

RESEARCH ARTICLE SUMMARY

BACTERIAL PILI

Architecture of the type IVa pilus machine

Yi-Wei Chang, Lee A. Rettberg, Anke Treuner-Lange, Janet Iwasa, Lotte Sogaard-Andersen, Grant J. Jensen*

INTRODUCTION: Type IVa pili are bacterial cell surface structures that perform critical functions in motility, surface adhesion, virulence, and biofilm formation. Type IVa pili are anchored in the cell envelope and pull cells forward through cycles of extension, adhesion to surfaces, and retraction, all powered by the type IVa pilus machine (T4PM). Although the structures and connectivities of the 10 core T4PM proteins and minor pilins have already been determined, the overall architecture of the

T4PM and its extension and retraction mechanisms have not.

RATIONALE: To elucidate the architecture of the intact T4PM, we directly imaged T4PMs within intact *Myxococcus xanthus* cells by cryo-electron tomography. Mutants that either lacked T4PM components or contained individual T4PM proteins fused to a tag were then imaged. Difference maps revealed the locations of all components of the T4PM machine. Hypothetical models

were then built by fitting the known atomic structures of the components together in their relative positions.

RESULTS: Both piliated and nonpiliated T4PMs are multilayered structures that span the entire cell envelope. T4PMs include an outer membrane pore, three interconnected periplasmic ring structures and another in the cytoplasm, a cytoplasmic disc and dome, and a periplasmic stem. The PilQ secretin forms the outer membrane pore; TsaP forms a periplasmic ring around PilQ; peri-

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plasmic domains of PilQ together with PilP constitute the mid-periplasmic ring; and the globular domains of PilO and PilN constitute the lower periplasmic ring and connect

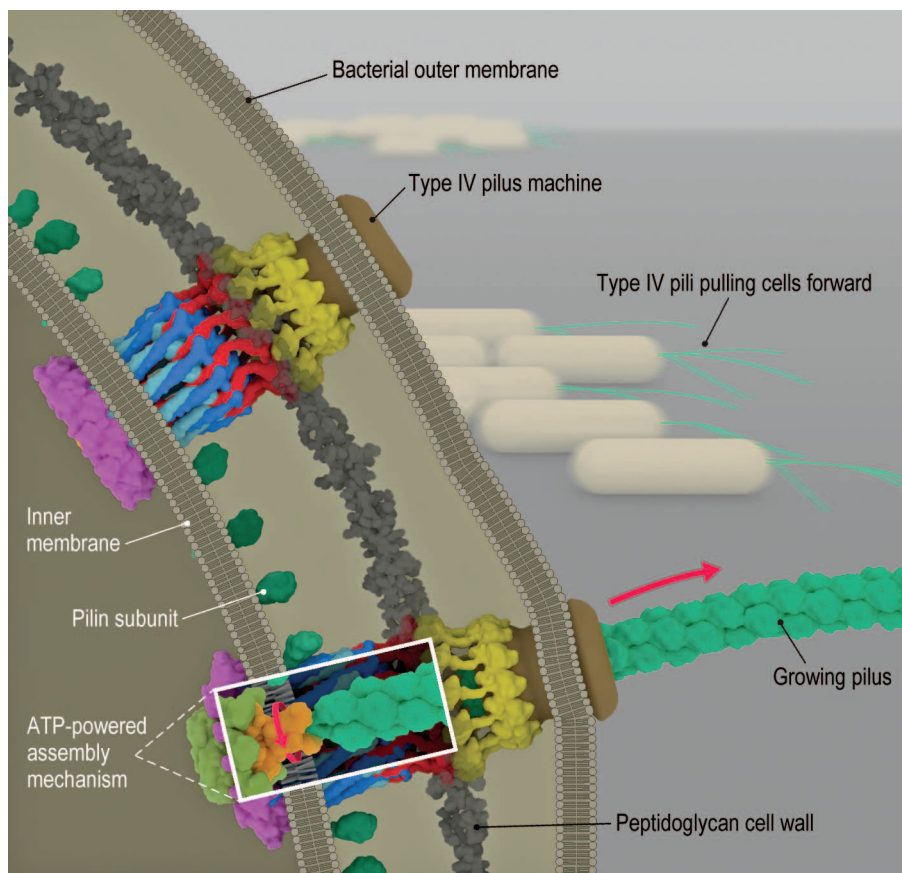
via coiled coils across the inner membrane to PilM, which forms the cytoplasmic ring. The cytoplasmic domains of the inner membrane protein PilC form the cytoplasmic dome on the T4PM axis inside the PilM ring. The short stem in the nonpiliated state is composed of minor pilins and PilA, the major subunit of the pilus. In the piliated state, the pilus extends from the cell exterior through the PilQ pore and the periplasmic rings to PilC in the inner membrane. In the piliated structure, the hexameric adenosine triphosphatases (ATPases) PilB and PilT bind in a mutually exclusive manner to the base of the T4PM, where they appear as the cytoplasmic disc during extensions and retractions, respectively.

