Two novel heteropolymer-forming proteins maintain the multicellular shape of the cyanobacterium Anabaena sp. PCC 7120

Benjamin L. Springstein1, Dennis J. Nürnberg2,*, Christian Woehle1,†, Julia Weissenbach1,‡, Marius L. Theune1, Andreas O. Helbig3, Iris Maldener4, Tal Dagan1 and Karina Stucken5

1 Institute of General Microbiology, University of Kiel, Germany
2 Department of Life Sciences, Imperial College, London, UK
3 AG Proteomics & Bioanalytics, Institute for Experimental Medicine, Christian-Albrechts-Universität zu Kiel, Germany
4 Interfaculty Institute of Microbiology and Infection Medicine Tübingen/Organismic Interactions, University of Tübingen, Germany
5 Department of Food Engineering, University of La Serena, Chile

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Correspondence
B. L. Springstein, Department of Microbiology, Blavatnik Institute, Harvard Medical School, 4 Blackfan Cir, Boston, MA, 02115, USA
Tel: +1 857 225 4160
E-mail: benjamin_springstein@hms.harvard.edu

K. Stucken, Department of Food Engineering, University of La Serena, Av. Raúl Bitran N°1305, La Serena, 1720010, Chile
Tel: +56 51 2204397
E-mail: kstucken@userena.cl

Present address
* Institute of Experimental Physics, Free University of Berlin, Berlin, Germany
† Max Planck-Genome-Centre Cologne, Max Planck Institute for Plant Breeding Research, Cologne, Germany
‡ Faculty of Biology, Technion-Israel Institute of Technology, Haifa, Israel

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Polymerizing and filament-forming proteins are instrumental for numerous cellular processes such as cell division and growth. Their function in stabilization and localization of protein complexes and replicons is achieved by a filamentous structure. Known filamentous proteins assemble into homopolymers consisting of single subunits – for example, MreB and FtsZ in bacteria – or heteropolymers that are composed of two subunits, for example, keratin and α/β tubulin in eukaryotes. Here, we describe two novel coiled-coil-rich proteins (CCRP) in the filament-forming cyanobacterium Anabaena sp. PCC 7120 (hereafter Anabaena) that assemble into a heteropolymer and function in the maintenance of the Anabaena multicellular shape (termed trichome). The two CCRPs – Alr4504 and Alr4505 (named ZicK and ZacK) – are strictly interdependent for the assembly of protein filaments in vivo and polymerize nucleotide independently in vitro, similar to known intermediate filament (IF) proteins. A ΔzicKΔzacK double mutant is characterized by a zigzagged cell arrangement and hence a loss of the typical linear Anabaena trichome shape. ZicK and ZacK interact with themselves, with each other, with the elongosome protein MreB, the septal junction protein SepJ and the divisome associate septal protein SepI. Our results suggest that ZicK and ZacK function in cooperation with SepJ and MreB to stabilize the Anabaena trichome and are likely essential for the manifestation of the multicellular shape in Anabaena. Our study reveals the presence of filament-forming IF-like proteins whose function is achieved through the formation of heteropolymers in cyanobacteria.

Abbreviations
CC, coiled-coil; CCRP, coiled-coil-rich proteins; CDF, cumulative distribution function; CSB, conserved syntenic blocks; DAPI, 4’,6-diamidino-2-phenylindole; FRAP, fluorescence recovery after photobleaching; PBP, penicillin-binding protein; PG, peptidoglycan; Van-FL, BODIPY™ FL vancomycin.
Introduction

Cytoskeletal proteins that polymerize to form protein filaments are paramount in bacterial cell biology where they play a role in cell division, alignment of bacterial microcompartments (BMCs), chromosome and plasmid segregation, organization of cell polarity and the determination of cell shape [1]. For example, FtsZ [2], the prokaryotic homolog to the eukaryotic tubulin [3], is a main component of the divisome [4], a multiprotein complex that governs cell division in bacteria and self-assembles into a proteinaceous ring (called Z-ring) at the midcell position [5]. Another key bacterial cytoskeletal protein is MreB [6], which is a homolog of the eukaryotic actin [2, 7] and a crucial component of the multiprotein complexes termed the elongosomes. This complex modulates cell elongation in many rod-shaped bacteria through regulating peptidoglycan (PG) biogenesis [8]. Both, FtsZ and MreB monomers assemble into filamentous strands (protofilaments), consisting of only one type of monomer, termed homopolymers [1]. The cell division in prokaryotes markedly contrasts the division of plastid organelles in photosynthetic eukaryotes that are of cyanobacterial origin [9]. Cell division in plastids is dependent on the cooperative function and heteropolymerization of two FtsZ homologs, FtsZ1 and FtsZ2 in the green lineages and FtsZA and FtsZB in the red lineage. However, each FtsZ homolog is also self-sufficient to form homopolymers (reviewed by [10]). In contrast, a likely horizontally transferred pair of tubulin homologs, BtubA and BtubB from Prothescobacter spp., exclusively assembles into heteropolymers in vitro [11], revealing similar properties than eukaryotic microtubules that are heteropolymers composed of α and β tubulin monomers [12]. Eukaryotic IF proteins, despite sharing substantially the same building blocks and a high degree of coiled-coil (CC) domains [13], which are considered excellent mediators of protein–protein interactions [14], only form heteropolymers with a subset of other IF proteins within their same assembly group but otherwise form obligate homopolymers [15].

Polymer-forming coiled-coil-rich proteins (CCRPs) have been shown to play a role also in multicellularity traits in myxobacteria and actinomycetes (reviewed by [1, 16]). Similar to eukaryotic IFs [13], many bacterial CCRPs perform cytoskeletal functions through their ability to self-assemble into filaments in vitro and in vivo in a self-sufficient and cofactor independent manner [17–20]. The CCRP Crescentin determines the Caulobacter crescentus typical curved morphology by aligning to the inner cell curvature and exuding local mechanical constraints on the PG biosynthesis, likely through cooperation with MreB [17, 21]. In analogy to eukaryotic IF proteins, Crescentin assembles into straight protein filaments with a width of 10 nm and displays a similar domain organization [17]. However, while Crescentin is often considered a prokaryotic homologue to eukaryotic IF proteins, its restricted distribution to only one identified organism questions real homologous relationships and rather suggests that it was acquired by horizontal gene transfer [22]. Multicellular actinomycetes, such as Streptomyces spp., grow by building new cell wall (i.e., PG) only at the cell poles, independent of MreB [23], a striking different cell growth than in most other bacteria [24]. This characteristic polar growth mode is organized by a cytoskeletal network of at least three CCRPs – DivIVA, Scy and FilP – that directly interact with each other to form the polarisome [18]. FilP and Scy independently self-assemble into filaments in vitro [18, 25], thereby fulfilling major IF-like criteria [1]. In vivo, however, Scy does not form filaments and instead accumulates as foci at future branching points [18], while FilP localizes as gradient-like caudates at the hyphal tips [26], instead of forming distinct filaments as observed for Crescentin [17].

