

GROWTH POLE RING protein forms a 200-nm-diameter ring structure essential for polar growth and rod shape in *Agrobacterium tumefaciens*

J. R. Zupan^a, R. Grangeon^a, J. S. Robalino-Espinosa^{a,b}, N. Garnica^a, and P. Zambryski^{a,1}

^aDepartment of Plant and Microbial Biology, University of California, Berkeley, CA 94720; and ^bDepartment of Molecular Cell Biology, Vrije Universiteit Amsterdam, 1081 HV Amsterdam, The Netherlands

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Polar growth in *Agrobacterium* pirates and repurposes well-known bacterial cell cycle proteins, such as FtsZ, FtsA, PopZ, and PodJ. Here we identify a heretofore unknown protein that we name GROWTH POLE RING (GPR) due to its striking localization as a hexameric ring at the growth pole during polar growth. GPR also localizes at the midcell late in the cell cycle just before division, where it is then poised to be precisely localized at new growth poles in sibling cells. GPR is 2,115 aa long, with two N-terminal transmembrane domains placing the bulk of the protein in the cytoplasm, N- and C-terminal proline-rich disordered regions, and a large 1,700-aa central region of continuous α -helical domains. This latter region contains 12 predicted adjacent or overlapping apolipoprotein domains that may function to sequester lipids during polar growth. Stable genetic deletion or riboswitch-controlled depletion results in spherical cells that grow poorly; thus, GPR is essential for wild-type growth and morphology. As GPR has no predicted enzymatic domains and it forms a distinct 200-nm-diameter ring, we propose that GPR is a structural component of an organizing center for peptidoglycan and membrane syntheses critical for cell envelope formation during polar growth. GPR homologs are found in numerous Rhizobiales; thus, our results and proposed model are fundamental to understanding polar growth strategy in a variety of bacterial species.

Agrobacterium | bacterial polar growth | apolipoprotein | rod-shape morphology | GROWTH POLE RING protein

How bacteria grow and divide is intensely investigated, given the importance of bacteria across all of biology, from microbes to man. Several model systems, including *Escherichia coli*, *Bacillus subtilis*, and *Caulobacter crescentus*, have provided a wealth of fundamental insights into lateral dispersed cell elongation (1). However, as more bacteria enter the ring of scrutiny, another mode of cell elongation, limited to one or both poles, has emerged (2–5). In gram-negative bacteria, a single growth pole is used by the Rhizobiales order of Alphaproteobacteria, which includes the plant pathogen/genetic engineer *Agrobacterium* and plant nitrogen-fixing symbiont *Rhizobium*, as well as the animal pathogens *Brucella* and *Bartonella* (2–14).

Little is known about the enzymes involved, their spatiotemporal expression, or what mechanisms regulate bacterial polar growth. To date, several molecular players for polar growth have been identified in *Agrobacterium tumefaciens*. First, *Agrobacterium*-specific homologs of the classic cell division factors FtsA and FtsZ localize to the growth pole during cell elongation and then migrate to the midcell during initiation and completion of septation (6, 9, 12, 13). At the division site, FtsA and FtsZ are then precisely located at the new growth poles in resulting sibling cells. Second, homologs of *Caulobacter* old and new pole-specific markers, PopZ_{Cc} (15, 16) and PodJ_{Cc} (17, 18), respectively, have opposite localizations in *Agrobacterium*. PopZ_{At} exclusively marks the growing pole (13). PodJ_{At} initially marks the old pole but then also accumulates at the growth pole later in the cell cycle; this relocation is correlated with the transition of the growth pole to an old pole, and PodJ_{At} is proposed to facilitate this transition (13). It

is likely that FtsZ also plays a role in the transition to an old pole, as cells depleted of FtsZ fail to divide but continue to grow (12). Genetic deletion or depletion of PodJ_{At} or PopZ_{At} causes severe growth defects (14, 19–21), attesting to their essential roles.

Notably, the amino acid sequences of PopZ_{At} and PodJ_{At} are distinct from their *Caulobacter* counterparts, each with only 23% identity (13). Nevertheless, as the search for *Caulobacter* homologs successfully identified critical pole-specific factors in *Agrobacterium*, we used this approach again to search for potential homologs of *Caulobacter* TipN, dubbed a landmark protein for establishing and perpetuating new pole polarity throughout most of its cell cycle (22, 23). We found an uncharacterized *Agrobacterium* protein with little amino acid identity to TipN but an overall similarity in secondary structural domains. Most remarkably, the *Agrobacterium* protein localizes into six subpolar foci arranged in a hexameric ring at the growth pole.

The lack of significant amino acid sequence similarity to TipN and its distinctive localization pattern led us to name the newly identified *Agrobacterium* protein GROWTH POLE RING (GPR). Here we describe the domain architecture of GPR, its localization during the *Agrobacterium* cell cycle, and genetic analyses revealing that GPR is essential for polar growth and rod shape morphology.