Next, we asked whether the known atomic structures of the proteins could be fit within the map where our imaging results indicated, while still satisfying all known constraints of size, connectivities, and interfaces. This successful effort resulted in “pseudo-atomic” working models of both states of the T4PM. The models suggest that through ATP hydrolysis, PilB rotates PilC, incrementally moving it into positions that facilitate incorporation of new PilA subunits one by one from the inner membrane onto the base of the growing helical pilus. Pilus retraction is driven by replacement of PilB with PilT, which rotates PilC into positions that promote PilA departure from the base of the pilus back into the membrane.

CONCLUSION: We determined the architecture of the T4PM in the piliated and nonpiliated states and mapped all known components onto this architecture, producing a complete structural map of the T4PM. The results illustrate how the structure and function of macromolecular complexes that defy purification and traditional structural approaches can nonetheless be interrogated through cryo-electron tomography of intact cells and model building. ■

The list of author affiliations is available in the full article online.

*Corresponding author. E-mail: jensen@caltech.edu
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Bacterial type IVa pilus machine (T4PM). Two T4PMs are depicted on the left, spanning the envelope of a Gram-negative bacterial cell. T4PMs extend and retract pili to pull cells forward. The structural data presented here support the hypothesis that ATP hydrolysis in the cytoplasm causes an adapter protein in the inner membrane to rotate, facilitating the transfer of pilin subunits from the inner membrane onto the growing pilus. The process is reversed during retraction.

RESEARCH ARTICLE

BACTERIAL PILI

Architecture of the type IVa pilus machine

Yi-Wei Chang,^{1,2} Lee A. Rettberg,² Anke Treuner-Lange,³ Janet Iwasa,⁴ Lotte Søgaard-Andersen,³ Grant J. Jensen^{1,2*}

Type IVa pili are filamentous cell surface structures observed in many bacteria. They pull cells forward by extending, adhering to surfaces, and then retracting. We used cryo-electron tomography of intact *Myxococcus xanthus* cells to visualize type IVa pili and the protein machine that assembles and retracts them (the type IVa pilus machine, or T4PM) in situ, in both the pilated and nonpilated states, at a resolution of 3 to 4 nanometers. We found that T4PM comprises an outer membrane pore, four interconnected ring structures in the periplasm and cytoplasm, a cytoplasmic disc and dome, and a periplasmic stem. By systematically imaging mutants lacking defined T4PM proteins or with individual proteins fused to tags, we mapped the locations of all 10 T4PM core components and the minor pilins, thereby providing insights into pilus assembly, structure, and function.

Type IVa pilus machines (T4PMs) are part of a superfamily of bacterial and archaeal multi-protein assemblies that also include type IVb pilus machines, type II secretion systems (T2SS), DNA uptake systems, and archaeal flagella (1, 2). T4PMs are involved in cell motility (3), host adhesion (4), predation (5), DNA uptake (6), biofilm formation (7, 8), and protein secretion (9). T4PMs consist of an extracellular pilus fiber (T4P) and a cell envelope-spanning complex that we refer to here as the basal body. A key feature of T4Ps is their ability to undergo cycles of extension and retraction (10, 11). Powered by a T4P-assembly adenosine triphosphatase (ATPase), the basal body extracts pilin monomers from the inner membrane (IM) and appends them onto the base of the helical pilus fiber, pushing the fiber outward. The fiber has a diameter of ~6 nm, can extend up to several micrometers in length, and can adhere to specific surfaces (12). Subsequently, the basal body, powered by a T4P-disassembly ATPase, extracts monomers from the pilus base back into the membrane (13). T4PMs retract at rates up to 1 μm/s and generate forces as high as 150 pN to pull the cell forward (14, 15), making T4PMs the strongest molecular motors known. T4Ps have been identified as important virulence factors in several human pathogens, including *Neisseria gonorrhoeae*, *N. meningitidis*, and *Pseudomonas aeruginosa* (16).