Among prokaryotes, cyanobacteria exhibit the largest morphological diversity, comprising unicellular species as well as complex cell-differentiating multicellular species [27]. For the model multicellular cyanobacterium Anabaena, it is imperative to form stable trichomes in order to cope with external influences such as shearing stress [28, 29]. Under nitrogen-deprived growth conditions, Anabaena develops specialized cell types for nitrogen fixation (heterocysts), which are evenly spaced among the Anabaena trichome and provide other vegetative cells with fixed nitrogen compounds like glutamine [30]. In Anabaena, proteinaceous cell-joining structures that allow intercellular transport (i.e., cell–cell communication) and function by gating are termed septal junctions (analogous to eukaryotic gap junctions [31, 32]). Septal junctions consist of several structural elements, an intracellular cap, a plug inside the cytoplasmic membrane formed by the septal junction protein FraD and a tube traversing the septum through nanopores in the peptidoglycan [31]. The correct positioning of the septal protein SepJ, which is involved in septum maturation and filament stability, among others, depends on the FtsZ-driven divisome component FtsQ [33], which links the early and late assembly components of the divisome [34]. FtsZ was shown to be essential for Anabaena viability and to assemble in a typical Z-ring structure at future septum sites in vegetative cells while being downregulated in heterocysts [35–37].
contrast, MreB is dispensable for Anabaena viability but determines its typical cylindrical cell shape, since ΔmreB mutant cells display a pronounced rounded and swollen morphology. This cell swelling, however, does not affect chromosome segregation, which was found to be governed, at least in part, by random segregation in Anabaena [38]. Maintenance of the cylindrical cell shape is furthermore dependent on a class B penicillin-binding protein (PBP) [39] and AmiC-type cell wall amidases in Anabaena [40,41], suggesting that loss of normal cell shape is commonly associated with defects in PG biogenesis.

In this work, we aimed to identify proteins that play a role in cyanobacterial morphology and multicellularity. Searching for IF-like CCRPs, we identified two novel CCRPs in Anabaena that are capable of assembly exclusively into a heteropolymer in vitro and in vivo and that have a putative role in the Anabaena linear trichome shape.

Results

CCRPs ZacK and ZicK from Anabaena are conserved in cyanobacteria

A computational survey of the Anabaena genome for protein-coding genes containing a high coiled-coil content [42] revealed two CCRPs Alr4504 and Alr4505; here, we term the two CCRPs ZicK and ZacK, respectively (i.e., zig and zag in German). ZicK is predicted to contain five distinct coiled-coil (CC) domains, while ZacK has four CC domains (Fig. 1A; File S1). Using PSORTb (v.3.0.2), both ZicK and ZacK are predicted to be cytoplasmic proteins, which is corroborated by the absence of detectable transmembrane domains (predicted using TMHMM v. 2.0). Since both proteins (and their homologs) are annotated as hypothetical proteins (File S2), we validated their transcription under standard (BG11) and diazotrophic (BG11o) growth conditions (Fig. S1B,C). The genomic neighbourhood of zicK and zacK motivated us to test for a common transcriptional regulation of both genes (i.e., an operon structure); however, we did not identify a common transcript (using RT–PCR; Fig. S1A,B). Searching for known proteins sharing structural similarities to ZicK/ZacK using I-TASSER revealed structural similarities between ZicK and the eukaryotic cytolinker protein plectin, and of ZacK with the cell division protein EzrA, a predicted structural similarity that was previously associated with other bacterial CCRPs, including Crescentin, HmpFSyn and HmpFsyn [42]. Further annotation using the NCBI conserved domain search [43] showed that ZicK and ZacK contain ‘structural maintenance of chromosomes’ (SMC) domains (Fig. 1B), similarly to what we previously identified in other self-polymerizing cyanobacterial CCRPs [42]. A search for ZicK/ZacK homologs by sequence similarity revealed that they are absent in pico-cyanobacteria (i.e., Synechococcus/Prochlorococcus) and generally rare in unicellular cyanobacteria. Otherwise, about 50% of the examined 168 cyanobacterial genomes have homologs for the two genes (Fig. 1C and File S2). Several heterocystous cyanobacteria lacking ZicK/ZacK homologs are characterized by a reduced genome (e.g., Nostoc azollae PCC 7008 and Richelia intracellularis). Additionally, Chlorogloeopsis spp. that forms multiseri- ate filaments is lacking the homologs and several strains of Fischerella spp., which form true branching filaments, harbour only a ZacK homolog. The protein sequence of ZicK and ZacK homologs is well conserved, with about 55% amino acids identity among the homologs. The number of CC domains, however, differ among ZicK and ZacK homologs: between 3–6 CC domains in ZicK homologs, and 3–10 domains in ZacK homologs (File S1). Notably, ZicK and ZacK are neighbours in 53 out 72 genomes; both proteins and their genomic neighbour- hood are highly conserved among heterocystous cyanobacteria (Fig. 1C,D).

ZicK and ZacK are interdependent for polymerization in vitro

As a prerequisite for proteins to be considered as IF-like proteins, it is imperative for them to be able to self-interact in vivo and to polymerize into long protein filaments in vitro [1]. To investigate the in vitro polymerization properties of ZicK and ZacK, we employed an in vitro polymerization assay that we previously established to test CCRPs’ polymerization properties [42]. As a positive control for our approach, we used Crescentin (Fig. 1A), which formed an extensive filamentous network in our in vitro assay (Fig. S2). As negative controls, we included empty vector-carrying Escherichia coli cells, GroEL1.2 from Chlorogloeopsis fritschii PCC 6912 (known to form oligomers [44]) and the highly soluble maltose-binding protein (MBP), all of which were tested negative for filament formation in vitro using our approach (Fig. S2). Purified ZicK protein formed into amorphous, nonfilamentous protein aggregates while for ZacK low levels of aggregated sheet-like structures could be observed (Fig. 2A). However, the vast majority of ZacK protein precipitated into clumps of aggregates upon renaturation, which resembled the structures observed for GroEL1.2, suggesting that ZacK has only a partial capacity to self-polymerize or, more likely, is highly unstable on its own in vitro. Inspired by the close
genomic localization of zicK and zacK, we next tested for a potential heteropolymerization of both proteins. This revealed that ZicK and ZacK co-assembled into a meshwork of protein heteropolymers upon corenatura-
tion (Fig. 2A, Fig. S3). While both, ZicK and ZacK renatured alone, formed aggregates in the dialysis tubes that were detectable with the naked eye (similar to our observations from GroEL1.2), corenatured ZicK/ZacK remained in solution, a common property of eukaryotic IFs [45]. Next, we tested whether the cofilamentation was dosage dependent and observed that distinct protein filaments could be detected in vitro only with equal amounts of ZicK and ZacK (Fig. S3). To further test for an in vivo self-interaction, we analysed the self-binding capacity ZicK and ZacK using the bacterial adenylate cyclase two-hybrid (BACTH) assay and found that ZicK and ZacK interact with themselves and also with each other (Fig. 2B) confirming the heterologous binding capacity of both proteins. Consequently, ZicK and ZacK fulfil a major characteristic of IF and IF-like proteins as they are able to self-assemble into filament-like structures in vitro, although, unlike other bacterial IF-like CCRPs, this assembly exclusively occurs as a heteropolymer.

**ZicK and ZacK are interdependent for polymerization in vivo**

To examine the in vivo localization pattern of ZicK and ZacK, we initially expressed translational GFP
fusions of both proteins from the replicative pRL25C plasmid, which is commonly used in experimental work in Anabaena [38,46]. The expression of ZicK-GFP and ZacK-GFP from their respective native promoters (as predicted using BPROM [47]) revealed no discernible expression of ZacK-GFP, while ZicK-GFP accumulated within the cells as dot-like aggregates (Fig. S4A). We assume that the lack of ZacK-GFP expression from its predicted native promoter is based on the uncertainty of the precise promoter site prediction. Alternatively, and although not expressed as an operon, the expression of \( zacK \) could be affected by the expression of \( zicK \) by so far unknown mechanisms. Consequently, we proceeded to investigate the in vivo localization of both proteins from the copper-regulated \( petE \) promoter (\( P_{petE} \)), which is commonly used to study the subcellular protein localization in Anabaena (e.g., FtsZ, MreB and SepI; [36,38,48,49]).