Results

Identification of GPR Protein. As polar growth in *Agrobacterium* is not well understood, it is critical to identify factors that localize

Significance

The extensively studied rod-shaped bacteria *Escherichia coli*, *Bacillus subtilis*, and *Caulobacter crescentus* grow by dispersed insertion of new cell wall material along the entire length of the cell. An alternative prokaryotic growth mode—polar growth—is used by some Actinomycetales and Proteobacteria. The latter phylum includes the family Rhizobiaceae, of which *Agrobacterium tumefaciens* is a prominent member. The factors responsible for polar growth are largely unknown. Here we describe GROWTH POLE RING (GPR) protein from *Agrobacterium* with a striking localization as a ring of six foci at the growth pole. GPR is potentially the polar growth organizer in *Agrobacterium*, as its absence leads to severe morphological defects, including loss of cylindrical cell shape, while its overproduction leads to ectopic growth poles.

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¹To whom correspondence should be addressed. Email: zambrysk@berkeley.edu.

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to the growth pole that may function to facilitate such growth. We used the *C. crescentus* new pole-localizing protein, TipN, as a query and identified an uncharacterized *Agrobacterium* protein, A9CJ72 (24), Atu1348 in the *A. tumefaciens* proteome, with low (~20%) amino acid sequence identity to TipN. For reasons described below, we named *Agrobacterium* A9CJ72 the GPR protein.

Fig. 1 diagrams the predicted structural domains of GPR (25); *SI Appendix, Fig. S1A* provides more detailed predictions of secondary structure according to PHYRE2 (26). Two predicted N-terminal transmembrane domains suggest that the bulk of the protein resides in the cytoplasm (Fig. 1 and *SI Appendix, Fig. S1A*). The N (80 aa) and C (110 aa) termini are enriched in proline residues and are predicted to be disordered. Strikingly, 82% of GPR, encompassing 1,720 residues (160–1,890), is predicted to be comprised of contiguous α -helices (*SI Appendix, Fig. S1A*). This large central α -helical region (blue area in Fig. 1) contains numerous nonoverlapping and overlapping subdomains homologous to eukaryotic apolipoproteins (ApoLPs; Pfam P01442) (25) with solved structures (27–30). *SI Appendix, Table S1* lists the locations of 12 predicted ApoLP domains in GPR, 8 of which have significant e-values and an average size of 204 aa. *Caulobacter* TipN and *Agrobacterium* GPR differ dramatically in length (882 aa vs. 2,115 aa), but have an overall similar placement of related structural domains, N-terminal membrane-spanning domains, N- and C-terminal disordered regions, and central regions of continuous α -helices (26) (*SI Appendix, Fig. S1A and B*). TipN is distinct from GPR, however, with a much shorter central α -helical region (520 aa) and no predicted ApoLP domains.

ApoLPs have been extensively studied in eukaryotes, as they are essential for the formation of cardioprotective high-density lipoproteins (reviewed in ref. 31), where α -helical domains of ApoLPs sequester lipid droplets (reviewed in ref. 32). ApoLPs also have been identified in prokaryotes, primarily as structural components of lipid droplets (33–35). Due to data mining of bacterial genomes, their numbers are increasing; in 2015, prokaryotic ApoLPs represented approximately 16% of the total PF01442 family members, while as of October 2018, this proportion had more than doubled, to 41% (*SI Appendix, Fig. S2*). Interestingly, 272 of 326 (83%) bacterial sequences reported to date that contain predicted ApoLPs (Pfam P01442) (25) are from the Rhizobiales order, which includes *Rhizobium* and *Agrobacterium* species. Most ApoLPs in the Rhizobiales are large proteins and have an overall architecture similar to GPR, including two transmembrane domains near the N terminus.

To date, no ApoLPs have been identified in the Caulobacteriales order, consistent with the absence of ApoLP domains in TipN. Furthermore, TipN and GPR likely have distinct functions, as they participate in distinctly different means of growth, dispersed in *Caulobacter* compared with polar in *Agrobacterium*. Potentially, the large central region of GPR carrying ApoLP domain in *Agrobacterium* represents an evolutionary diversification in the Rhizobiales order compared with the Caulobacteriales order, which creates a new function (36–38).

GPR Localization. We created GFP fusions to the N and C termini of *Agrobacterium* GPR and visualized their localization. GFP

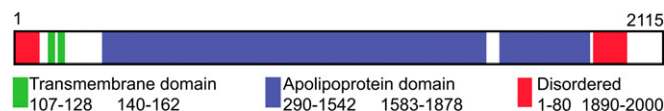


Fig. 1. GPR domain structure. Predicted transmembrane regions are shown in green, disordered regions are in red, and overlapping and/or adjacent ApoLP domains (*SI Appendix, Table S1* lists exact locations and lengths) are in blue. Other areas (white) are predominantly α -helical according to Pfam (25). *SI Appendix, Fig. S1* provides more details.

