GTP Binding Induces Filament Assembly of a Recombinant Septin

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Summary

The septins are a family of GTPases involved in cytokinesis in budding yeast, Drosophila, and vertebrates (see [1] for review). Septins are associated with a system of 10 nm filaments at the S. cerevisiae bud neck, and heteromultimeric septin complexes have been isolated from cell extracts in a filamentous state [2–4]. A number of septins have been shown to bind and hydrolyze guanine nucleotide [2, 5, 6]. However, the role of GTP binding and hydrolysis in filament formation has not been elucidated. Furthermore, several lines of evidence suggest that not all the subunits of the septin complex are required for all aspects of septin function [3, 7]. To address these questions, we have reconstituted filament assembly in vitro by using a recombinant Xenopus septin, XI Sept2. Filament assembly is GTP dependent; moreover, the coiled-coil domain common to most septins is not essential for filament formation. Septin polymerization is preceded by a lag phase, suggesting a cooperative assembly mechanism. The slowly hydrolyzable GTP analog, GTP–γ–S, also induces polymerization, indicating that polymerization does not require GTP hydrolysis. If the properties of XI Sept2 filaments reflect those of native septin complexes, these results imply that the growth or stability of septin filaments, or both, is regulated by the state of bound nucleotide.

Results and Discussion

To obtain large amounts of septin protein, a Xenopus septin, XI Sept2, was expressed in E. coli, and the protein was purified by affinity column chromatography. The purity of the column eluate exceeded 90%, as judged by Coomassie-stained acrylamide gels (Figure 1A). Furthermore, the preparation ran as a single peak of apparent molecular weight of 87 kDa on a gel filtration column, compatible with it being a monomer or a very stable dimer (Figure 1B). Like all known septins, XI Sept2 contains a P loop motif and other sequences characteristic of GTP binding proteins [8]. The human gene most related to XI Sept2 is now known as Sept2 (BAA05893) [9]; that of budding yeast is Cdc3p. No protein bound nucleotide was detected in our septin preparations (as assessed by HPLC over a MonoQ column; our unpublished data). We therefore tested the ability of recombinant XI Sept2 to bind and hydrolyze exogenous guanine nucleotide. Using a filter binding assay, we found that XI Sept2 bound to radiolabeled GTP, whereas the BSA control showed no detectable binding (Figure 1C). Based on these experiments, we estimate that 40%–50% of the septin in our preparation is competent for GTP binding. Binding is specific for GTP; unlabeled GTP, but not ATP or GDP, efficiently compete with the radiolabeled guanine nucleotide. Unlabeled GTP–γ–S competed for binding as efficiently as GTP (Figure 1E). The rate of GTP binding is slower than would be expected for binding of a small molecule. This may indicate that a conformational change is required for the protein to bind nucleotide; it is also possible that although the recombinant protein is soluble, it is not perfectly folded and the slow binding is due to partial refolding. To assess nucleotide hydrolysis, we incubated XI Sept2 with [α–32P]GTP or GTP–γ–S, and the production of radioactive GDP or free thiophosphate was followed by thin layer chromatography (TLC). GTP, but not GTP–γ–S, was efficiently hydrolyzed (Figure 1D). On the other hand, addition of 10 units of alkaline phosphatase to the GTP–γ–S-containing sample resulted in >70% hydrolysis within 60 min (unpublished data). Thus, XI Sept2 readily binds and hydrolyzes exogenous GTP.

Filament-forming proteins such as actin, tubulin, and FtsZ polymerize in a nucleotide-dependent manner. To determine whether XI Sept2 can assemble into filaments in the presence of guanine nucleotide, we incubated the protein with GTP or GTP–γ–S and subsequently adsorbed it to glass coverslips. The coverslips were then fixed and stained with anti-XI Sept2 antibody. Under these conditions, immunofluorescence microscopy revealed filamentous structures of up to 7 μm in length (Figure 2A). These filaments were of uniform intensity along their lengths, suggesting a constant width.Filaments were not observed when XI Sept2 was incubated with ATP or GDP (Figure 2A). The structure of XI Sept2 filaments was examined by transmission electron microscopy (Figure 2B). Short filaments (with a length between 200 and 1000 nm) could be detected in reactions that had been incubated with GTP for 5 min, whereas longer incubation periods resulted in filaments >1000 nm long (Figure 2B, upper left panel). The width of the filaments was estimated to be approximately 20 nm. In the absence of added nucleotide or in the presence of ATP, small amorphous aggregates were observed, confirming that filament formation requires GTP (Figure 2B, upper right). At high magnification, a narrow region of electron-dense material was visible along the filament lengths (Figure 2B, lower left), suggesting that the 20 nm-wide structures are composed of two 8–9 nm-wide filaments in close apposition. This can be compared with the reported width (7–9 nm) of septin filaments.
Figure 1. *Xl* Sept2 Binds and Hydrolyzes GTP

