the subunits within the cognate protein filament. Because the superstructure of a linear filament of identical subunits consists of a repeated vector, it must form a helix of tighter or looser curvature. Close clustering in space by a linear filament will therefore, by definition, be achieved through the formation of tight helical structures.

Although different TubZRC systems have so far been observed to have considerable similarity (25, 26), our *Bt* and *Bm* TubR complexes exhibit different curvatures. *Bt* TubRC crystallized as an extended DNA–protein filament with protein wrapping helically around the outside of the DNA, whereas the *Bm* TubR–DNA complex appears to form a ring or short helix with external DNA. Whereas it is possible that the two TubRC structures represent entirely diverged forms of the adapter complex, we think this unlikely; the *Bt* TubRC filament was clearly quite flexible in vitro, and it is in fact possible to reconcile the two structures as “flexible” and “curled” modes of a similar complex. The 12-bp repeat between major groove binding sites in the *Bt* TubRC filament is approximately conserved when DNA is superimposed upon the *Bm* TubR helix, so the binding periodicity would be similar (11–13 bp). The main structural change in DNA binding between the two would be rearrangement of the minor groove binding β -hairpins. These are roughly paired in both structures; however, the difference being that their orientation is reversed, providing a convenient switch between the two states. This proposed closure would neatly wrap the DNA and close up the space between adjacent TubR dimers (Fig. 4 *A* and *B* and Fig. S7).

Why then would the two DNA–protein filaments have different curvatures in vitro? One possible cause could be that binding

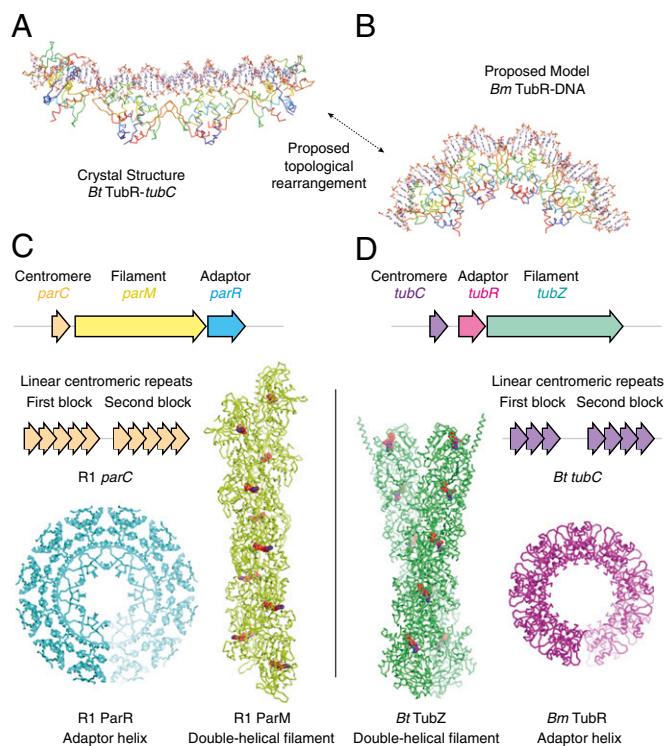


Fig. 4. *Bm* TubR helix suggests further convergent evolution of type II and III partitioning systems. (A) Structure of *Bt* TubR (α ribbon representation, blue at N terminus, red at C terminus) bound to *tubC* (stick representation, C in white/CPK colors). (B) Structure of *Bm* TubR (α ribbon representation, blue at N terminus, red at C terminus) shown with the DNA (stick representation, C in white/CPK colors) from PDB ID code 1HW2 after superimposition of the protein chains. (C and D) Comparison of operon structure (3, 10, 12), centromeric structure (20, 21), filament superstructure (9, 14, 16), and adapter complex superstructure (18, 19, this study) for the (actin-like) ParMRC and (tubulin-like) TubZRC plasmid partitioning systems.

to the TubZ filament is required to stabilize the TubRC complex in one state or another. It seems quite plausible that interactions with the linear filament stabilize the relatively flexible structure of the TubRC centromeric complex. The internally clustered TubR dimers in a ring-like structure would seem more capable of making a protein-protein contact than the sparsely distributed dimers in the flexible, extended form, as this form does not present TubR on a consistent side of the filament. The internal diameter of the *Bm* TubR crystal structure, at ~ 6.5 nm, is smaller than the diameter of the TubZ filament (14), at ~ 10 nm; however, the rings observed by electron microscopy exhibit internal diameters between 5 and 10 nm, allowing sufficient flexibility to encompass TubZ filaments in theory. A switch between a flexible and a ring-like form might allow the TubRC complex to clamp onto a growing TubZ filament during partitioning. Further structural studies of the TubZRC system will be required to understand the mechanism of binding and it is worth mentioning that the interaction sites between TubZ and TubR currently remain unknown.

Convergent Evolution of Type II and III Centromeric Complexes? It has not escaped our notice that an intriguing parallel emerges from this work. The structure of *Bt tubC* is strongly reminiscent of that of R1 *parC*, both consisting of two series of linear repeats separated by a linker, whereas the ring-like structure of *Bm* TubR is similar in superstructure to the known ParRC complex (Fig. 4C). This similarity adds to our earlier finding that TubZ, being tubulin related, forms actin- (and ParM)-like double helical filaments that are the smallest closed-symmetrical polymers possible and seem exquisitely tailored to the plasmid segregation task at hand (14). The observation of such similar superstructures,

produced to perform similar functions by entirely unrelated proteins in different systems, suggests that convergent evolution may have taken place, driving both systems toward the same solution to the problem of binding a cytomotive filament, with entirely different protein and DNA building blocks.

Materials and Methods

Detailed information is provided in *SI Materials and Methods*. Briefly, all proteins were produced in *E. coli* and purified by chromatography; DNA was either synthesized (*Bt tubC24*), produced by PCR (all other linear DNA), or purified as a plasmid (*ptubC*); the *Bt tubC* aptamer microarray experiment was performed in conjunction with LC Sciences; electron microscopy was carried out using a 2% (wt/vol) uranyl acetate negative stain in an FEI T12 electron microscope; the sequence of *Bt tubC24* was TTTAAGTTAACTTTAGTTTACA; *Bt* TubRC24 crystals were solved at 7 Å by molecular replacement and confirmed by selenomethionine SAD, whereas *Bm* TubR crystals were solved at 3.5 Å by selenomethionine SAD; 90° light scattering was carried out at 400 nm using 1.25 μ M *Bt* TubZ, 500 nM TubR, and 125 nM of TubR binding sites in each *tubC* DNA, except for *tubC* repeats 4–7, where the concentration was doubled to make the difference from *tubC24* clearly visible.

ACKNOWLEDGMENTS. We thank Rachel Larsen, Joseph Pogliano (both University of California at San Diego), and Katherine Michie [Medical Research Council Laboratory of Molecular Biology (MRC-LMB)] for providing materials; Sebastian Eustermann (MRC-LMB) for his tetraloop sequence; Fabrice Gorrec and Sonja Kuhlman for their help at MRC-LMB crystallization facility; Chen Shaoxia, Qing Wang, and Colin Palmer for their aid with electron microscopy at MRC-LMB; Sonja Dunbar for summer work and Ramona Duman (both MRC-LMB) for help at the European Synchrotron Radiation Facility (ESRF); and LC Sciences, the ESRF, and Diamond Light Source for their excellent service and support. This work was supported by Medical Research Council Grant U105184326.

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