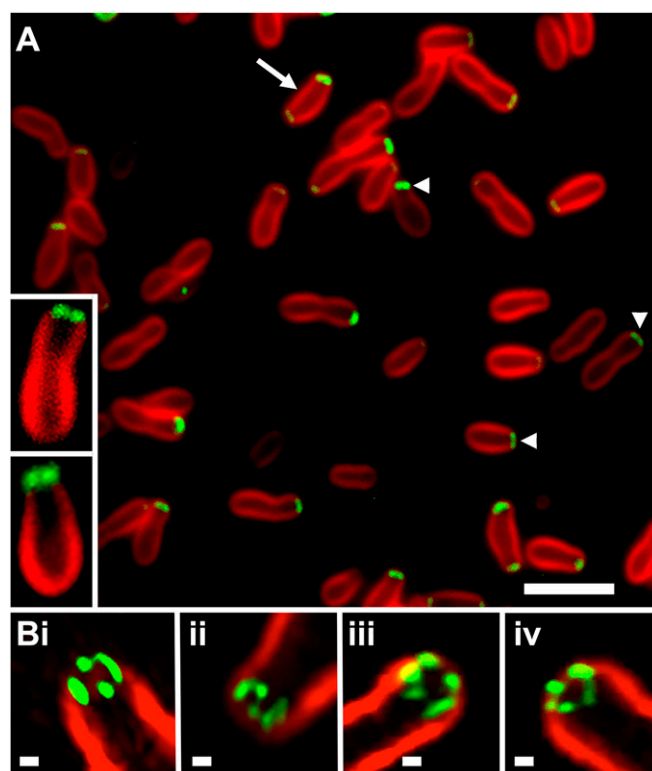


Fig. 2. Localization of GFP-GPR at the growth pole. (A) On WFM, most cells show a single broad polar focus of GFP (arrowheads), and a few cells (arrow) show additional weak localization at the opposite pole. (Insets) Cells with paired polar foci. (Scale bar: 3 μ m.) (B) SIM 3D reconstructions show subpolar localization and multiple foci. *Movies S1–S4* correspond to Fig. 2B, *i–iv*, respectively. FM4-64 stains the older nongrowing pole; weak or no FM4-64 staining occurs at the growth pole. (Scale bar: 100 nm.)

fused to the N terminus reveals a broad, almost flat region of fluorescence near the growing pole (Fig. 2A; arrowheads) that differs from the tight focus of fluorescence observed with fusions to other growth pole-specific factors, such as FtsZ, FtsA, and PopZ (6, 13). On image enlargement, this broad band appears as paired foci very near, but not at, the tip of the growth pole (Figs. 2A, *Insets* and 3A).

As paired foci in single optical sections are indicative of a potential ring structure, we used high-resolution structured illumination microscopy (SIM) to better resolve these polar foci. Indeed, GPR resolves into a ring of four to six foci subpolar to the growth pole (Fig. 2B). Some foci are round and likely represent single foci, while other foci likely represent overlapping fluorescence from two adjacent round foci, as exemplified by the cell shown in Fig. 2B, *i*. *Movies S1–S4* show different rotations of the four cells shown in Fig. 2B. These multiple foci with a unique ring-like localization at the growth pole are the basis for naming *Agrobacterium* protein A9CJ72 the GPR protein.

GFP-GPR occasionally exhibits bipolar localization (Fig. 2A, arrow), and we performed time-lapse studies to further analyze this observation (see below). C-terminal GFP fusions to GPR resulted in abnormal growth patterns with multiple ectopic growth poles (*SI Appendix, Fig. S3*) suggesting that GFP blocks an essential function at the C terminus of GPR.

GPR Function Is Sensitive to Levels of Ectopic GFP-GPR. For the images presented in Fig. 2 and throughout, we used very low levels of induction of GFP-GPR expression. Here, as in previous work, we cloned genes of interest (as fusions to GFP) into a low-copy number plasmid carrying a tightly regulated *lac* promoter (39),