(A) A Coomassie-stained acrylamide gel showing affinity-purified *Xl* Sept2. The major band corresponds to the expected *Xl* Sept2 molecular weight of 41 kDa.

(B) Gel filtration of purified *Xl* Sept2 and a set of molecular weight standards (Pharmacia) on a Superdex 200 column.

(C) [*H]GTP binding of *Xl* Sept2 and Δcc-Sep2, as determined by a filter binding assay. A total of 1 μM septin was incubated with 4.4 μM GTP and 5 mM MgCl2.

(D) Nucleotide hydrolysis by *Xl* Sept2 and Δcc-Sep2, as assayed by TLC. BSA was used as a negative control in both experiments.

(E) Competition of GTP binding. A total of 200 pmoles *Xl* Sept2 (1 μM) was incubated with 4.4 μM [*H]GTP and the indicated amounts of nonlabeled GTP, GTP–γ–S, GDP, or ATP.

purified from *Drosophila* and yeast cell extracts [2, 3]. Moreover, structures reminiscent of the paired filaments described here have been observed when the purified yeast septin complex is dialyzed into a low-salt buffer; in this case, filaments within a doublet were spaced by a variable (2–20 nm) gap without sign of a bridging structure [3]. A minority of *Xl* Sept2 filaments associated laterally to form small bundles (Figure 2B). The presence of guanine nucleotide is therefore sufficient to induce assembly of a single septin into filamentous structures. Many (but not all) of the known septins contain C-terminal sequences predicted to form a coiled-coil. However, the importance of these domains in septin function(s) has not been addressed previously. We have generated a septin protein lacking the coiled-coil domain (Δcc-Sep2) and have characterized its GTPase- and filament-formation activities. Δcc-Sep2 maintained GTPase activity, albeit with a reduced GTP binding efficiency (Figures 1C and 1D). This could indicate that a larger fraction of the truncated protein fails to fold correctly in the bacterial cell as compared to the full-length septin. Importantly, purified Δcc-Sep2 was able
In order to gain insights into the septin polymerization mechanism, we examined the kinetics of XI Sept2 filament assembly by right-angle light scattering. This method has been used for studying the polymerization of filament-forming proteins such as actin and FtsZ. It has been shown that the intensity of scattered light is linear relative to polymer mass and that this relationship is largely independent of filament length [10–12]. Scattered-light measurements are sensitive to changes in filament width that could be caused by filament bundling (e.g., [13]; however, light and electron microscopy of septin filaments indicate that only a minor fraction of septin filaments are bundled [see Figure 2]). When GDP or ATP was added to XI Sept2, the level of scattered light remained stable for the entire duration of the experiment over a wide range of XI Sept2 concentrations (1–20 μM). Moreover, coassembly between full-length XI Sept2 and a biotinylated form of Δcc-Sep2 was also observed by immunofluorescence, for which anti-XI Sept2 antibody was used in conjunction with fluorophore-coupled streptavidin (our unpublished data). When polymerized under equivalent conditions, Δcc-Sep2 filaments were less abundant than XI Sept2 filaments (compare GTP panels in Figures 2A and 3A). Thus, the filament formation ability of XI Sept2 is diminished, but not abolished, by deletion of the coiled-coil domain.
Figure 4. Quantification of *Xl* Sept2 Filament Assembly by Right Angle Light Scattering
(A) Time courses of filament assembly in the presence of 0.5 mM GTP and GTP-γ-S at different *Xl* Sept2 concentrations. After nucleotide addition, the cuvette was repositioned in the spectrometer at time 0 min.
(B) Estimation of the critical concentration for filament assembly in the presence of 0.5 mM GTP and GTP-γ-S.
(C) Analysis of nucleotide bound to septin filaments. Septin polymer was isolated by sedimentation, and the nucleotide present in the total reaction and that bound to polymer were extracted and analyzed by TLC. Since the pellets were too fragile to wash, some nucleotide was recovered in the negative control; however, significantly more nucleotide pelleted in the presence of *Xl* Sept2 polymer.
(D) A Coomassie gel of the sedimentation reaction demonstrating that sedimentation of septin is GTP dependent.

ization mechanism, in which initiation of filament assembly is the rate-limiting process and is followed by a relatively faster elongation of polymer nuclei [14].

