Assembly properties of the bacterial tubulin homolog FtsZ from the cyanobacterium Synechocystis sp. PCC 6803

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The tubulin homolog FtsZ is the major cytoskeletal protein in the bacterial cell division machinery, conserved in almost all bacteria, archaea, and chloroplasts. Bacterial FtsZ assembles spontaneously into single protofilaments, sheets, and bundles in vitro, and it also accumulates at the site of division early during cell division, where it forms a dynamic protein complex, the contractile ring or Z-ring. The biochemical properties of FtsZ proteins from many bacteria have been studied, but comparable insights into FtsZs from cyanobacteria are limited. Here, using EM and light-scattering assays, we studied the biochemical and assembly properties of SyFtsZ, the FtsZ protein from the cyanobacterial strain Synechocystis sp. PCC 6803. SyFtsZ had a slow GTPase activity of ~0.4 GTP/FtsZ molecule/min and assembled into thick, straight protofilament bundles and curved bundles, designated toroids. The assembly of SyFtsZ in the presence of GTP occurred in two stages. The first stage consisted of the assembly of single-stranded straight protofilaments and opened circles; in the second stage, the protofilaments associated into straight protofilament bundles and toroids. In addition to these assemblies, we also observed highly curved oligomers and minirings after GTP hydrolysis or in the presence of excess GDP. The three types of protofilaments of SyFtsZ observed here provide support for the hypothesis that a constriction force due to curved protofilaments bends the membrane. In summary, our findings indicate that, unlike other bacterial FtsZ, SyFtsZ assembles into thick protofilament bundles. This bundling is similar to that of chloroplast FtsZ, consistent with its origin in cyanobacteria.

FtsZ, a tubulin homologue, is a crucial protein in bacterial cell division and is well-conserved in almost all bacteria, archaea, and chloroplasts. It assembles spontaneously into single protofilaments (pf), sheets, and bundles in vitro, and it accumulates in vivo at the site of division early in the cell division process to form a dynamic protein complex, the contractile ring or Z-ring (1–3).

FtsZ from Escherichia coli (EcFtsZ) assembles in vitro into mostly single pf with an average length of 200 nm (4). EcFtsZ pfS are highly dynamic, turning over with a half time of 5–8 s in vitro (5, 6) and in vivo (7). The mechanism of dynamics is now established to be treadmilling both in vitro (8, 9) and in vivo (10, 11). Treadmilling involves small patches of membrane-bound pfS adding subunits at one end and disassembling at the other.

FtsZ from different species can have different dynamic properties. The Z-ring in Bacillus subtilis has a similar dynamic turnover rate as in E. coli (7); however, the Z-ring from the slow-growing pathogen Mycobacterium tuberculosis has a slower turnover rate of 25-s half-time (12). Chloroplast Z-rings have unique and much slower dynamic properties (13–15). Chloroplasts have two copies of FtsZ, designated FtsZ1 and FtsZ2 (from green algae and plants) and FtsZA and FtsZB (from red algae). Both FtsZs co-localize in the middle of the chloroplast and are required for division (16–18). In vitro, they assemble into thick straight bundles of pfS, either separately or as a mixture (19, 20). Previous results suggested that one FtsZ might assemble as the primary scaffold and another might co-assemble to enhance polymer disassembly (20). The slow dynamics of FtsZ bundles might be related to the large diameter of the chloroplasts.

The chloroplast originated from a cyanobacterial endosymbiont and maintained FtsZ for its division. In contrast to the two FtsZs in chloroplasts, there is only one copy of FtsZ in cyanobacteria, similar to other bacteria. It should be interesting to compare the biochemical and assembly properties of cyanobacterial FtsZ with FtsZ from chloroplasts and bacteria. Here, we studied the biochemical properties of FtsZ from the cyanobacterium Synechocystis sp. PCC 6803 (SyFtsZ).

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This article contains Figs. S1–S3.