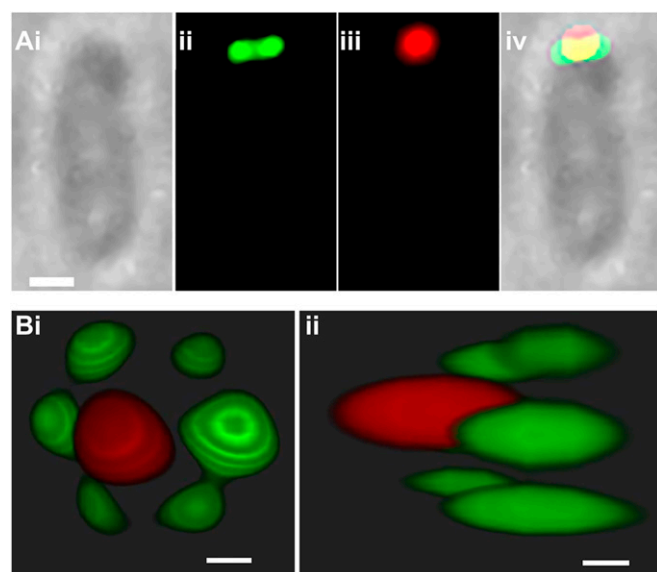


Fig. 3. Localization of PopZ-GFP and GFP-GPR. (A) PopZ-RFP fluorescence overlaps with GFP-GPR. Some PopZ-RFP extends beyond GFP-GPR toward the tip of the growth pole, while the bulk of PopZ overlaps with GPR. Compare bright field (i), green (ii), and red (iii) fluorescence with both green and red fluorescence in the merged image (iv). (Scale bar: 500 nm.) (B) SIM confirms that PopZ localizes at the tip of growth pole, while GFP-GPR is subpolar and peripheral. (i) growth-pole view; (ii) lateral view. [Movie S5](#) shows an animation of the 3D reconstruction of PopZ and GPR localization from SIM images. BF, bright field. (Scale bar: 50 nm.)

and induced protein expression by adding 2.5–10 mM isopropyl β -D-1-thiogalactopyranoside (IPTG) (6, 9, 13, 14, 19). For example, 10 mM and 5 mM IPTG resulted in expression of FtsZ-GFP or FtsA-GFP at approximately 10% of the levels of endogenous FtsZ or FtsA, respectively; these levels of ectopic induction did not alter cell morphology or cell cycle timing (6, 9, 13, 14, 19). However, for cells to remain viable during ectopic expression of GFP-GPR, it was essential to reduce the IPTG concentration to only 0.25 mM. [SI Appendix, Fig. S4](#) shows cells carrying GFP-GPR induced for expression with 0.25 mM, 0.5 mM, 1.0 mM, and 2 mM IPTG. As the concentration of IPTG increased, the fluorescence of GFP-GPR increased, as expected; however, with more GFP-GPR expression, cells produced more ectopic growth poles. Ectopic growth poles were first observed with 0.5 mM IPTG induction, but not with 0.25 mM IPTG induction, and 0.25 mM induction was sufficient to allow observation of GFP-GPR by fluorescence microscopy.

The foregoing data suggest that WT GPR function is very sensitive to ectopic expression of GFP-GPR, potentially because GFP-GPR was induced to higher levels than expected. We tested the relative protein levels of GFP-GPR compared with WT GPR under different IPTG induction conditions. [SI Appendix, Fig. S5](#) shows that 0.1 mM IPTG was not sufficient to induce GFP-GPR expression, 0.25 mM IPTG resulted in GFP-GPR expression at approximately 20% of WT levels, and 0.5 mM or 1.0 mM IPTG resulted in GFP-GPR expression at 50% or higher compared with WT. Thus, higher levels of IPTG significantly increased the ratio of the ectopic GFP fusion protein compared with endogenous WT GPR, and this may account for the altered cell shape and ectopic poles observed.

GPR Forms a Large Hexameric Ring at the Growth Pole. Wide-field fluorescence microscopy (WFM) revealed GFP-GPR as paired foci near the growth pole tip (Fig. 2). Similar WFM during coexpression of GFP-GPR with PopZ-RFP showed PopZ-RFP

closer to the tip of one end of the cell and GFP-GPR just below and overlapping PopZ-RFP (Fig. 3A). SIM imaging confirmed this localization and further revealed that GFP-GPR forms six distinct foci (Fig. 3B), as suggested by images in Fig. 2B; this notable localization is further evidenced in [Movie S5](#). Each GFP-GPR focus is approximately 50 nm in diameter, similar in size to the PopZ-RFP focus. As the space between each focus is also $\sim$ 50 nm, the ring has a circumference of $\sim$ 600 nm with a diameter of $\sim$ 200 nm.

Localization of GPR During the *Agrobacterium* Cell Cycle. Fig. 4A presents images of a single cell expressing GFP-GPR following time-lapse observation. GFP-GPR is at the growing pole until cell division (at 90 min) and then is located at the new growth poles, resulting from midcell constriction in sibling cells (at 120 min). The previous new pole has become an old pole (4), yet GFP-GPR remains at this site, reflecting that GPR is membrane-anchored. That GPR remains at this old growth pole even until the next cell division (at 180 min) may suggest the time it takes for its removal and/or degradation. These results imply that GFP-GPR observed in the sibling new cell poles is likely to be newly synthesized; that is, it is unlikely that a membrane-anchored protein would easily migrate to the new poles. [Movie S6](#) shows the same cell shown in Fig. 4.

Notably, how long GFP-GPR remains at the new old pole varies. [Movie S7](#) shows two different cells, one cell in which GFP-GPR remains at the new old pole and another cell in which it disappears relatively soon after the first cell division. The daughter that contains the original old pole does not contain GFP-GPR at its (very) old pole; in contrast, the daughter that originates from polar growth at the new pole contains GFP-GPR at its (new) old pole. These results also underscore that daughter cells are not equivalent.