The expression of ZicK-GFP and ZacK-GFP from \( P_{petE} \) in Anabaena independently did not reveal distinct or filamentous structures but resulted in the formation of inclusion body-like aggregates within the cells (Fig. 3A), similar to those observed for \( P_{zicK}::zicK-gfp \) (Fig. S4A). We could not detect any structures when we expressed ZicK and ZacK N-terminally fused to GFP, suggesting that the N terminus is key for proper protein folding. Consequently, we next proceeded to co-express the two proteins with different fluorophores from \( P_{petE} \): ZicK C-terminally fused to eCFP (ZicK-eCFP) and ZacK C-terminally fused to GFP (ZacK-GFP). Heterologous co-expression revealed the formation of a ZicK/ZacK heteropolymer filament-like structure that usually localized along the longitudinal cell axis and in rare occasions also perpendicular to the cell axis (Fig. 3B). The prominent formation of the ZicK/ZacK heteropolymer was furthermore evident as
electron-dense filament-like structures in ultrathin sections using electron microscopy (Fig. 3C). Nonetheless, we could not identify similar filamentous structures in Anabaena WT electron micrographs. Thus, our results identify the general potential of ZicK and ZacK to form a heteropolymer in vivo, yet, the frequency and scale of the ZicK/ZacK heteropolymer in the wild-type remains an open question. To confirm that the localization of fluorophore-tagged ZicK and ZacK is not affected by the wild-type (WT) zicK or zacK alleles, we additionally localized both proteins individually or together in a ΔzicKΔzacK double mutant. These experiments revealed the same localization pattern as in Anabaena WT, suggesting that natively present ZicK or ZacK proteins do not affect the formation of ZicK/GFP, ZacK-GFP or the ZicK-eCFP/ZacK-GFP heteropolymer (Fig. S4B,C). The intracellular localization of the ZicK/ZacK heteropolymer in Anabaena indicates that the polymer is either anchored at the cell poles or specifically split during cell division, as ZicK/ZacK filament-like structures were never observed to cross cell–cell borders and only traversed through not yet fully divided cells (Fig. 3B inlay and Fig. 3C). To further explore whether the ZicK/ZacK heteropolymer assembly is restricted to Anabaena, we proceeded to analyse ZicK and ZacK in vivo in an unrelated heterologous system using the E. coli split GFP assay [50]. Clearly, discernible filamentous-like structures (reminiscent of FilP-GFP [20]) could be detected upon co-expression of ZicK and ZacK C-terminally fused to the split GFP products (ZicK-NGFP and ZacK-CGFP; Fig. S5). This is in agreement with the lack of discernible structures upon expression of N-terminally GFP-fused ZicK and ZacK in Anabaena. Nonetheless, some indications for heteropolymerization were also present upon co-expression of NGFP-ZicK with ZacK-CGFP, which is in agreement with the BACTH results that indicated that both, N and C-terminal fusions of ZicK and ZacK, are potentially able to interact with each other (Fig. 2B). The different

![Fig. 3. ZicK and ZacK form a heteropolymer in vivo. (A, B) Merged GFP or eCFP-fluorescence and chlorophyll autofluorescence (red) and bright-field micrographs of Anabaena WT cells expressing (A) ZicK-GFP, ZacK-GFP or (B) co-expressing ZicK-eCFP and ZacK-GFP from PpetE. (B) Inlay shows that ZicK/ZacK filaments only cross not yet fully divided cells. (A, B) Results show representative figures from three independent experiments. Scale bars: (A, B) 5 µm, (B inlay) 2.5 µm. Notably, overexpression of ZicK-eCFP and ZacK-GFP does not lead to morphological alterations in Anabaena; however, it is likely that the fluorescently-tagged proteins are nonfunctional (as indicated by the lack of mutant complementation; see Fig. S4). Consequently, one should consider the ZicK/ZacK heteropolymer localization as observed here with a grain of salt. We suggest that the observed filaments are indicative of ZicK and ZacK heteropolymerization in the cell. (C) Electron micrographs of ultrathin sections of Anabaena WT and Anabaena cells co-expressing ZicK-eCFP and ZacK-GFP. Black arrows indicate electron-dense structures coinciding with the ZicK/ZacK heteropolymer observed in B. Figure C shows representative result from several independent trichomes investigated in a single experiment. Scale bars: 1.6 µm.](image-url)
heteropolymerization phenotype of ZicK/ZacK polymer in *E. coli* and *Anabaena* suggests that there are other so far unknown factors that modulate the specific ZicK/ZacK heteropolymerization phenotype, as highlighted in the following sections.

**Deletion of zicK and zacK leads to defects in trichome and cell shape and *Anabaena* viability**

In contrast to the obtained ΔzicKΔzacK double mutant, single ΔzicK or ΔzacK mutant strains could not be generated, suggesting that the presence (or absence) of ZicK or ZacK alone is lethal for *Anabaena*. Further investigating the ΔzicKΔzacK mutant phenotype revealed an altered trichome and cell shape phenotype and a reduced trichome viability (Fig. 4). Unlike the linear trichome growth pattern of the WT, the ΔzicKΔzacK mutant strain mostly grew as zigzagged trichomes (Fig. 4A,B), a phenotype that could be rescued by heterologous expression of P<sub>pZicK::zicK-zacK</sub> from pRL25C but not from P<sub>pZicK::zicK-ecfp+zacK-gfp</sub> (Figs S4C and S6A). Additionally, ΔzicKΔzacK cells were significantly larger (WT: 27.42 ± 14.75 µm<sup>3</sup>; ΔzicKΔzacK: 32.52 ± 12.54 µm<sup>3</sup>; *P* < 0.0001; Student’s *t*-test) and significantly more round (defined as the mean of 4*cell area*/π*major_axis<sup>2</sup>); WT: 0.8063 ± 0.1317; ΔzicKΔzacK: 0.8530 ± 0.1130; *P* < 0.0001; Student’s *t*-test) in comparison with the WT (Fig. 4C,D), reminiscent of the ΔmreB mutant strain [38]. The round and swollen cell phenotypes of the ΔzicKΔzacK mutant strains are indicative of an impairment in cell wall integrity and/or defects in PG biogenesis as well as an elevated sensitivity to turgor pressure [51,52]. Consequently, we tested for an elevated sensitivity of the ΔzicKΔzacK mutant to cell wall damaging enzymes. This showed...
that the ΔzicKΔzacK mutant is slightly more sensitive to Proteinase K treatment but was unaffected by lysozyme treatment and still retained the ability to grow diazotrophically (i.e., on BG110 plates) (Fig. 4E) and to form heterocysts (Fig. S6B). More importantly, however, we found that the ΔzicKΔzacK mutant lost the ability to grow in liquid culture (with and without agitation; Fig. 4F), hinting for an elevated sensitivity to fluid shear stress or turgor pressure. This defect could be complemented by heterologous expression of PzicK::

ZicK and ZacK interact with proteins involved in cell shape and trichome integrity