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The abbreviations used are: pf, protofilament; SyFtsZ, FtsZ from the cyanobacterium Synechocystis sp. PCC 6803; EcFtsZ, FtsZ from E. coli; SyFtsZ-TS, double mutant SyFtsZ-A271T/A274S; MtbFtsZ, M. tuberculosis FtsZ; SpnFtsZ, S. pneumoniae FtsZ; GMPCPP, guanylyl-(α,β)-methylene-diphosphonate.
Assembly properties of cyanobacterial FtsZ

Figure 1. EM images of negatively stained SyFtsZ pfs after adding 0.5 mM GTP in HMK buffer (50 mM HEPES, 100 mM KAc, 5 mM MgAc) at pH 7.5. 10 μM SyFtsZ assembled into pfs (some isolated pfs are seen in the background), which further associate into straight bundles and toroids. The diameter of the toroids is 200–300 nm. Transitions between straight bundles and toroids are indicated by arrows. Bar, 200 nm.

Results

SyFtsZ assembles into straight bundles of pfs and toroid-like circle bundles

Negative-stain EM images of SyFtsZ filaments assembled in GTP showed some isolated pfs, but mostly large bundles of pfs that were either straight or curved into toroids (Fig. 1). The diameter of the toroids ranged from 200 to 300 nm. Straight bundles and toroids were mostly isolated, but occasionally, a transition between straight and curved bundles was observed (Fig. 1, arrows). In contrast to EcFtsZ, which can assemble in the absence of Mg^{2+} (6, 21), SyFtsZ showed no assembly without Mg^{2+} or when using GMPCPP to replace GTP (Fig. S1).

Assembly was then monitored by light scattering, which showed that it proceeded in two stages (Fig. 2). The light scattering reached a final plateau at ~300 s (Fig. 2A). Amplification of the first 100 s (Fig. 2B) showed an early stage where a 15–20-s lag was followed by a rise to an intermediate plateau at 30–50 s. After 50 s, the second stage began as light scattering accelerated, leading to the final plateau. Negative-stain EM showed that the first stage resulted in the assembly of isolated short pfs and thin bundles of pfs curved into partial circles (Fig. 2C). The curved filaments consisted of apparently more than single filament, and their diameters were 200–300 nm, similar to toroid-like circle bundles, but mostly too short to form a complete circle. At the final plateau following the second stage of assembly, the EM showed predominantly large bundles of pfs and toroids (Fig. 2D).

We repeated the assembly with a limited amount of GTP (0.1 mM GTP) and found that the light-scattering signal rose to a plateau at 500 s and then slowly dropped back to the base after 1 h (Fig. 2E). This suggested that the large pbf bundles disassembled after the GTP was used up. Fig. 2F shows an assay of GTase as a function of FtsZ concentration. GTase activity was 0.41 ± 0.2 GTP/FtsZ/min above a critical concentration around 1.6 μM (Fig. 2F). This is ~10 times less than EcFtsZ but close to that of chloroplast FtsZ. At this rate of hydrolysis, the 10 μM SyFtsZ in Fig. 2E should consume the 100 μM GTP in about 30 min. However, the polymers in that experiment started to disassemble after 10 min, well before the GTP should be exhausted. Small and Addinall (22) documented for EcFtsZ that GDP inhibited assembly, suggesting that the early decrease in light scattering seen here was due to inhibition by the buildup of GDP.

Previous results showed that high salt concentration caused an increased level of bundling of EcFtsZ (6, 23, 24). We obtained similar results with SyFtsZ. Fig. 3 shows that high concentrations of potassium and magnesium accelerated the assembly and enhanced the light scattering. Without potassium or magnesium, SyFtsZ showed no assembly in pH 7.5 buffer (Fig. 3, A and B). In most experiments, the solution we used contained 100 mM KAc and 5 mM MgAc. Increasing KAc to 350 mM increased the rate and final plateau of light scattering (Fig. 3, A and C). EM showed that in addition to straight and circular bundle structures, some helical structures could be observed. The average diameter and pitch of these helical structures were about 145 and 185 nm, respectively (Fig. 3D). Polymerization of SyFtsZ is also sensitive to MgAc concentration. When the concentration of MgAc was reduced to 2.5 mM, the light-scattering signal was much weaker and had a longer lag time (Fig. 3, B and C). EM showed that there were more single pfs and thin circles and fewer straight and circular bundles (Fig. 3F). When the concentration of MgAc increased from 5 to 10 mM, the light-scattering signal increased and the lag time decreased (Fig. 3B). EM showed more straight and circular bundles (Fig. 3E).