To independently assess GPR localization during the cell cycle, we performed a quantitative assessment of the localization of GFP-GPR in cells coexpressing FtsZ-RFP or PopZ-RFP that each exhibit distinct time frames for their relocation from the

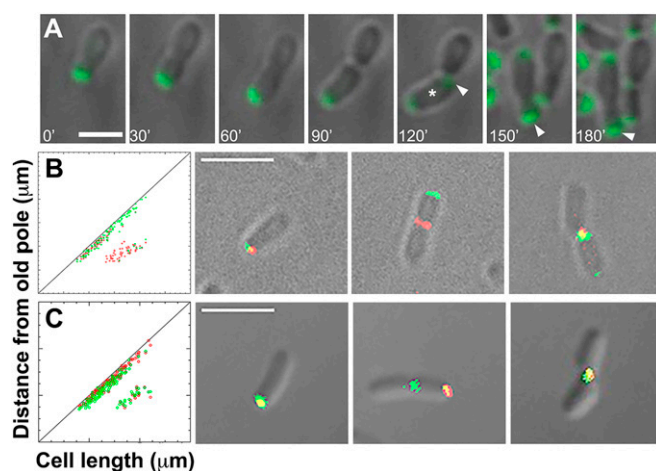


Fig. 4. GPR localization during the cell cycle and quantification of GPR, FtsZ, and PopZ localization to the growth pole or the midcell. (A) Time-lapse localization of GFP-GPR. Note that between 120 and 150 min, the lower cell (white asterisk) rotates ([Movie S7](#)) approximately 120° (clockwise), so that its growth pole faces downward at 150 min. The old pole is indicated by an arrowhead. (B and C) Coexpression of GFP-GPR and FtsZ-RFP (B) and GFP-GPR and PopZ-RFP (C). Red and green foci at the growth pole and the midcell were plotted along the diagonal or in the region between the diagonal and the x-axis, respectively, in 127 cells (B) and 100 cells (C). The tick marks on the x- and y-axes correspond to 0.5 μ m. Representative cell images are shown to the right of each graph. While red and green foci occur together at the growth pole, FtsZ-RFP can occur at the midcell without GFP-GPR (but not vice versa), and GFP-GPR can occur at the midcell without PopZ-RFP (but not vice versa). (Scale bar: A, 2 μ m; B and C, 3 μ m.)

growth pole to the midcell. FtsZ marks the new growth pole in *Agrobacterium* and then relocates to the midcell region approximately 60–70 min before septation (13), while PopZ remains at the growth pole until septation (13). Instead of elapsed time, we used cell length as a measure of progression through the cell cycle, and more than 100 green or red fluorescent foci were plotted relative to cell length (Fig. 4 *B* and *C*); foci that localize at the growth pole are plotted on the diagonal, and foci that occur at the midcell are plotted between the diagonal and the *x*-axis. GPR and FtsZ colocalize at the growth pole in shorter cells, corresponding to early stages of the cell cycle, and GPR remains at the growth pole as cells elongate up to 4 μ m. In contrast, in cells ranging from 2.6 to 4 μ m, FtsZ is rarely found at the growth pole and instead occurs at the midcell.

Coexpression of GFP-GPR and PopZ-RFP confirms that PopZ remains at the growth pole throughout the cell cycle (13) and in the longest cells, as expected. However, GPR relocates to the midcell just slightly before PopZ, as (*i*) there are a few more green foci in cells 2–3.5 μ m long, (*ii*) we detect green foci at the midcell without red foci, and (*iii*) we rarely detect red PopZ foci at the midcell without green GPR foci.

Deletion or Depletion of GPR Causes Severe Growth Defects. To test whether GPR is essential for polar growth, we created a genetic deletion of *gpr* (*SI Appendix, Fig. S6* shows deletion strategy). Δgpr cells grow slowly, with a doubling time of 4–5 h in liquid cultures (vs. 90 min for WT); however, cell cycle length varies between cells (see below). Strikingly, Δgpr cells are coccoid and vary in diameter and overall shape (Fig. 5*A*). Few cells are elongated but misshapen (Fig. 5*A*, arrows). There is little evidence of normal polar growth, except some cells produce round projections (Fig. 5*A*, encircled). A few cells produce two small “buds” (Fig. 5*A*, arrowheads) that may lead to mini cells.

Nevertheless, at least some Δgpr cells grow in liquid culture and form small colonies on agar plates. Time-lapse imaging (Fig. 5*B* and *Movie S8*) shows that round cell 1 produces cell 2, which produces cell 3; after a delay cell 2 produces cell 4, and then cell 5 forms. Thus, Δgpr daughter cells are not synchronous in their cell cycle progression. Normally, two cells should produce four cells and then eight cells; instead, we observe cell number 3 before cell number 4, and cell number 5 before cell number 6.