We sought to examine the role of GTP hydrolysis in the polymerization kinetics of *Xl* Sept2. To this aim, we used the GTP analog GTP-γ-S, which binds to *Xl* Sept2 with the same affinity as GTP but fails to be hydrolyzed (see Figures 1D and 1E). The use of GTP-γ-S in polymerization reactions did not significantly alter the length of the lag phase (compare plots in Figure 4A). However, equal concentrations of *Xl* Sept2 yielded a higher degree of polymerization at steady state in the presence of GTP-γ-S than in the presence of GTP. Hence, the equilibrium between monomeric and polymeric septin appears to be shifted toward polymer formation when nucleotide hydrolysis is inhibited. Although filament bundling could also account for this relative increase, EM analysis did not indicate a significant increase in bundling in the presence of GTP-γ-S (Figure 2). Several alternative mechanisms could account for the differences seen between GTP- and GTP-γ-S-induced polymerization. Septin filaments bound to GDP could be less stable than filaments bound to GTP (or GTP-γ-S). Additionally, monomeric GDP bound *Xl* Sept2 may accumulate in the reaction and have a lower affinity for filament ends than GTP bound monomer. Indeed, we have found that under the conditions used to measure polymerization in Figure 4A, up to 70% of *Xl* Sept2 is bound to GDP 60 min after nucleotide addition (see Experimental Procedures). Therefore, our data are compatible with either of these two mechanisms (or both) accounting for the different polymerization capabilities of *Xl* Sept2 in the presence of GTP and GTP-γ-S. Importantly, these data confirm that filament assembly does not require GTP hydrolysis.

Although *Xl* Sept2 can polymerize in the presence of GTP-γ-S, which is not hydrolyzed to a detectable extent, the data in Figure 1 indicate that *Xl* Sept2 does hydrolyze GTP. Therefore, we determined the state of the nucleotide bound to polymeric *Xl* Sept2. Septin filaments that were assembled in the presence of 32P-γ-GTP (25 μM Sept2, 500 μM GTP) were sedimented through a glycerol cushion to isolate polymeric *Xl* Sept2. SDS-PAGE analysis showed that under these conditions septin protein sediments in a GTP-dependent manner (Figure 4D). The nucleotide bound to polymeric septin was extracted and resolved by TLC. We found that the GDP:GTP ratio was 2:1 both in the unfractionated septin reaction and in the fraction containing polymeric septin (Figure 4C). Although significant amounts (30%) of GMP were also present in the unfractionated reaction, no GMP was present in the pellet. Thus, both GDP and GTP are present in polymeric septin. These data further confirm that...
poymerization is not directly coupled to nucleotide hydrolysis, which is consistent with the observation that GTP-γ-S also supports polymerization.

If XI Sept2 forms nucleated polymers, there ought to be a "critical concentration" for filament assembly. This is defined as the concentration below which nucleation cannot occur and only monomers (and a small number of short oligomers) exist. Above the critical concentration, nucleated polymers assemble and are in equilibrium with the monomeric form. The most reliable method of measuring the critical concentration is to dilute preassembled polymer to a variety of concentrations and to determine the concentration below which all filaments disassemble. This experimental procedure is preferred over initiating assembly at different concentrations because near the critical concentration, the long lag phases and low elongation rates make it difficult to assess when steady state has been reached. We therefore assembled XI Sept2 filaments to steady state in the presence of either GTP or GTP-γ-S, diluted them to different protein concentrations, and let them equilibrate to a new steady state. Measurements of the polymer content by light scattering showed that filaments were still present at all the concentrations tested (0.8–4.2 μM). The data points fitted a linear plot that, when extrapolated to zero polymer content, predicts a critical concentration of 0.2 μM and 0.07 μM in the presence of GTP and GTP-γ-S, respectively, although these are rough estimates (Figure 4B). Because of the low signal-to-noise ratio at lower protein concentrations, we were unable to directly measure the critical concentration with this assay. However, we have followed depolymerization by indirect immunofluorescence of XI Sept2 filaments and have established that septin filaments assembled in the presence of GTP or GTP-γ-S and diluted below 0.05 μM disassemble; in the case of GTP-assembled filaments, filaments are no longer detectable after a few minutes at room temperature, suggesting that the critical concentration is different from zero (our unpublished data).