Assembly properties of SyFtsZ change with pH

Next, we examined the assembly of SyFtsZ in buffers of different pH. The light-scattering assay showed two effects of lowering pH: an accelerated assembly and increased plateau (Fig. 4A). The assemblies displayed typical cooperative assembly kinetics with a lag time. The two-stage assemblies in pH 7.2 and pH 6.5 buffers were not as clear as we observed at pH 7.5 buffer, perhaps because the first stage was too short or too weak. EM of the pH 7.2 assembly at 40 s showed short straight bundles and toroids already assembled, mixed with short pfs and partial circles (Fig. S2). EM of assemblies at the final plateau showed a mixture of straight pbf bundles and toroids at pH 7.5, pH 7.2, and pH 6.8 (Fig. 4, B–D). However, only straight bundles were observed at pH 6.5 (Fig. 4E). The increased light scattering at reduced pH is consistent with the straight bundles being larger in diameter and/or longer.

In contrast to the large changes in light scattering and bundle morphology at lower pH, the GTase was hardly changed. The GTase activity of FtsZ was 0.41 ± 0.2 GTP/FtsZ/min at pH 7.5, 0.50 ± 0.1 at pH 7.2, and 0.36 ± 0.2 at pH 6.5 with critical concentrations of 1.5–2 μM (Fig. 4F).

SyFtsZ assembled into highly curved oligomers and miniring structures in the presence of GDP

Previous studies have shown that FtsZ from E. coli and Pseudomonas aeruginosa could assemble into highly curved short oligomers or miniring structures with an average diameter of 23 nm in the presence of GDP (25–28). Minirings were especially favored in the presence of ZipA protein (27). We also found highly curved oligomers and miniring structures in the EM images of SyFtsZ pfs during GTP hydrolysis (Fig. 5A) or in the presence of GDP (Fig. 5, B and C) in both pH 7.5 and pH 6.5 solutions. The diameters of these highly curved pfs or minirings were ~20 ± 2 nm. Sedimentation results confirmed assembly...
of these oligomers. Using a Beckman TLA-100 rotor, we selected two centrifuge speeds: 50,000 rpm and 90,000 rpm. Only large bundles assembled in GTP could be spun down at 50,000 rpm (around 100,000 \( \times g \)); in contrast, both bundles in GTP and partly short oligomers and/or minirings could be spun down at 90,000 rpm (around 310,000 \( \times g \)). Fig. 5 (D and E) and Fig. S3 show the sedimentation results analyzed by SDS-PAGE. At 50,000 rpm, there was little protein in pellet with GDP. At 90,000 rpm with GDP, 21% of FtsZ was pelleted at pH 7.5 at both 5 and 10 \( \mu M \) FtsZ, and 49% of 10 \( \mu M \) FtsZ was pelleted at pH 6.5. The pellets are consistent with the short oligomers and minirings observed by EM.

**Mutant AXXA to TXXS in the H8 helix reduces bundle formation and enhances GTPase activity**

FtsZ comprises two globular subdomains, which are connected by a central core helix (H7 helix) and a synergy loop (T7 loop). The T7 loop is located between the H7 helix and H8 helix, and it reaches into the active site of an adjacent subunit to activate its GTPase activity (29–31) (Fig. 6A). The amino acids Asn-207, Asp-209, and Asp-212 of EcFtsZ are highly conserved in all FtsZs. Mutation of Asp-209 in the T7 loop or Asp-212 in the H8 helix of EcFtsZ severely impaired its GTPase activity and function (32, 33). Following the very conserved sequence DFADV(R/K), we have identified two amino acids in the H8