Δgpr daughter cells have an additional phenotype, failure to separate following division, so one can easily distinguish the order of formation of cells 3, 4, and 5 (Fig. 5*B*); such images are representative of 20% of Δgpr cells with round protrusions (Fig. 5*A*). In addition, at least 40% of the cells are unusually large and spherical; such cells grow in all directions over a long period (14 h) and produce a quartet of daughter cells that remain attached (*Movie S9*).

The coccoid phenotype of Δgpr disappeared following ectopic expression of WT GPR under control of the *lac* promoter (*SI Appendix, Fig. S7*). Surprisingly, rescue occurred following extremely low levels (0.025 mM) of IPTG (10-fold less IPTG than used previously to induce GFP-GPR expression), suggesting that cells do not require high amounts of GPR to survive. In support, higher levels of IPTG resulted in abnormal branched cells with ectopic poles. Thus, GPR levels must be tightly controlled in WT cells.

Since stable genetic deletion of *gpr* caused such a strong phenotype, we next used a riboswitch strategy (*SI Appendix, Fig. S8*) to make *RS::gpr*. Without theophylline, *RS::gpr* exhibits a coccoid phenotype identical to Δgpr , and WT cell shape is restored on addition of theophylline (Fig. 6*A* and *B*). Notably, inducible expression of GFP-GPR with low levels of IPTG (0.025 mM) also rescues *RS::gpr* grown in the absence of theophylline (Fig. 6*C*).

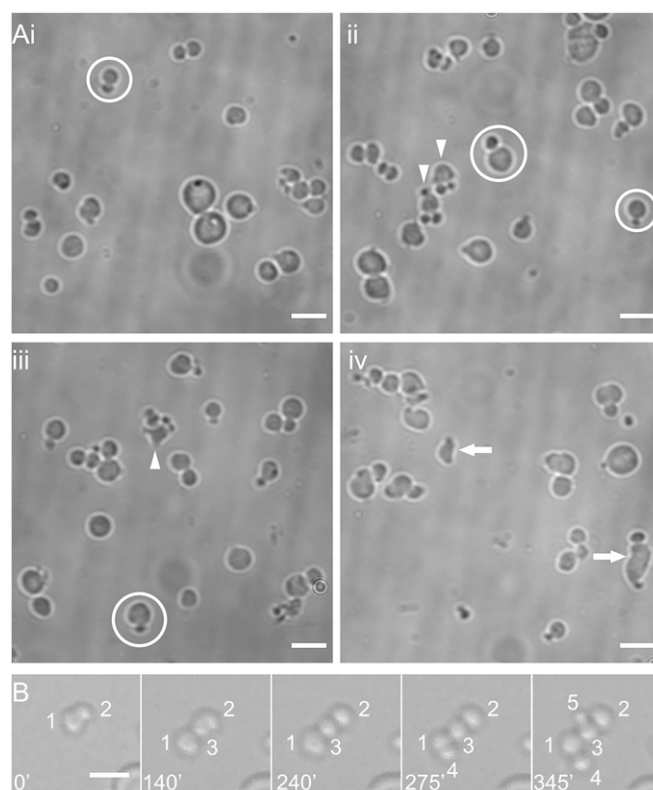


Fig. 5. Deletion of GPR results in spherical cells. (*A*) Δgpr cells are mostly coccoid. Some cells appear to be making new round cells (encircled), a few cells produce two buds (arrowheads), and a few cells are elongated but misshapen (arrows). (*B*) Time-lapse imaging of two dividing cells, labeled 1 and 2 (0 min), giving rise to cells numbered 3, 4, and 5 (images derived from *Movie S8*). The times shown were chosen to illustrate newly arising or dividing cells and do not represent a constant time difference between each panel. (Scale bar: 3 μ m.)

Loss of GPR Impacts Localization of PopZ. PopZ localizes very close to GPR (Fig. 3), and GPR relocates to the midcell just before PopZ. To test whether PopZ position is dependent on GPR, we monitored PopZ localization following depletion of GPR in *RS::GPR* (Fig. 6*D* and *E*). In GPR-depleted cells, PopZ-GFP appears mostly diffuse, with a few random foci in the misshapen cells (Fig. 6*D*). WT PopZ-GFP localization is rescued when this strain is grown in the presence of theophylline (Fig. 6*E*). Thus, PopZ can still form foci on its own, but that proper localization to the cell pole requires rod shape morphology, dependent on GPR. These data demonstrate the utility of *RS::gpr* to assay phenotypic effects following depletion of GPR.