Considering the impact of the deletion of zicK and zacK on cell and trichome shape and the assumed septal docking of the ZicK/ZacK heteropolymer, we next wanted to investigate whether both, ZicK and ZacK, physically interact with other proteins known to function in cell shape determination and cell–cell communication. Using the BACTH assay, we found that both, ZicK and ZacK, interacted with the divisome-associated septal protein SepI [49], the septal protein SepJ [54], the cell shape-determining protein MreB as well as an elongosome associated protein (ZipM; covered in more detail in subsequent study) and the Anabaena homolog to HmpF (here named HmpFAna), whose homologs were shown to be involved in motility in Nostoc punctiforme ATCC 29133 [55] and Synechocystis sp. PCC 6803 [42] (Fig. 5A). No interactions were found with FtsZ, FraC and FraD (Fig. S7). We attempted to further confirm our interaction results with affinity co-elution experiments but found that N-NTA-bound ZicK and ZacK purified from E. coli readily precipitated upon transfer from denaturing to native buffer conditions, precluding further co-elution studies. Additionally, we observed that nondenaturing conditions failed to purify overexpressed CCRPs from E. coli, confirming their inherent insoluble nature, a property known to eukaryotic IFs [56]. Instead, we surveyed for further interaction partners by anti-GFP co-immunoprecipitation experiments of Anabaena cells expressing ZicK-GFP and analysed coprecipitated proteins by LC-MS/MS analytics (all 27 identified possible interactors are listed in File S3). This analysis confirmed that ZicK and ZacK interact with each other in vivo and further strengthened the observed association of ZicK with MreB (Fig. 5B). Furthermore, ZicK coprecipitated ParA (Fig. 5B), a Walker A-type ATPase, involved in chromosome and plasmid partitioning [57].

Deletion of zicK and zacK affects the localization of MreB and the chromosomes

As our BACTH analysis identified SepI and SepJ as interaction partners, and both proteins are involved in intercellular transport and cell–cell communication in Anabaena [49,58], we proceeded to test whether the ΔzicKΔzacK mutant is also affected in solute diffusion using fluorescence recovery after photobleaching (FRAP) experiments of calcine-stained ΔzicKΔzacK mutant. However, this analysis did not reveal any defect in cell–cell communication in the ΔzicKΔzacK mutant (Fig. S8A–C), and hence, ZicK and ZacK do not affect septal junction functionality. Additionally, electron microscopy of ultrathin sections of the ΔzicKΔzacK mutant did not show any discernible differences in the ultrastructure of the cells compared to cells of Anabaena WT (Fig. S8D). In accordance with a lack of interaction between FtsZ and ZicK/ZacK, FtsZ placement was unaffected in the ΔzicKΔzacK mutant as shown using anti-FtsZ immunofluorescence (Fig. S9A). Following the lead of ZicK/ZacK interaction partners, we next analysed the localization of MreB in the Anabaena WT and the ΔzicKΔzacK mutant using a functional PpRL25C::gfp-mreB fusion [38]. In Anabaena WT, we predominantly observed GFP-MreB filaments throughout the cells without any directional preferences (96% or 202 of 211 counted cells) and rarely cell pole-localized foci (4% or 9 of 211 counted cells) (Fig. 6A). Even though GFP-MreB filaments were present in 40% (81 of 199 counted cells) of the ΔzicKΔzacK mutant cells expressing GFP-MreB (Fig. 6A inlay), we only detected those filaments in nonrounded cells that seemingly had a WT-like phenotype (Fig. 6A). In contrast, in rounded/swollen cells of zigzagged trichomes, GFP-MreB was mostly (60% or 118 of 199 counted cells) restricted to the cell poles (Fig. 6A). WT-like nonrounded cells accounted for 24% of the ΔzicKΔzacK mutant cells expressing GFP-MreB (245 of 1040 cells counted), whereas rounded/swollen cells of zigzagged trichomes accounted for 1% of the ΔzicKΔzacK mutant cells (795 of 1040 counted cells; note that not all cells counted for cell shape characteristics also contained GFP-MreB signals, thus explaining the discrepancies between the different counted cell numbers. This is likely due to the copy number variation of the pRL25C plasmid). To further investigate the potential
effect of zicK and zacK deletion on MreB and hence elongosome function, we stained sites of active cell wall biosynthesis using a fluorescent vancomycin derivative (Van-FL; [59]). The staining pattern between the WT and the ΔzicKΔzacK mutant was indistinguishable but the fluorescence intensity levels were slightly decreased in the ΔzicKΔzacK mutant (Fig. S9B,C). Nonetheless, this is likely accounted for by the reduced growth rate of the ΔzicKΔzacK mutant (Fig. 4E and general observation on growth plates).

Considering the interaction of ZicK/ZacK with ParA, we further tested for a function of ZicK and ZacK in DNA placement and compared the DNA distribution in the WT and the ΔzicKΔzacK mutant as measured by distribution of 4’,6-diamidino-2-phenylindol (DAPI) staining intensity (Fig. 6B,C). For that, we calculated the width of the DAPI focal area as the range of DAPI staining around the maximum intensity focus (±10 grey intensity in arbitrary units). This revealed that the staining focal area size was significantly different among the WT and the ΔzicKΔzacK mutant. The DAPI signal observed in the ΔzicKΔzacK mutant appears more condensed, and indeed, the ΔzicKΔzacK mutant focal DAPI area was smaller than the WT (Fig. 6C). Overall, our results may suggest the involvement of ZicK/ZacK in DNA distribution and segregation in dividing cells.

**Discussion**

Here, we provide evidence for the general capacity of two *Anabaena* CCRPs, which we termed ZicK and ZacK, to form polymers *in vitro* and *in vivo*. While the previously described prokaryotic filament-forming CCRPs formed homopolymers [17,19,20,60], ZicK and ZacK exclusively assembled into a heteropolymer *in vitro* and *in vivo*, thus revealing a new property of bacterial CCRPs. The inherent heteropolymerization tendency of ZicK and ZacK was confirmed in a heterologous and evolutionary distant *E. coli* system, which was previously used to investigate other known CCRPs such as Scc from *Leptospira biflexa* [61] or Crescentin [62]. Although heteropolymerization has previously been described for prokaryotic cytoskeletal proteins, none of those polymerization pairs both belonged to the group of CCRPs. BacA and BacB, members of the widely conserved class of bactofilins, both independently polymerize into filaments *in vitro*, colocalize *in vivo* in *C. crescentus* and interact directly with each other as indicated by co-
immunoprecipitation analysis. Unlike CCRPs, whose self-interaction is based on the high degree of CC domains, in bactofilins, the DUF583 domain is proposed to mediate protein polymerization [63].

Fig. 6. Effect of zicK and zacK deletion on MreB localization and DNA distribution. (A) Merged GFP fluorescence and chlorophyll autofluorescence micrographs of Anabaena WT and ΔzicKΔzacK mutant expressing GFP-MreB from PetE. Cells were grown on BG11 plates. Scale bars: 5 µm. Inlay shows clear MreB filaments in cells displaying WT-like cell shapes of the ΔzicKΔzacK mutant. (B) DAPI fluorescence and merged bright-field and chlorophyll autofluorescence micrographs of Anabaena WT and ΔzicKΔzacK mutant on BG11 plates. White arrows indicate strings of DNA that seemingly traverse from one cell to the other. Notably, no such strings are observed in dividing cells (white star), suggesting that it is an effect that occurs after complete cell division. Although we note that high-resolution microscopy would be needed to fully resolve this observation. (A, B) Images presented are representatives from duplicate experiments. Scale bars: 5 µm. (C) Plot profile showing a cumulative distribution function (CDF) of the DAPI signal intensities of pixels (grey value) along the long axis of Anabaena WT and ΔzicKΔzacK mutant cells (n = 151 for each strain) in arbitrary units (a.u.) and arranged to the respective peak maxima. The focal area size in the ΔzicKΔzacK mutant was smaller in comparison with the Anabaena WT (P = 6.8 × 10⁻¹¹⁷, using Wilcoxon test). Notably, the comparison of cell length among the strains reveals a similar result: the ΔzicKΔzacK mutant cell size was smaller in comparison to the Anabaena WT (P = 7.2 × 10⁻¹¹⁷, using Wilcoxon test). Consequently, we compared the area of the focal DAPI staining decided by the cell size among the strains. This, however, revealed that this ratio is not significantly different between the ΔzicKΔzacK mutant and the Anabaena WT.
BacA and BacB has not been studied in vitro. Another interesting pair of potential copolymerizing cytoskeletal proteins that both independently assemble into homopolymers but also co-align in vivo and affect each other’s properties are Crescentin and the CtpS enzyme from C. crescentus [62]. However, again, the co-assembly in vitro is not reported in the literature.