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**Figure 2.** The assembly of SyFtsZ assayed by light scattering shows two stages in HMK buffer at pH 7.5 (A and B). A shows that light scattering reaches a final plateau at 300–600 s. B amplifies the first 100 s and shows that light scattering first approaches an intermediate plateau at 30–50 s and then accelerates to the much higher final plateau. Note that light scattering does not measure polymer, but is greatly enhanced by bundling. The signal here is primarily a measure of bundle formation. C and D, EM of 10 \( \mu M \) SyFtsZ shows short single pfs and partial circles at the first stage assembly, at 30 s (C), and assembly into straight bundles and toroids at the second stage of assembly, at 300 s (D). Bar, 200 nm. E, light-scattering assay of 10 \( \mu M \) SyFtsZ in HMK buffer assembled in 0.1 mM GTP. Light scattering reached a plateau at 500 s and then slowly decreased to the base after 1 h. F, the GTPase activity of SyFtsZ in HMK buffer was 0.41 ± 0.2 GTP/min/FtsZ above a critical concentration of 1.6 \( \mu M \). Error bars, S.D.
helix that are conserved as Thr and Ser in fast-growing bacteria (Thr-215 and Ser-218 of EcFtsZ, or Thr-216 and Ser-219 of FtsZ from B. subtilis). In most cyanobacteria and chloroplast FtsZs, the corresponding amino acids are Ala (Ala-271 and Ala-274 of SyFtsZ). Interestingly, there are several cyanobacterial FtsZs, such as FtsZ from Prochlorococcus and Synechococcus, where these are still Ser and Thr. Both of these belong to marine picocyanobacteria, whose cell diameter is less than 1 μm. This size is similar to that of bacteria and much smaller than other cyanobacteria. Also, some slow-growing bacterial FtsZs have only one Thr or Ser. For example, in M. tuberculosis FtsZ (Mtb-FtsZ), the first Thr is changed to Gly, and in S. pneumoniae FtsZ (SpnFtsZ), the second Ser is changed to Ala. Previous results suggested that both MtbFtsZ and SpnFtsZ assembled into small bundles. MtbFtsZ slowly assembled into two-stranded pfs at pH 6.5 (12, 34), and SpnFtsZ assembled into mostly double- or multiple-strand pfs, with a high critical concentration around 3–6 μM (35, 36).

To address whether these two amino acids are related to bundle formation, we constructed the double mutant SyFtsZ-T215A/S218A (SyFtsZ-TS). The GTPase activity of this mutant increased ~80%; from 0.4 to 0.7 GTP/FtsZ/min in pH 7.5 buffer (Fig. 7A) and from 0.5 to 0.9 GTP/FtsZ/min in pH 7.2 buffer (Fig. 7B). The increased GTPase of the mutant correlates with reduced bundle formation, suggesting that bundling results in reduced subunit turnover and GTPase. Fig. 7 (C and D) highlights the difference of their assembly properties measured by the light scattering at different pH. In pH 7.5 solution, the assembly of mutant SyFtsZ only had the fast initial stage in contrast to the two-stage assembly curves of WT SyFtsZ assembly (Fig. 7C; compare with Fig. 2B). Negative-stain EM confirmed that the assembled pfs of SyFtsZ-TS were mixtures of single pfs and thin straight and curved bundles, including some closed circles (Fig. 7E). There were no large bundles in pH 7.5 buffer for FtsZ up to 15 μM. In pH 7.2 solution, the assembly showed two stages, but they were much slower than WT (Fig. 7D). The first stage of assembly, which plateaued from 50 to 200 s, showed short pfs and circles typical of this stage (Fig. 7F). The second stage started around 200 s, rose to a much higher light-scattering signal, and approached a plateau at 900 s (Fig. 7D). EM showed mostly thick straight bundles and a few circles at pH 7.2 (Fig. 7G), and no toroid-like circular bundles were observed. Evidently, mutant proteins reduced bundle formation.