Discussion

We have identified a previously unknown protein, which we named GROWTH POLE RING (GPR), with a striking localization as a ring of six foci at the growth pole in *A. tumefaciens*. GPR plays a major role in polar growth, as its absence leads to severe morphological defects, including loss of cylindrical cell shape. In corollary, its overproduction leads to ectopic growth poles; the finding that overproduction only slightly above WT levels leads to multiple growth poles implies that GPR expression must be tightly regulated. While prokaryotic proteins with single ApoLP domains (associated with lipid droplets) have been reported (33), to date, GPR is the first prokaryotic protein identified with an extended multiple ApoLP domain structure anchored in the inner membrane. GPR potentially represents a

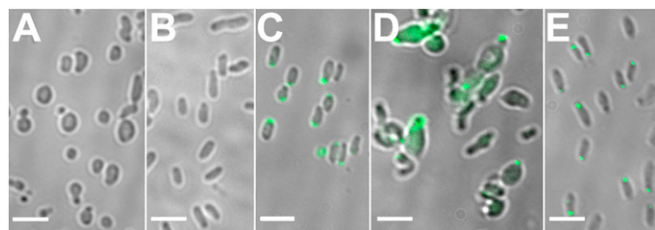


Fig. 6. Riboswitch-controlled depletion of GPR mimics stable genetic deletion of GPR and alters PopZ-RFP localization. *RS::gpr* cells were imaged at 24 h after growth without theophylline (A) or at 24 h after growth in 0.5 mM theophylline (B). Expression of GFP-GPR was induced with 0.025 mM IPTG for 24 h in the *RS::gpr* strain grown in the absence of theophylline (C). PopZ-RFP was introduced into *RS::gpr* and grown in the presence of 0.25 mM IPTG without theophylline (D) or 0.25 mM IPTG plus 0.5 mM theophylline (E).

new class of proteins with a critical role in polar growth and cell morphogenesis in the Rhizobiales.

We propose that GPR functions as a scaffold for assembly of polar growth machinery essential for peptidoglycan (PG) and/or phospholipid (PL)-dependent membrane biogenesis (Fig. 7A). Polar growth necessitates a mechanism to ensure that the growing pole is not amorphous or unstructured. This need is compounded by the fact that *A. tumefaciens* lacks an underlying cytoskeleton, such as provided by the actin homolog MreB in model systems like *E. coli* and *B. subtilis*, where MreB serves as a scaffold for circumferential PG synthesis during cell elongation by interspersed growth (40–42). Indeed, the numerous ApoLP domains in GPR support the hypothesis that GPR provides a scaffold for membrane assembly, as ApoLPs are well known to sequester lipids in eukaryotes (32). The dramatically abundant α -helical regions of GPR, including a continuous stretch of more than 1,700 amino acids, may provide a hydrophobic environment for lipid assembly.

Mycobacterium also grows by polar growth and produces a polar complex that contains at least three PG synthetic enzymes (MurG, GlfT2, and Pks13) (43). This complex resides in a band called the subpolar space, 0.5 μ m from the tip of the growth pole, and the diameter of the complex corresponds to the diameter of the mature cylindrical cell. In striking contrast, the GPR ring complex localizes approximately 100 μ m below the tip of the new pole, and the diameter of the GPR ring is only approximately 200 nm. These latter data reveal that *Agrobacterium* polar growth initially occurs in a much narrower ring of activity. Indeed, independent measurements of new pole cell diameter in different-sized new cells demonstrated that approximately one-third of the upper part of the new cell is of significantly narrower diameter during the early stages of polar growth, and later, PG synthesis increases the new cell diameter until it corresponds to the diameter of the old cell compartment just before division (9). *Mycobacterium* and *Agrobacterium* present surprisingly different examples of what might comprise the geometry of cell wall synthesis complexes during polar growth. Nevertheless, they both suggest a general mechanism for polar growth via a ring structure near the growth pole(s).

How might GPR foci be organized to be ~50 nm apart in a GPR hexamer of ~200 nm diameter? An extended linear string of 1,700 helically arranged amino acids would be ~600 nm long. More likely, GPR ApoLP domains fold into specific 3D shapes. The ApoLP structure is very dynamic, and monomers fold as bundles of helices or extended curved structures in the presence of lipids (44, 45). ApoLPs form two types of complexes of ~9–10 nm diameter when associated with lipids: (i) three ApoLPs arranged in a trefoil with their alpha helical regions on the surface of lipid spheres or (ii) two stacked ApoLPs around a lipid disk (reviewed in ref. 32).

Our data suggest that the 1,700-aa central region of GPR contains approximately eight ApoLPs, each ~200 aa long. These eight ApoLPs may provide an extremely flexible region that may undergo significant rearrangements from bundled to extended configurations during the assembly of membranes and PG. Based on the structure of ApoLP A4 (44), Fig. 7B graphically depicts two ApoLP domains in GPR, alternating between helix bundles and extended conformations. It is highly likely that additional, as-yet unidentified proteins may be structural and functional components of the GPR hexamer. Future investigations using a variety of approaches likely will lead to abundant and informative discoveries regarding GPR function during polar growth in the Rhizobiales.

Materials and Methods

Strains and Cell Growth Conditions. The strains and plasmids used in this study are listed in *SI Appendix, Table S2*. WT *A. tumefaciens* C58 was transformed with the relevant plasmids and grown at 28 °C in Luria Broth (LB) medium containing appropriate antibiotics.

