Inside Out: Tubulin Cytomotive Filaments versus Microtubules

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In this issue of Structure, Zehr and colleagues describe a structure of a three-stranded PhuZ tubulin cytomo
tive filament determined at 8.6 Å resolution. This reveals an assembly mechanism different from that of
microtubules, leading to a hypothesis explaining cytomotive-filament dynamics.

The prokaryotic homologs of eukaryotic tubulins can polymerize into thin cytomo
tive filaments in vivo and in vitro (Löwe and Amos, 2009). The tubulin cytomatic
filaments are helical bundles of small numbers of protofilaments (Aylett et al., 2010, 2013; Kraemer et al., 2012; Pilhofer et al., 2011), in contrast to the microtubule structure in eukaryotic cells containing 13 tubulin protofilaments (tubulin molecule chains). The tubulin cytomatic filaments display nucleotide-dependent behaviors like those of microtubules in eukaryotic cells despite the difference in high-order architecture. The dynamic properties of cytomatic filaments are thought to be essential to their physiological functions in prokaryotic cells. The mechanism and structural nature underlying prokaryotic tubulin assembly and dynamics were unclear up to now. In this issue of Structure, Zehr et al. (2014) report the structural map of three-stranded cytomatic fila
ment at 8.6 Å resolution, which clearly reveals secondary structural features. The high-quality density map enabled the authors to build a pseudo-atomic model, which has provided good insight into the assembly and dynamics of prokaryotic tubulin filaments.

To understand the assembly, let’s first look into the crystal structures of the bacteriophage-encoded PhuZ tubulin-like proteins (Aylett et al., 2013; Kraemer et al., 2012; Oliva et al., 2012). The core structure of the PhuZ tubulins is very similar to that of eukaryotic tubulins. The major difference presents at their C-terminal regions. In PhuZ, the C-terminal region forms an unusually long helix (H11) followed by an extended loop (marked in green on the right side of Figure 1). In eukaryotic tubulins, however, the H11 is relatively short, and the extended loop folds back to the surface of the core structure forming the helices H12' and H12 (see the left side of Figure 1 and Figure 3E in Sui and Downing, 2010). In the new study, Zehr et al. (2014) demonstrate that the distinct C-terminal regions underlie the architectural and assembly differences between tubulin cytomatic filaments and eukaryotic microtubules. Zehr et al. (2014) first confirmed the intraprotofilament interaction (the head to tail interaction) of the PhuZ tubulin subunits is different from that in eukaryotic microtubules (Aylett et al., 2013; Kraemer et al., 2012; Nogales et al., 1999). Within a protofilament, the helix H11 and the C-terminal loop extend into the adjacent subunit. In addition, Zehr et al. (2014) note the elongated C-terminal region also defines the lateral interaction between the protofilaments in the three-stranded PhuZ filament. The elongated C-termini face inward and hold the protofilaments together as shown in Figure 1 (marked in green with springs on the right). This is completely different to microtubules where the lateral interaction between the protofilaments is defined by the H1'-S2, H2-S3 loops of one tubulin molecule and the M loop of the neighboring tubulin molecule (Sui and Downing, 2010). The C-terminal region of the eu
karyotic tubulin forms an additional helix, H12, (marked in green on the left in Figure 1) and faces outward on the microtubule. This region is responsible for binding with microtubule motors (on β-tubulins) and some microtubule associated proteins. The overall architecture of the three-stranded PhuZ cytomatic filament resembles an inverted configura
tion of microtubules with a smaller num
ber of component protofilaments.

Zehr et al. (2014) propose a C-terminal “spring model” to explain the dynamic assembly and disassembly of PhuZ cytomatic filaments. In this hypothesis, the elongated C-terminal regions are in a compacted form and tightly hold the adja
cent PhuZ tubulin molecules longitudinally in the GTP state (see Figure 7 in Zehr et al., 2014). The compacted PhuZ subunits are assembled into the twisted protofilament in the cytomatic filament. After hydrolysis, the compacted subunits cannot revert to an extended form within the helical filament lattice. Therefore, energy as strain is stored, resulting in highly dynamic metastable filaments. This intriguing hypothesis can explain the dynamic behavior for cytomatic filaments with the same concept to the “GTP cap model” for the dynamic microtubules.

There is other important information in the paper: the nucleus for the PhuZ tubulin polymerization is proposed to contain six monomers (organized into a trimer of dimers) based on the growth kinetics measured by right-angle light scattering. From the structural model, the authors identified conserved salt-bridge-forming residues. Polymerization characterization after a series of point mutations confirmed their importance in PhuZ cytomatic filament assembly. This, in turn, demonstrates the credibility of the proposed structural model determined with a density map revealing sec
ondary structural features.

The new study of three-stranded PhuZ filament provides structural insight into the assembly of prokaryotic tubulin fila
ments. Can we derive from it a universal
understanding that the C-terminal region defines the lateral interaction in all types of prokaryotic tubulin filaments? While this paper was in press, the same research group published a separate paper showing that the two-stranded filament of the bacterial tubulin TubZ-Bt can transit into a four-stranded filament upon GTP hydrolysis (Montabana and Agard, 2014). In that paper, the C-terminal regions of TubZ-Bt face inward in a two-stranded filament, which is consistent with the architecture of the three-stranded PhuZ filament structure. However, in the reported stable four-stranded TubZ-Bt filament, the protofilaments are rotated outward, exposing the C-terminal regions on the outside, an orientation similar to that in the microtubule. Based on the two papers, the lateral interactions in various cryomotive filaments of prokaryotic tubulins can be complex. Nevertheless, the changeable lateral interaction indicates that intraprotofilament interaction should be more stable than the interprotofilament interaction in cryomotive filaments of prokaryotic tubulins. Because both the protofilament number and the lateral interaction changed with nucleotide states for the TubZ-Bt filaments, the “spring model” from the PhuZ tubulin family may not represent a comprehensive view for all the prokaryotic tubulin filaments. With excellent electron microscopes and a state-of-the-art direct electron detector, better structural details of these cryomotive filaments are anticipated, which may help to develop a detailed and integrated understanding about prokaryotic tubulin assembly.

Purified prokaryotic tubulins can polymerize into a variety of filamentous structures under different conditions. Another remaining question therefore becomes: which of the reported in vitro structures naturally present in the bacteria? Resolving this issue will help to clarify if both forms of the reported lateral interactions play physiological functional roles in bacteria.

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REFERENCES