We then tested the effect on EcFtsZ of changing the Thr and Ser to Ala. We constructed two single mutants (EcFtsZ-T215A and EcFtsZ-S218A) and a double mutant EcFtsZ-T215A/S218A. Negative-stain EM showed a modest increase in bundles assembled for the mutant EcFtsZ (Fig. 8A–D). These were mostly double pfs or small bundles, and no large bundles were seen. These modest bundling filaments were similar to the assembly of MtbFtsZ (12, 34) and SpnFtsZ (35), which had one of the two amino acids changed. Interestingly, GTPase activities of mutant EcFtsZ were only slightly changed (Fig. 8E). The GTPase was reduced from 4.6 to 3.5 GTP/FtsZ/min for EcFtsZ-S218A and little changed for EcFtsZ-T215A and EcFtsZ-T215A/S218A. The critical concentration increased from 1 to around 2.5 μM for EcFtsZ-T215A and EcFtsZ-T215A/S218A.

Discussion

Perhaps the most striking observation in the present work was that SyFtsZ assembled into thick bundles of pfs that could
be either straight or curved into a toroid of 200–300-nm diameter. Assembly of similar straight and curved pf bundles was reported previously for EcFtsZ and MtbFtsZ, but only in the presence of crowding agents polyvinyl alcohol or methylcellulose (37, 38). A recent study of FtsZ from the cyanobacterium Anabena showed a mixture of straight pf bundles and toroids essentially the same as those we show here for SyFtsZ. In that study, removal of the 51 N-terminal amino acids (this N-terminal peptide is much longer in cyanobacteria than the 9–15 amino acids in most bacteria) eliminated the toroids but left the thick straight pf bundles. FtsZ from the chloroplasts of Galdieria sulfuvaria, GsFtsZA and GsFtsZB, also formed thick straight bundles. In this case, the assembly used the globular domains only, truncating the N- and C-terminal extensions. The morphology of SyFtsZ bundles, straight versus toroids versus helices, is sensitive to the pH value and salt concentration. Similar bundle morphologies have been observed for EcFtsZ assembled in the presence of crowding agents (38). Because the internal pH of cyanobacterial cells is above 7, SyFtsZ should assemble preferentially curved filaments and bundles in vivo.

The thick bundles seem to be a property of cyanobacterial FtsZ and the derivative chloroplast FtsZ. However, the relevance of the bundling for in vivo function is not clear. It is noted that the diameter of cyanobacterium Synechocystis sp. PCC
6803 is around 2–3 μm, and the chloroplast average of 5 μm is much larger than the ~1 μm of the bacterial cell. These results are consistent with previous studies of chloroplast FtsZ (20), suggesting that bundling of FtsZ might benefit the cell division of large cells. The in vitro bundles appear to be three-dimensional, but because all FtsZ subunits have the C-terminal peptide that mediates membrane tethering, the in vivo bundle may be limited to a two-dimensional ribbon, with all pfs tethered to the membrane. The amount of FtsZ in cyanobacteria has not been determined, but in other bacteria, it is typically present at ~5,000 copies (2). This could make a ribbon of 7 pfs encircling the 1-μm diameter in E. coli. Where only 30% of the FtsZ is in the Z-ring, the ribbon would average only 3 pfs wide. There is not enough FtsZ in a single cell to make a large bundle.

We found that bundling was reduced significantly when we mutated Ala-271 to Thr and Ala-274 to Ser, which were the amino acids at the corresponding positions in E. coli and other rapidly dividing bacteria. Interestingly, in the FtsZ from marine picocyanobacteria, such as Prochlorococcus and marine Synechococcus, these amino acids are still Ser and Thr, suggesting that they may reduce the bundling activity. Those marine picocyanobacteria have small sizes with diameters of less than 1 μm, compared with the 2–3-μm diameter of Synechocystis. Previous results revealed that marine Synechococcus strains and Prochlorococcus were sister clades, which are distantly related to other cyanobacteria, including freshwater Synechococcus strains, such as Synechococcus elongates (39–42). This is consistent with our analysis. One possibility is that the intrinsic bundling activity is useful for the stable Z-ring formation and benefits from the larger diameter of most cyanobacteria and chloroplasts. The biochemical properties of marine picocyanobacterial FtsZ need to be further investigated. Also, we do not have in vivo data to support this possibility, so further investigations need to be conducted.