Despite the numerous independently confirmed heteropolymerization properties of ZicK and ZacK, we note that the results from our in vivo experiments are based on artificial and heterologous expression of the two CCRPs and thus in itself could lead to artefacts. We hypothesize that the absence of a ZicK/ZacK heteropolymer in strains expressing ZicK-GFP or ZacK-GFP alone (with the WT zicK and zacK alleles still present) may be due to a dosage-dependent effect, where the presence of unequal concentration of ZicK or ZacK in the cell leads to protein aggregates. Our observation of ZicK-GFP or ZacK-GFP aggregates upon expression alone in the ΔzicKΔzacK mutant strain supports the dosage effect hypothesis. Also, in our in vitro polymerization assay, ZicK and ZacK only formed clear and distinct filament-like structures when both proteins are present in equal concentrations. Nonetheless, co-expression of ZicK-eCFP and ZacK-GFP could not complement the ΔzicKΔzacK mutant. Attempts to express both proteins fused to a fluorophore from the native promoter remained unsuccessful, possibly a result of the close genomic proximity. Furthermore, the genomic neighbourhood of zicK and zacK suggests that the ZicK/ZacK heteropolymer formation could be relying on cotranslational assembly (e.g., as observed for LuxA/LuxB [64]). Cotranslational assembly of natively present ZicK and ZacK would lead to an efficient binding of the two subunits such that the additional expression of one unit only in excess (i.e., ZicK-GFP or ZacK-GFP alone) would lead to the formation of aggregates. As such, it remains to be elucidated to what extent the ZicK/ZacK heteropolymer actually exists in Anabaena WT. Although we could not identify any protein filaments in our ultrathin sections from Anabaena WT, other studies have previously described filamentous strings and even longitudinal cell-spanning polymers in multicellular Anabaena and Nostoc strains [65,66]. Despite compelling evidence for the existence of a cyanobacterial Z-ring structure during cell division [48,67], no Z-ring ultrastructures have yet been identified, and consequently, the absence of longitudinal ZicK/ZacK filaments in ultrathin sections does not rule out that they exist but could also indicate that they could not be visualized yet.

Our results indicate that ZicK and ZacK are associated with the elongasome (through their interaction with MreB) and proteins in the septal cell wall (through the interaction with SepJ and SepI) and might affect cellular DNA placement (Fig. 7). A function of ZicK/ZacK in chromosome segregation would be in concert with the identified interaction of ZicK with ParA; this, however, remains to be elucidated as it could also be an indirect consequence of the swollen/rounded cell shape in the ΔzicKΔzacK mutant, similar to the indirect effect of mreB deletion of carboxysome alignment in Synechococcus elongatus [68]. Nonetheless, so far, no chromosome partitioning system has yet been identified in multicellular cyanobacteria. In Anabaena, deletion of mreB did not affect chromosome segregation but induced a swollen cell phenotype [38], similar to the ΔzicKΔzacK mutant. Consequently, MreB and ZicK/ZacK likely share functional properties but are not exclusively involved in the same cellular processes. Swollen cell morphotypes were also described for Anabaena or Synechocystis mutants lacking penicillin-binding proteins (PBP), which are enzymes that are directly involved in cell wall biogenesis through the modification of the PG layer [39]. This presumed link of ZicK/ZacK to the actin-like MreB cytoskeleton and the PG biogenesis apparatus is also indicated by the altered localization of GFP-MreB and the decreased PG staining intensity in the ΔzicKΔzacK mutant strain. Consequently, ZicK and ZacK might indirectly act to positively regulate PG biogenesis, although we cannot exclude that the reduced staining intensity in the ΔzicKΔzacK mutant simply reflects the slower growth rate of this strain. An interaction or involvement of prokaryotic filament-forming CCRPs with MreB and

Fig. 7. Interaction network of known divisome, elongasome and septal proteins in Anabaena. A model for a partial divisome, elongasome and septal junction network in Anabaena as deduced from BACTH and co-IP analyses. Black arrows indicate interactions that have been previously described by [33,49,67]. Red arrows indicate interactions identified in the current analysis.
PG synthesis was previously observed in other bacteria. Examples are the gliding motility in *Mycobacterium xanthus*, where a multiprotein complex, including the filament-forming CCRP AglZ and MreB, was found to coordinate type A-motility [69]. Similarly, the curved morphology of *C. crescentus* is induced by Crescentin, which functionally associates with MreB and likely modulates PG biogenesis by exuding local mechanical forces to the cell membrane [16,21]. Other aspects like a decreased cell envelope permeability of the ΔzicKΔzacK mutant are also conceivable, although we did not detect any cell wall defects in the ΔzicKΔzacK mutant. MreB and the elongosome are the main determinants of the PG exoskeleton, which provides the cell with structural integrity and resistance to turgor pressure [70]. The lack of liquid growth of the ΔzicKΔzacK mutant would also argue for a defect in the resistance to turgor pressure.

Together with the cell shape-determining protein MreB, ZicK and ZacK could possibly contribute to normal cell shape and relay trichome shape-stabilizing properties to neighbouring cells in the trichome by means of their association with the filament stabilizing protein SepJ (Fig. 7). As such, they are important for maintaining the linear *Anabaena* trichome phenotype. ZicK and ZacK polymers might constitute stabilizing platforms or scaffolds for other proteinaceous structures, similarly to the stabilizing function of the eukaryotic cytoskeleton for cell–cell contacts (i.e., desmosomes). Furthermore, ZicK shares in silico predicted structural similarities with the spectrin repeats of plectin, a well-described eukaryotic cytolinker protein. Plectin links the three eukaryotic cytoskeletal systems (actin filaments, microtubules and IFs), thereby contributing to the resistance to deformation of vertebrate cells [12]. They stabilize desmosomes and are hence directly involved in cell–cell contact integrity [71]. An analogous cytolinker function of ZicK could explain why ZacK alone did not form properly folded protein filaments on its own and suggests that ZacK requires ZicK as the linking protein for polymerization. As plectin not only stabilizes but also dynamically disassembles IF protein filaments (i.e., vimentin) in a concentration-dependent manner [72], this would further support a dosage-dependent effect of ZicK and ZacK for heteropolymerization.

The ubiquity of ZicK and ZacK in heterocystous cyanobacteria that form linear trichomes (Fig. 1C, File S2) indicates that the presence of ZicK and ZacK contributes to the maintenance of a linear trichome shape. The ΔzicKΔzacK mutant had a zigzagged phenotype and was unable to grow in liquid culture. We hypothesize that the loss of linear trichome shape led to an increase in accessible surface for the acting mechanical forces in liquid [73], including fluid shear stress [74], ultimately resulting in forces that cannot be endured by the abnormal mutant trichomes. Nonetheless, the absence of ZicK and/or ZacK in several heterocystous cyanobacteria species, including the fern-symbiont *N. azollae* and ramified cyanobacteria, may suggest that these species are less sensitive to mechanical stress (i.e., due to their interaction with the host tissues or complex filament formation), and hints for the presence of additional, yet unknown, mechanisms for the determination of filament shape.