Bioinformatics. We used TipN protein from *C. crescentus* (CC_1485 from strain CB15) as the query in a BLAST search of the Uniprot database Microbial Proteomes with default search parameters (24). GPR (Atu1348) was identified from a manual inspection of the list of hits generated by this search. GPR domains were identified using Pfam (25) and Phyre2 (26).

Molecular Cloning and Strain Construction. Standard molecular cloning techniques were used (46). To construct pJZ251 (*P_{lac}::gpr-gfp*), *gpr* was amplified by PCR from *A. tumefaciens* genomic DNA with NdeI sites at each end and subcloned into pCR2.1-Topo (Thermo Fisher Scientific), and then *gpr* was isolated from NdeI digestion and ligated to similarly digested pJZ210. A similar strategy was used to construct pJZ253 (*P_{lac}::gfp-gpr*), except that *gpr* was PCR-amplified with 5' AvrII and 3' KpnI sites.

Δgpr and *RS::gpr_{At}* were created as described previously (14, 19). In brief, C58 was transformed with pJZ298 (Δgpr) or pJZ274 (*RS::gpr*) (*SI Appendix, Table S2*), followed by selecting for a single cross-over into the genome by growth on carbenicillin and then selecting for a second recombination by growth on sucrose. Δgpr and *RS::gpr_{At}* were verified by PCR amplification of the relevant genomic region and sequencing.

Time-Lapse and Fluorescence Microscopy. Unless noted otherwise, lactose-inducible expression was achieved by diluting overnight cultures to 10^8 cells/mL and adding 0.25 mM IPTG for 4–5 h before single-time or time-lapse

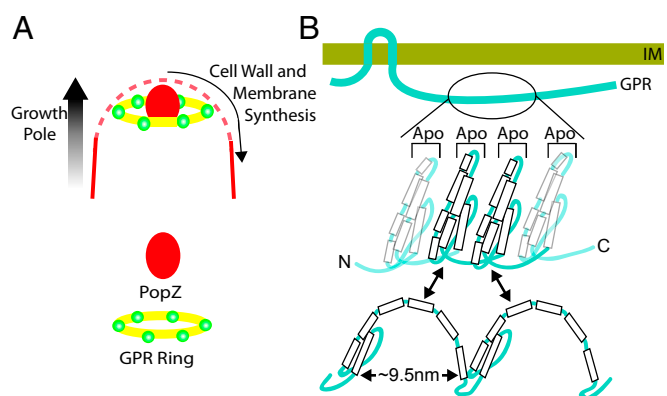


Fig. 7. Models of the GPR hexamer ring (A) and monomer secondary structure (B). (A) Six GPR complexes (green spheres) form a (yellow) ring-shaped scaffold ~100 nm from the tip of the growth pole to organize membrane and PG synthesis. The solid red line indicates outer membrane (OM) staining with red lipophilic dye FM4-64; the dashed red line indicates OM stains less intensively at the growth pole (6). (B) Flexible/reversible structure (based on a model for ApoLP A1; figure 5B in ref. 44) of GPR ApoLP domains (extended vs. compact) may facilitate a variety of configurations between adjacent GPR monomers that maintain the diameter of the GPR scaffold. C, C terminus; IM, inner membrane; N, N terminus.

imaging by deconvolution fluorescence microscopy as described previously (13). Long-term time-lapse imaging of Δgpr was done using the CellASIC ONIX system (EMD Millipore) on a Zeiss Axio Observer Z1 microscope. For quantitative analyses, C58 cells were cotransformed with pJZ253 (GFP-GPR) and pJZ269 (PopZ-RFP) or with pJZ253 and pTC077 (FtsZ-RFP); cell length and position of the fluorescent foci were determined using ObjectJ (47). All images were processed using Fiji software (48).

Superresolution Microscopy. Superresolution images were captured using a Zeiss Elyra PS.1 structured illumination microscope (SIM) equipped with a Zeiss Plan-Apochromat 100 $\times$ /1.46 oil immersion objective lens and a pco.edge scientific complementary metal-oxide semiconductor camera with a 1.6 $\times$ tube lens. GFP-GPR fluorescence was detected with 488-nm laser excitation. FM4-64 or RFP fluorescence was detected with 561-nm laser excitation. The lateral pixel size was 41 nm $\times$ 41 nm in the recorded images. Z-stacks were acquired by capturing 20 slices with a 0.1- μ m step size. The 3D SIM images were reconstructed using ZEN 2012 (black edition) (Zeiss) and processed with Imaris 8.1 (Bitplane).

Protein Analysis. Preparation of *Agrobacterium* cell lysates, gel electrophoresis, and Western blot analysis were performed as described previously (49). The primary antibody was commercially prepared rabbit polyclonal anti-GPR antibody (Genscript) generated against the peptide FEEKPETRKATKPK (amino acids 27–41) and used at a dilution of 1:1,000.

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