The physiological function of the circle bundles of cyanobacterial FtsZ is still unclear. We observed that SyFtsZ assembled into both straight and circular bundles when pH levels were above 6.8; meanwhile, only straight bundles assembled when pH was 6.5. It was reported that cyanobacterial cells maintained a narrow range of survivable intracellular pH of 7.1–7.3 even when they were exposed to a low, growth-inhibiting external pH below 6 (43). This indicates that the FtsZ circles might be useful for its function. Recently, Corrales-Guerrero et al. (44) reported that deletion of N terminus of Anabaena FtsZ eliminated the circular bundles and produced only straight bundles in vitro. The mutant was nonfunctional as the sole source of FtsZ and was toxic when co-expressed with WT FtsZ. These
data suggest that formation of circular bundles might be necessary for its function; however, the possibility that deletion of the N terminus affects binding of other association proteins cannot be excluded \((44)\).

The 200–300-nm curvature of the toroids has been designated the intermediate curved conformation \((45)\). Osawa and Erickson \((46)\) have suggested that this intermediate curved conformation, rather than the highly curved miniring, may be responsible for generating the constriction force on the membrane. It is not clear what triggers the transformation from straight to intermediate curved. GTP hydrolysis is not required, because intermediate curved pf s can be obtained in nonhydrolyzable GMPCPP and by the EcFtsZ mutant D212A \((2)\). Also, Popp et al. \((37)\) observed toroids and spiral pf bundles assembled in GMPCPP and in GTP plus EDTA. Previous studies have provided evidence that a pf bundle can transition from straight to curved, in a process called spooling \((47, 48)\). A transition from straight to curved pf s has been invoked for treadmilling pf
patches (49). Fig. 1 provides some of the clearest images of the spooling transition for SyFtsZ.

Another type of curved pf is defined as highly curved pfs and minirings. The miniring with a diameter around 20–30 nm was first reported when EcFtsZ was assembled onto cationic lipid monolayers without GTP or after GTP hydrolysis (25, 50). The miniring structures of EcFtsZ could also be observed in solution (27, 28, 51), especially following the addition of ZipA (27), a FtsZ-associated protein conserved in only /H9253-proteobacteria. Similar miniring structures were further observed in the studies of Caulobacter crescentus FtsZ when stabilized by the regulator protein FzlA (52) and B. subtilis FtsZ with the addition of ZapA (53). It is concluded that GTP favored the assembly of straight or intermediate curved pfs, and GDP favored highly curved pfs. It is possible that these two curved conformations of FtsZ together provide a continuous force to bend and constrict the membrane at the site of cell division.

Experimental procedures
Plasmid construction and protein purification

cDNA for SyFtsZ from cyanobacterium Synechocystis sp. PCC 6803 was inserted into pET15b using PCR and engineered Ndel/BamHI digestion sites. Mutants SyFtsZ-TS, EcFtsZ-T215A, EcFtsZ-S218A, and EcFtsZ-T215A/S218A were generated by point mutagenesis.