Conclusions

In this paper, we have shown that a single vertebrate septin assembles into filaments upon addition of GTP. This finding has been documented by immunofluorescence, electron microscopy, and light-scattering measurements. Filament assembly does not require GTP hydrolysis; indeed, the extent of filament assembly is greater in the presence of GTP-γ-S. In addition, we show that filament assembly is stimulated by, but not wholly dependent on, the coiled-coil domain that is found in most, but not all, septin proteins. Finally, kinetic analysis of polymerization suggests that septin filaments assemble as a nucleated polymer.

Biochemical studies have established that septins exist in heteromultimeric complexes [2–4]. In budding yeast, however, the CDC10 septin can be deleted without compromising cell viability [3]. Furthermore, in Drosophila, though two additional septins copurify with the Pnut septin, recent evidence indicates that Sep2, but not Sep1, still localizes to the leading edge of cleavage furrows in the absence of Pnut [7]. Therefore, not all the subunits in the septin complex are required for all aspects of septin function in these organisms. In this paper, we have shown that a single septin can efficiently polymerize in vitro. Thus, filament assembly in vitro does not require septin heteromultimers. Whether a single septin subunit can polymerize in vivo remains to be determined. It is conceivable that the general properties of septin polymerization that we have described by examining a single subunit may be generally applicable to the biochemistry of septin heteromultimers, though future experiments are necessary to establish this point.

The relationship between septin polymerization and septin function has been addressed previously. Cells carrying a deletion of the CDC10 septin gene maintain many septin-dependent functions (localization of Bni4p, Bud4p, and the other septins to the bud neck, efficient cytokinesis) but lack neck filaments. Although the relationship between neck filaments and septin filaments is not entirely clear, this result indicates that highly ordered arrays of septin filaments are not required for septin function [3]. In vitro experiments showed that septin complexes purified from wild-type yeast cells associate into long, paired filaments upon dialysis into low-salt buffer. On the other hand, complexes purified from cells lacking the septin Cdc10p failed to assemble into filaments under the same conditions [3]. These data may indicate that septin function may not require septin filament assembly [1, 3]. Alternatively, septin complexes lacking Cdc10p could assemble into filaments that do not form highly ordered arrays in vivo and are incompetent to assemble under the conditions used in these in vitro experiments. The evolutionary conservation of the motifs required for GTPase activity strongly suggests that this activity is required for septin function. In this report, we have presented evidence that GTP binding, though not GTP hydrolysis, is intimately linked to septin polymerization. It is therefore likely that GTP-dependent filament assembly is critical for septin function in vivo.

Although the results presented here shed light on some aspects of septin filament assembly, many key issues remain to be addressed. Most importantly, it is still unclear how the filamentous state of septins correlates with their biological function. The ability to generate septin mutants, characterize their ability to assemble into filaments in vitro, and then monitor the phenotype of such mutations in vivo should help elucidate this problem.

Experimental Procedures

Purification of XI Sept2

The coding sequence of XI Sept2 (GenBank accession number AF212298) and the coding sequence minus the predicted coiled-coil domain (amino acids 309–356; identified according to [15]) were cloned in the pTYB4 vector (NEB) as an N-terminal in-frame fusion with the intein-CBD (chitin binding domain) coding sequence. Extracts of BL21(DE3) cells expressing the XI Sept2-intein constructs were loaded into a chitin-agarose column, and affinity purification was performed according to the manufacturer’s instructions. After DTT-induced cleavage of the intein-CBD moiety, the protein was eluted from the column, dialyzed against buffer S (20 mM Tris [pH 8.0], 20 mM NaCl, 20% glycerol, 1 mM MgCl2, 0.1% Triton X-100), and stored at –80°C at a concentration of 1–2 mg/ml.