The key hallmarks of permanent bacterial multicellularity are morphological differentiation and a well-defined and reproducible shape, termed patterned multicellularity [75]. In sporulating actinomycetes, this is exemplified by the highly reproducible cell division, cell proliferation and cell differentiation pattern [76]. In filamentous cyanobacteria, the reproducible linear trichomes are considered a major contributor to the cyanobacterial patterned multicellularity [75], manifesting a selective advantage to biotic and abiotic environmental factors [77,78]. Our results indicate that ZicK and ZacK serve as regulators of the typical linear *Anabaena* trichome and as such as regulators of *Anabaena* patterned multicellularity. The evolution of patterned multicellularity is considered an important step towards a sustainable division of labour and the development of cell differentiation [75]. Our study provides initial evidence for a role of two heteropolymer-forming CCRPs in the evolution and maintenance of cyanobacterial multicellular forms.

**Materials and methods**

**Bacterial strains and growth conditions**

*Anabaena* sp. PCC 7120 was obtained from the Pasteur Culture Collection (PCC) of cyanobacteria (France). Cells were grown photoautotrophically in BG11 or without combined nitrogen (BG110) at constant light with a light intensity of 30 µmol·m$^{-2}$·s$^{-1}$ in liquid culture or on agar plates (1.5% w/v agar). When appropriate, 5 µg·mL$^{-1}$ spectinomycin (Sp), 5 µg·mL$^{-1}$ streptomycin (Sm) or 30 µg·mL$^{-1}$ neomycin (Nm) was added to strains carrying respective plasmids or chromosomal insertions. In some cases, basal copper-regulated petE-driven expression of gene candidates in *Anabaena* cells was lethal or growth inhibiting, therefore these strains were grown in BG11 without copper and protein expression was later induced by the addition of CuSO$_4$ at indicated concentrations to the culture. *E. coli* strains DH5α, DH5αMCR, XL1-blue and HB101 were used for cloning and conjugation by triparental mating. BTH101
was used for BACTH system, and BL21 (DE3) was used for expression of His6-tagged proteins in E. coli. All E. coli strains were grown in LB medium containing the appropriate antibiotics at standard concentrations. Tables S1–S4 list all used bacterial strains, plasmids and oligonucleotides.

**Prediction of coiled-coil-rich proteins**

Genome sequence of *Anabaena* (GCA_000009705.1) was analysed by the COILS algorithm [79] as previously described [20]. The algorithm was run with a window width of 21, and the cut-off for amino acids in coiled-coil conformation was set to ≥ 80 amino acid residues. The resulting set of protein candidates was further manually examined with online available bioinformatic tools, including NCBI conserved domain search [80], NCBI BLAST [81], TMHMM [82] and I-TASSER [83]. Protein candidates exhibiting BLAST hits described [20]. The algorithm was run with a window width of 21, and the cut-off for amino acids in coiled-coil conformation was set to ≥ 80 amino acid residues. The resulting set of protein candidates was further manually examined with online available bioinformatic tools, including NCBI conserved domain search [80], NCBI BLAST [81], TMHMM [82] and I-TASSER [83]. Protein candidates exhibiting BLAST hits described [20]. The algorithm was run with a window width of 21, and the cut-off for amino acids in coiled-coil conformation was set to ≥ 80 amino acid residues. The resulting set of protein candidates was further manually examined with online available bioinformatic tools, including NCBI conserved domain search [80], NCBI BLAST [81], TMHMM [82] and I-TASSER [83]. Protein candidates exhibiting BLAST hits involved in cytoskeletal processes or similar domain architectures as known IF and IF-like proteins like Crescentin, FilP, vimentin, desmin or keratin were selected, and enzymatic proteins as well as proteins predicted to be involved in other cellular processes were excluded.

**Distribution of homologs in cyanobacteria**

Homologs to the *Anabaena* proteins were extracted from precalculated cyanobacterial protein families [42]. Conserved syntenic blocks (i.e., gene order) were identified using CSBFinder-S [84].

**RNA isolation and cDNA synthesis**

RNA from *Anabaena* WT was isolated using the Directzol™ RNA MiniPrep Kit (Zymo Research, Freiburg, Germany) according to the manufacturer’s instructions. RNA was isolated in technical triplicates from 10 mL cultures. Isolated RNA was treated with DNA-free™ Kit (2 units rDNAs/reaction; Thermo Fischer Scientific, Dreieich, Germany), and 200 ng RNA was reverse transcribed using the qScript™ cDNA Synthesis Kit (Quanta Biosciences, Beverly, MA, USA). RT–PCR of cDNA samples for *rnpB, zicK, zacK* and *zicK-zacK* was performed using primer pairs #1/#2, #3/#4, #5/#6 and #3/#8, respectively.

**Transformation**

*Anabaena* was transformed by triparental mating as previously described [85]. Briefly, 100 µL of overnight cultures of DH5α carrying the conjugal plasmid pRL443 and DH5αMC carrying the cargo plasmid and the helper plasmid pRL623, encoding for three methylases, were mixed with 200 µL *Anabaena* culture (for transformation into the ΔzicKΔzacK mutant, cells were scraped from the plate and resuspended in 200 µL BG11). This mixture was directly applied onto sterilized nitrocellulose membranes (Amer sham Protran 0.45 NC, Cytiva Europe, Freiburg, Germany) placed on top of BG11 plates supplemented with 5% (v/v) LB medium. Cells were incubated in the dark at 30 °C for 6–8 h with subsequent transfer of the membranes to BG11 plates, and plates were placed under standard growth conditions. After 24 h, membranes were transferred to BG11 plates supplemented with appropriate antibiotics.

**Plasmid construction**

Ectopic expression of *Anabaena* protein candidates was achieved from a self-replicating plasmid (pRL25C [86]) under the control of the copper-inducible petE promoter (PpetE) or the native promoter (predicted by BPROM [47]) of the respective gene. All constructs were verified by Sanger sequencing (Eurofins Genomics, Ebersberg, Germany). Plasmids were created using standard restriction enzyme-based techniques or Gibson assembly. Information about precise plasmid construction strategies is available from the authors upon request.

**Anabaena mutant strain construction**

The ΔzicKΔzacK mutant strain was generated using the pRL278-based double homologous recombination system employing the conditionally lethal sacB gene [87]. For this, 1500 bp upstream and downstream of *zicK-zacK* was generated by PCR from *Anabaena* gDNA. Upstream region of *zicK* was amplified using primers #97/#98, and downstream region of *zacK* was amplified using primers #99/#100. The respective upstream and downstream homology regions flanking the CS.3 cassette (amplified with primer #95/#96 from pCSEL24) were then inserted into PCR-amplified pRL278 (using primer #93/#94) by Gibson assembly, yielding pTHS166. *Anabaena* transformed with pTHS166 plasmids was subjected to several rounds of restreaking on new plates (about 5–8 rounds). To test for fully segregated clones, colony PCRs were performed. For this, *Anabaena* cells were resuspended in 10 µL sterile H2O of which 1 µL was used for standard PCR with internal *zicK* and *zacK* gene primers #3/#6. Correct placement of the CS.3 cassette was then further confirmed using CS.3 cassette primers with binding sites outside of the 5’ and 3’ flanks used for homologous recombination (primers #95/#102 and #101/#96).