Plasmids containing SyFtsZ or EcFtsZ were transformed into BL21 E. coli and purified. Briefly, the soluble His6 proteins were first purified using a Talon column (Clontech Lab, Inc.). After elution with 250 mM imidazole, the samples were concentrated using an Amicon Ultra-15 centrifugal filter (Millipore Sigma) and exchanged into LTK50 buffer (50 mM Tris, pH 7.5, 50 mM KCl, 1 mM EDTA, 10% glycerol). Thrombin was added to 20 units/ml and incubated at room temperature for 2–4 h to remove the His6 tag. After digestion by thrombin, three amino acid residues, Gly-Ser-His, remain at the N terminus, which should have limited impact on assembly. Further purification was over a Resource Q column (GE healthcare) with a linear 50 mM to 100 mM KCl gradient in LTK buffer. Purified proteins were stored at −80 °C. Protein concentrations were determined by using a BCA assay with a corrected factor for a presumed 70% color ratio (54). Proteins were thawed and dialyzed into HMK buffer (50 mM Hepes, 100 mM KAc, 5 mM MgAc) for assembly at pH 7.5 and 7.2 or MMK buffer (50 mM MES, 100 mM KAc, 5 mM MgAc) for assembly at pH 6.8 and 6.5.

Electron microscopy

Negative-stain EM was used to visualize FtsZ polymerization. FtsZ proteins were incubated with GTP or GDP for 2–10 min. Then ~10 μl was applied to a carbon-coated copper grid and quickly withdrawn. Grids were immediately stained with 2% uranyl acetate, and specimens were imaged with a Philips 420 electron microscope.

Light-scattering measurement

The kinetics of FtsZ assembly was measured by a light-scattering assay using a Shimadzu RF-5301 PC spectrofluorometer, with both excitation and emission at 350 nm as described previously (20, 55). The light-scattering assay is especially sensitive to the assembly of large bundles. Each measurement was repeated two or three times, with consistent results.

Sedimentation assay

Assembly of SyFtsZ was also assayed by sedimentation. 5 or 10 μM SyFtsZ was incubated with or without 1 mM GTP in

Figure 7. Comparison of assembly properties of WT SyFtsZ and SyFtsZ-TS (SyFtsZ-A271T/A274S). A and B, the double mutant increased the GTPase somewhat at both pH 7.5 (A) and pH 7.2 (B). C, light scattering suggests that the SyFtsZ-TS at pH 7.5 assembled only to the first stage and was missing the large rise in turbidity attributed to the second stage (cf. Fig. 2). D, at pH 7.2, the first stage of assembly was prolonged, but the second stage rise in turbidity began at ~300 s. E–G, EM confirms that SyFtsZ-TS only assembles into single straight pfs and thin curved bundles at pH 7.5, 300 s (E) and at pH 7.2, 100 s (F). These assemblies are characteristic of stage 1 (see Fig. 2). At pH 7.2, 600 s large thick bundles characteristic of stage 2 are seen (G). Bar, 200 nm. All buffers contained 50 mM HEPES, 100 mM KAc, 5 mM MgAc, and 0.5 mM GTP.

Assembly properties of cyanobacterial FtsZ
HMK buffer, pH 7.5, or MMK buffer, pH 6.5, at room temperature for 10 min and centrifuged at 50,000 or 90,000 rpm for 30 min at 25 °C in a Beckman TLA100 rotor. The supernatant was carefully removed, and the pellet was resuspended in the same volume of buffer. The protein in the pellet and supernatant was assayed by 10% SDS-PAGE. The ratio of supernatant and pellet was measured using ImageJ software.

GTPase measurement

GTP turnover was measured after the assembly had reached its plateau of light scattering. The GTPase turnover was measured using a regenerative coupled GTPase assay as described previously (55, 56). Our assay mixture included 1 mM phosphoenolpyruvate, 0.8 mM NADH, 20 units/ml pyruvate kinase and lactate dehydrogenase (Sigma-Aldrich), and 0.5 mM GTP. After GTP hydrolysis by FtsZ, GDP in solution is rapidly regenerated into GTP, in a reaction that consumes one NADH per GDP. The NADH concentration was measured by the absorption at 340 nm in a Shimadzu UV-2401PC spectrophotometer, using the extinction coefficient 6,220 M⁻¹ cm⁻¹. The absorbance showed a linear decrease over time, giving the hydrolysis rate for each FtsZ concentration. These rates were plotted versus FtsZ concentration, and the overall hydrolysis rate in GTP/min/FtsZ was the slope of the line above the critical concentration. Each assay was repeated two or three times.

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