GTP Binding and Hydrolysis Assays

For GTP binding, 1 μM XI Sept2 (or Δcc-Sep2) in S buffer was incubated with 0.1 mg/ml BSA, 5 mM MgCl2, and 4.4 μM [3H]GTP.
Filament Immunofluorescence and Negative Staining For GTP hydrolysis, 1/2 "H[GTP and various concentrations of unlabeled GTP, GDP, GTP-γ-S, and ATP for 60 min at room temperature before filtering as described above. For GTP hydrolysis, 1 μM XI Sept2 was incubated with [γ-32P]GTP (800 Ci/mmol, 10 μCi/ml) or GTP-γ-[35S] (1333 Ci/mmol, 1 μCi/ml, Amersham) as described above. At time intervals, aliquots were diluted in denaturing solution (1% SDS, 20 mM EDTA), heated at 65°C for 10 min, and spotted on a PEI-cellulose plate (Macherey-Nagel). TLC plates were developed in 1 M LiCl, air-dried, and GTP, GDP, and Pi spots were quantified by phosphorimaging. The fraction of hydrolyzed nucleotide was calculated as GDP/(GTP + GDP) or Pi/(GDP + Pi). Nucleotide Rf values were confirmed by visualizing unlabeled standards with UV light. The mobility of the thio-phosphate group released from GTP-γ-[35S] was confirmed by treatment with alkaline phosphatase (Boehringer) and phosphorimaging.

In order to identify the nucleotide bound to XI Sept2, [γ-32P]GTP was included in the binding assays, the NC filters were treated with denaturing solution, and the released nucleotide was analyzed by TLC as described above.

Analysis of Nucleotide Associated with Polymeric Septin Precleared XI Sept2 (25 μM) was incubated with [γ-32P]GTP (500 μM) for 60 min. Assembly reactions (25 μl) were centrifuged through 100 μl of a 60% glycerol cushion (279,000 × g, 20 min at 4°C). The supernatant and cushion were aspirated, and the pellet was resuspended in 1% SDS/20 mM EDTA. Totals and pellet fractions were analyzed by SDS-PAGE, followed by Coomassie staining and by TLC described as above.

Filament Immunofluorescence and Negative Staining Polymerization reactions were carried out in buffer S in the presence of 5 mM MgCl₂ and the indicated nucleotide. Before nucleotide addition, all samples were prespun in an ultracentrifuge (279,000 × g, 20 min at 4°C). For immunofluorescence, aliquots of the reaction (25 μM septin, 100 μM nucleotide) were diluted in buffer S to a final septin concentration of 3.3 μg/ml, and 100 μl drops were spotted on aminopropylsilane-treated coverslips. One minute after sample application, the coverslips were washed with 2 ml buffer S, fixed in -20°C methanol, and stained with a polyclonal rabbit antibody raised against a peptide comprising the 16 amino-terminal residues of XI Sept2 (M.G. and A.A.H., unpublished data). Images were acquired by using a Spot II cooled CCD camera (Diagnostic Instruments) mounted on a Zeiss Axiosplan II microscope and were processed with Metamorph software (Universal Imaging) or Adobe PhotoShop. For electron microscopy, samples (25 μM septin, 500 μM nucleotide) were applied to carbon-coated grids for 5 min. The grids were rinsed with a drop of S buffer, stained with two drops of 1% uranyl acetate, and viewed in a CM120 BioTwin electron microscope (Phillips).

Light Scattering XI Sept2 in S buffer was centrifuged as above and was placed in a quartz cuvette in a LS50B spectrometer (Perkin Elmer). The sample was illuminated with 350-nm light, and the scattered light at the same wavelength was collected at a 90° angle. After a stable baseline was obtained, the cuvette was removed from its position in order to add 0.5 mM nucleotide and MgCl₂ to 5 mM, and its content was rapidly mixed before repositioning the cuvette. The temperature in the cuvette was maintained at 20°C throughout all of the experiments by using a water cooling system. For estimation of the critical concentration, steady-state assembly reactions were subjected to serial dilutions in S buffer containing 0.5 mM nucleotide and were placed in a water bath at 20°C for 2 hr. Scattering measurements of each dilution were performed for 20 or more min to ensure that a stable polymerization level had been reached. All intensity measurements are given in arbitrary units; the light scattered by buffer alone was subtracted from the plotted values. Data points were collected every second and were analyzed with WinLab (Perkin Elmer).

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