**Fluorescence microscopy**

Bacterial strains grown in liquid culture were either directly applied to a microscope slide or previously immobilized on a 2% (w/v) low-melting agarose in PBS (10 mM Na3HPO4, 140 mM NaCl, 2.7 mM KCl, 1.8 mM KH2PO4, pH 7.4) agarose pad and air-dried before microscopic analysis.
Epifluorescence was done using an Axio Imager.M2 light microscope (Carl Zeiss, Oberkochen, Germany) equipped with Plan-Apochromat 63x/1.40 Oil M27 objective and the AxioCam MR R3 imaging device (Carl Zeiss). GFP, Alexa Fluor 488 and BODIPY™ FL Vancomycin (Van-FL) fluorescence was visualized using filter set 38 [Carl Zeiss; excitation: 470/40 nm band-pass (BP) filter; emission: 525/50 nm BP]. Chlorophyll autofluorescence was recorded using filter set 15 (Carl Zeiss; excitation: 546/12 nm BP; emission: 590 nm long pass). When applicable, cells were previously incubated in the dark at RT for about 5 min with 10 µg·mL⁻¹ DAPI (final concentration) to stain intracellular DNA. For visualization of DAPI fluorescence filter set 49 (Carl Zeiss; excitation: G 365 nm; emission: 455/50 nm) was employed. For confocal laser scanning microscopy, the LSM 880 Axio Imager 2 equipped with a C-Apochromat 63x/1.2 W Korr M27 objective and an Airyscan detector (Carl Zeiss) was used and visualization of GFP, eCFP and chlorophyll autofluorescence was done using Zen black smart setup settings.

Transmission electron microscopy

For ultrastructure analysis, Anabaena trichomes were fixed with 2.5% (v/v) glutaraldehyde, immobilized in 2% (w/v) agarose, treated with 2% (v/v) potassium permanganate and dehydrated through a graded ethanol series [88]. The fixed cells were infiltrated by ethanol: EPON (2 : 1 to 1 : 2 ratio) and embedded in pure EPON. Ultrathin sections were prepared with a Leica UC6i Ultramicrotome, transferred to formvar-coated copper grids (Science Services GmbH München) and poststained with uranyl acetate and lead citrate [89]. Micrographs were recorded at a Philips Tecnai10 electron microscope (FEI, Hillsboro, OR, USA) at 80 kV.

Calcein labelling and fluorescence recovery after photobleaching experiments

For FRAP experiments, Anabaena WT and ΔzicKΔzacK mutant strain were grown on BG11 plates, resuspended in BG11 liquid media and washed three times in 1 mL BG11 (3000 g, 5 min). Cells were then resuspended in 0.5 mL BG11 and incubated with 10 µL calcein-AM (Cayman Chemical, Ann Arbor, MI, USA; 1 mg·mL⁻¹ in DMSO) for 1 h at 30 °C in the dark. To remove excess staining solution, the cells were washed four times with 1 mL BG11. Subsequently, the cells were spotted on BG11 agar for visualization by confocal laser scanning microscopy (Leica TCS SP5; HCX PL APO 63x 1.40-0.60 OIL CS). Calcein was excited at 488 nm and fluorescence emission monitored in the range from 500 to 530 nm with a maximally opened pinhole (600 µm). FRAP experiments were carried out by an automated routine as previously described [58]. After recording an initial image, selected cells were bleached by increasing the laser intensity by a factor of 5 for two subsequent scans and the fluorescence recovery followed in 0.5-s intervals for 30 s was recorded using the Leica LAS X software. Exchange coefficients (E) were then calculated as previously described [58,90].

BODIPY™ FL vancomycin (Van-FL) staining

Van-FL staining of strains grown on BG11 plates was essentially performed as described previously [91,92]. Briefly, cells were resuspended in BG11 medium, washed once in BG11 by centrifugation (6500 g, 4 min, RT) and incubated with 5 µg·mL⁻¹ Van-FL (dissolved in methanol; Thermo Fischer Scientific). Cells were incubated in the dark for 1 h at 30 °C, washed three times with BG11 and immobilized on an agarose pad. Van-FL fluorescence signals were then visualized using epifluorescence microscopy with an excitation time of 1 ms. Arithmetic mean fluorescence intensities were recorded from the septa between two cells with a measured area of 3.52 µm² using the histogram option of the zen blue 2.3 software (Carl Zeiss).

Alcian blue staining

Anabaena WT and ΔzicKΔzacK cells were grown on BG11₀ plates, resuspended in BG11₀ liquid medium and stained with 0.05% (w/v) alcian blue (final concentration). Polysaccharide staining of cells immobilized on an agarose pad was observed with an AxioCam E Rc5s colour camera (Carl Zeiss).

Data analysis

Cell volume and roundness were determined using the imaging software imagej [93], and a perfect circle is defined to have a roundness of 1. Cell volume was calculated based on the assumption of an elliptic cell shape of Anabaena cells using the Major Axis and Minor Axis values given by imagej and the formula for the volume of an ellipsoid

\[
V = \frac{4}{3} \pi abc = \frac{4}{3} \pi \left( \frac{\text{Major Axis}}{2} \right)^2 \frac{\text{Minor Axis}}{2}.
\]

Angles between neighbouring cells were determined from chlorophyll autofluorescence micrographs from individual Anabaena trichomes in imagej. For this, the trichomes were separated from the background using the ‘set autothreshold’ command. The adjustable Watershed plug-in was used to separate individual cells in one trichome (tolerance = 1). After selecting cells with the analyse particle command (size = 200–2000 pixel; circularity = 0.20–1.00), the centre of each cell was determined as XY coordinates using the centroid measurement setting. The angles between three of
these points, representing the angle change in the trichome at each cell, were then calculated using basic algebra in Microsoft excel.

Distribution of DAPI fluorescence signal intensity was analysed in ImageJ with the Plot Profile option along the long axis of 151 single cells with the rectangle tool. The resulting grey values were arranged according to the maximum intensity focus, and the width of the DAPI focal area resulting grey values were analysed in ImageJ with the Plot Profile option along the maximum (±10 grey value in arbitrary units). Statistical tests were performed with MATLAB (MathWorks; Natick, MA, USA) or GraphPad Prism v.8 (GraphPad Software, San Diego, CA, USA).

Bacterial two-hybrid and beta-galactosidase assays

Chemically competent E. coli BTH101 cells were cotransformed with 5 μg of plasmids carrying the respective T18 and T25 translational fusion constructs, plated onto LB plates supplemented with 200 μg·mL⁻¹ X-gal, 0.5 mm IPTG, Amp, Km and grown at 30 °C for 24–36 h. Interactions were quantified by beta-galactosidase assays from three independent colonies. For this aim, cultures were grown for 2 days at 20 °C in LB Amp, Km, 0.5 mm IPTG and beta-galactosidase activity was recorded as described in the manufacturer’s instructions (Euromedex, Souffelweyersheim, France; BACTH System Kit Bacterial Adenylate Cyclase Two-Hybrid System Kit) in a 96-well plate format as previously described [94].

GFP-fragment reassembly assay

Chemically competent E. coli BL21 (DE3) were cotransformed with indicated plasmid combinations, plated on LB Amp, Km and grown over night at 37 °C. Liquid overnight cultures of single colonies of the respective plasmid-bearing E. coli strains were then diluted 1 : 40 in the same medium the following day. Cells were grown for 2 h at 37 °C, briefly acclimated to 20 °C for 10 min, and protein expression was induced with 0.05 mm IPTG and 0.2% (w/v) L-arabinose. Pictures of induced cultures grown at 20 °C were taken 48 h after induction.

Co-immunoprecipitation

About 20–30 mL of the respective Anabaena cultures was pelleted by centrifugation (4800 g, 10 min, RT), cells were washed twice by centrifugation (4800 g, 10 min, RT) with 40 mL PBS and then resuspended in 1 mL lysis buffer [PBS-N: PBS with 1% (v/v) NP-40] supplemented with protease inhibitor cocktail (PIC; cOmplete™, EDTA-free Protease Inhibitor Cocktail; Sigma-Aldrich, Steinheim, Germany). Cells were lysed using the VK05 lysis kit (Bertin Instruments, Frankfurt am Main, Germany) in a Precellys® 24 homogenizer (3 strokes for 30 s at 6500 r.p.m.), and cell debris was pelleted by centrifugation (30 min, 21 100 g, 4 °C). 50 μL μMACS anti-GFP MicroBeads (Miltenyi Biotec, Bergisch Gladbach, Germany) were added to the resulting cell-free supernatant and incubated for 1 h at 4 °C with mild rotation. Subsequently, the sample was loaded onto μColumns (Miltenyi Biotec), washed two times with 1 mL lysis buffer and eluted in 50 μL Elution Buffer [50 mm Tris/HCl pH 6.8, 50 mm DTT, 1% (w/v) SDS, 1 mm EDTA, 0.005% (w/v) bromophenol blue, 10% (v/v) glycerol; Miltenyl Biotec]. Samples were stored at −80 °C until further use.

Mass spectrometry

Mass spectrometry of coprecipitated proteins was performed as previously described [49].

Immunofluorescence

Immunolocalization of FtsZ in Anabaena WT and ΔzieKΔzacK mutant was essentially performed as previously described [33]. For this, strains were scraped off from growth plates (BG11 and BG11p plates), resuspended in a small volume of distilled water and air-dried on Polysine® adhesion slides (Menzel) at RT followed by fixation and permeabilization with 70% ethanol for 30 min at −20 °C. Cells were allowed to air dry for 30 min at RT and then washed two times with PBST (PBS supplemented with 0.1% (v/v) Tween-20) for 2 min. Unspecific binding sites were blocked for 30 min at RT with blocking buffer (1× Roti®-ImmunoBlock in PBST; Carl Roth, Karlsruhe, Germany), and afterwards, rabbit anti-FtsZ (Agrisera; raised against Anabaena FtsZ; 1 : 150 diluted) antibody in blocking buffer was added to the cells and incubated for 1.5 h at RT in a self-made humidity chamber followed by five washing steps with PBST. 7.5 μg·mL⁻¹ (final concentration) Alexa Fluor 488-conjugated goat anti-rabbit IgG (H+L) secondary antibody (Thermo Fischer Scientific) in blocking buffer was added to the cells and incubated for 1 h at RT in the dark in a self-made humidity chamber. Subsequently, cells were washed five times with PBST, air-dried and mounted with ProLong™ Diamond Antifade Mountant (Thermo Fischer Scientific) overnight at 4 °C. Immunolocalization of FtsZ was then analysed by epifluorescence microscopy.

Spot assays

For spot assays, Anabaena WT and ΔzieKΔzacK mutant strain was grown on BG11 growth plates, resuspended in BG11 liquid medium and adjusted to an OD₅₇₀ of 0.4. Cells were then spotted in triplicates of 5 μL onto the respective
growth plates containing either no additives (BG11 or BG110), 50 μg·mL⁻¹ proteinase K or 100 μg·mL⁻¹ lysozyme in serial 1/10 dilutions and incubated under standard growth conditions until no further colonies arose in the highest dilution.

**Protein purification and in vitro filamentation assays**

For protein purification, *E. coli* BL21 (DE3) cells carrying His-tagged protein candidates were grown in overnight cultures at 37 °C and 250 r.p.m. The next day, overnight cultures were diluted 1 : 40 in the same medium and grown at 37 °C until they reached an OD₆₀₀ of 0.5–0.6. Protein expression was induced with 0.5 mM IPTG for 3–4 h at 37 °C and 250 r.p.m. Afterwards, cell suspensions of 50 mL aliquots were harvested by centrifugation, washed once in PBS and stored at −80 °C until further use. For *in vitro* filamentation assays, cell pellets were resuspended in urea lysis buffer (ULB: 50 mM NaH₂PO₄, 300 mM NaCl, 25 mM imidazole, 6 mM urea; pH 8.0) and lysed in a Precellys® 24 homogenizer (3× 6500 r.p.m. for 30 s) using the 2 mL microorganism lysis kit (VK01; Bertin) or self-packed Precellys tubes with 0.1-mm glass beads. The resulting cell debris was pelleted by centrifugation at 21 000 g (10 min, 4 °C), and the supernatant was incubated with 1 mL His-Pur™ Ni-NTA resin (Thermo Fischer Scientific) for 1 h at 4 °C in an overhead rotator. The resin was washed five times with 4× resin-bed volumes ULB and eluted in urea elution buffer (UEB: ULB supplemented with 225 mM imidazole). Total protein concentration was measured using the Qubit® 3.0 Fluorometer (Thermo Fischer Scientific). Filament formation of purified proteins was induced by overnight dialysis against polymerization buffer (PLB: 50 mM PIPES, 100 mM KCl, pH 7.0; HLB: 25 mM HEPES, 150 mM NaCl, pH 7.4; or 25 mM HEPES pH 7.5) at 20 °C and 180 r.p.m. with three bath changes using a Slide-A-Lyzer™ MINI Dialysis Device (10K MWCO, 0.5 mL or 2 mL; Thermo Fischer Scientific). Purified proteins were stained with an excess of NHS-fluorescein (dissolved in DMSO; Thermo Fischer Scientific), and *in vitro* filamentation was analysed by epifluorescence microscopy. The resulting protein samples were mass spectrometry analysed by ultrastructure analyses. AOH analysed protein samples by mass spectrometry. BLS, TD and KS performed the experimental work with contributions from MLT and JW. CW and TD performed comparative genomics analysis. DJN performed FRAP assays, and IM carried out ultrastructure analyses. AOH analysed protein samples by mass spectrometry. BLS, TD and KS drafted the manuscript with contributions from all co-authors.

**Conflict of interest**

The authors declare no conflict of interest.

**Author contributions**

BLS and KS designed the study. BLS established and performed the experimental work with contributions from MLT and JW. CW and TD performed comparative genomics analysis. DJN performed FRAP assays, and IM carried out ultrastructure analyses. AOH analysed protein samples by mass spectrometry. BLS, TD and KS drafted the manuscript with contributions from all co-authors.

**Peer Review**

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**Data accessibility**

All data generated or analysed during this study are included in this published article (and its Supporting Information).

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**Supporting information**

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Fig. S1. *zieK* and *zacK* are expressed at standard growth conditions.

Fig. S2. *In vitro* polymerization assay controls.

Fig. S3. Co-polymerization of ZicK and ZacK is dosage-dependent.

Fig. S4. Ectopic expression of ZicK and ZacK.

Fig. S5. Host-independent heteropolymerization of ZicK and ZacK upon heterologous expression in *E. coli*.

Fig. S6. Complementation of the zigzagged growth phenotype of the ΔzieKΔzacK mutant and its ability to grow under diazotrophic conditions.

Fig. S7. ZicK and ZacK do not interact with other components of the septal junctions besides SepJ.

Fig. S8. The ΔzieKΔzacK mutant is not defective in intercellular transport and cellular ultrastructures.

Fig. S9. PG biogenesis and Z-ring placement are largely unaffected in the ΔzieKΔzacK mutant.

Table S1. Cyanobacterial strains.

Table S2. *E. coli* strains.

Table S3. Plasmids.

Table S4. Oligonucleotides.