T4PMs exhibit a multilayered structure spanning the cell envelope of *M. xanthus*

To determine the structure of the T4PM in vivo, we used cryo-electron tomography (also known

as electron cryotomography) to image the cell poles of *Myxococcus xanthus*, a ubiquitous rod-shaped soil-residing predatory bacterium (Fig. 1A). (Examples of a tilt series and a cryotomogram are shown in movies S1 and S2, respectively). T4Ps with diameters of ~6 nm were easily recognized on the cell surface with their basal bodies spanning the cell envelope (Fig. 1B, white arrows). Basal bodies exhibited three layers of densities in the periplasm and a fourth in the cytoplasm (Fig. 1C). In the vicinity of pilated T4PMs, we also observed “empty” nonpilated basal bodies with similar structures but without long fibers attached (Fig. 1B, black arrow, and Fig. 1D).

To reveal details, we generated subtomogram averages of the pilated T4PM and nonpilated basal bodies with local resolution between ~2.5 nm and 4.5 nm (Fig. 1F and fig. S1, A and B). We later determined that a $\Delta pilB$ mutant (which lacks the PilB T4P-assembly ATPase and cannot assemble T4P) produced better-quality images of the empty basal body (likely due to the basal bodies being stalled in a “pilus preassembly” state and therefore more structurally homogeneous than in the wild type; see below), so we used the average from this strain for subsequent structural interpretation (Fig. 1G).

We found that the structure of the pilated T4PM basal body comprises an outer membrane (OM)-spanning pore including a ring immediately below the OM, two distinct rings in the periplasm, another ring in the cytoplasm surrounding a disc-like structure, and a long stem originating at the IM and passing through the periplasmic rings and OM pore (Fig. 1F). The structure of the empty basal body shows several similar structural features, including an OM pore with a ring immediately below the OM, two periplasmic rings, and a cytoplasmic ring, as well as a gate density in the OM

pore, connections between the OM pore and the mid-periplasmic ring, connections between the lower periplasmic ring and the IM, a much shorter stem, and no cytoplasmic disc (Fig. 1G). In the absence of the cytoplasmic disc, a cytoplasmic dome is also apparent. Aligning the averages of pilated and empty basal bodies with the IM reveals clear conformational changes upon piliation (movie S3). Going from the nonpilated to the pilated state, the OM pore is ~2 nm farther away from the mid-periplasmic ring (13.9 to 15.7 nm). Also, the diameter of the cytoplasmic ring in the pilated state is greater by ~4 nm (18.4 versus 14.4 nm), possibly because of its association with the cytoplasmic disc (Fig. 1, E and H).

Mapping components in the molecular envelope by imaging T4PM mutants

Ten highly conserved proteins are known to constitute the T4PM (16, 17). PilA, the major pilin protein, contains an N-terminal hydrophobic α helix and alternates between being anchored individually in the IM or bundled with other PilA N-terminal α helices to form the central pilus core (18). The remaining nine proteins are divided into three subgroups according to their location and function: the OM pore complex (PilQ and TsaP); the alignment complex (PilM, PilN, PilO, and PilP); and the motor complex (PilB, PilT, and PilC) (19). To systematically localize each component within the basal body, we imaged a series of *M. xanthus* mutants with individual T4PM proteins either missing or fused to a superfolder green fluorescent protein (sfGFP) (20) tag. Difference maps between the resulting subtomogram averages of the T4PM mutants and the wild-type structures were then calculated (Fig. 2). Combined with information already available about the accumulation, subcellular localization, and incorporation of individual T4PM proteins into the T4PM basal bodies of these mutants (fig. S2) and their connectivities and structures (fig. S3) (17, 18, 21–40), these maps allowed us to pinpoint each component within the T4PM (Figs. 2 and 3A).

PilQ

A previous study showed that knocking out *pilP* accelerates degradation of PilM, PilN, and PilO and causes mislocalization of PilC, whereas PilQ and TsaP remain stable and at the cell pole, forming a rudimentary T4PM basal body consisting only of PilQ and TsaP (fig. S2) (21). This suggests that PilP is crucial for stabilizing and linking other components to the OM pore complex. We therefore first imaged the $\Delta pilP$ mutant to obtain the structure of the OM pore complex (PilQ and TsaP) alone (Fig. 2, B1). The average showed a cylindrical channel in the OM with a clear gate and a large periplasmic vestibule. The overall structure is reminiscent of single-particle reconstructions of secretion channel complexes in T4PM, T2SS, and type III secretion systems (T3SS) (41–44) (fig. S4). Superposing the highly conserved gate and periplasmic vestibule regions, our in vivo structure was seen to be markedly longer in the transmembrane region than the structures generated from single-particle analyses, probably because the detergent

¹California Institute of Technology, Pasadena, CA 91125, USA.

²Howard Hughes Medical Institute, Pasadena, CA 91125, USA.

³Max Planck Institute for Terrestrial Microbiology, 35043 Marburg, Germany. ⁴University of Utah, Salt Lake City, UT 84112, USA.

*Corresponding author. E-mail: jensen@caltech.edu

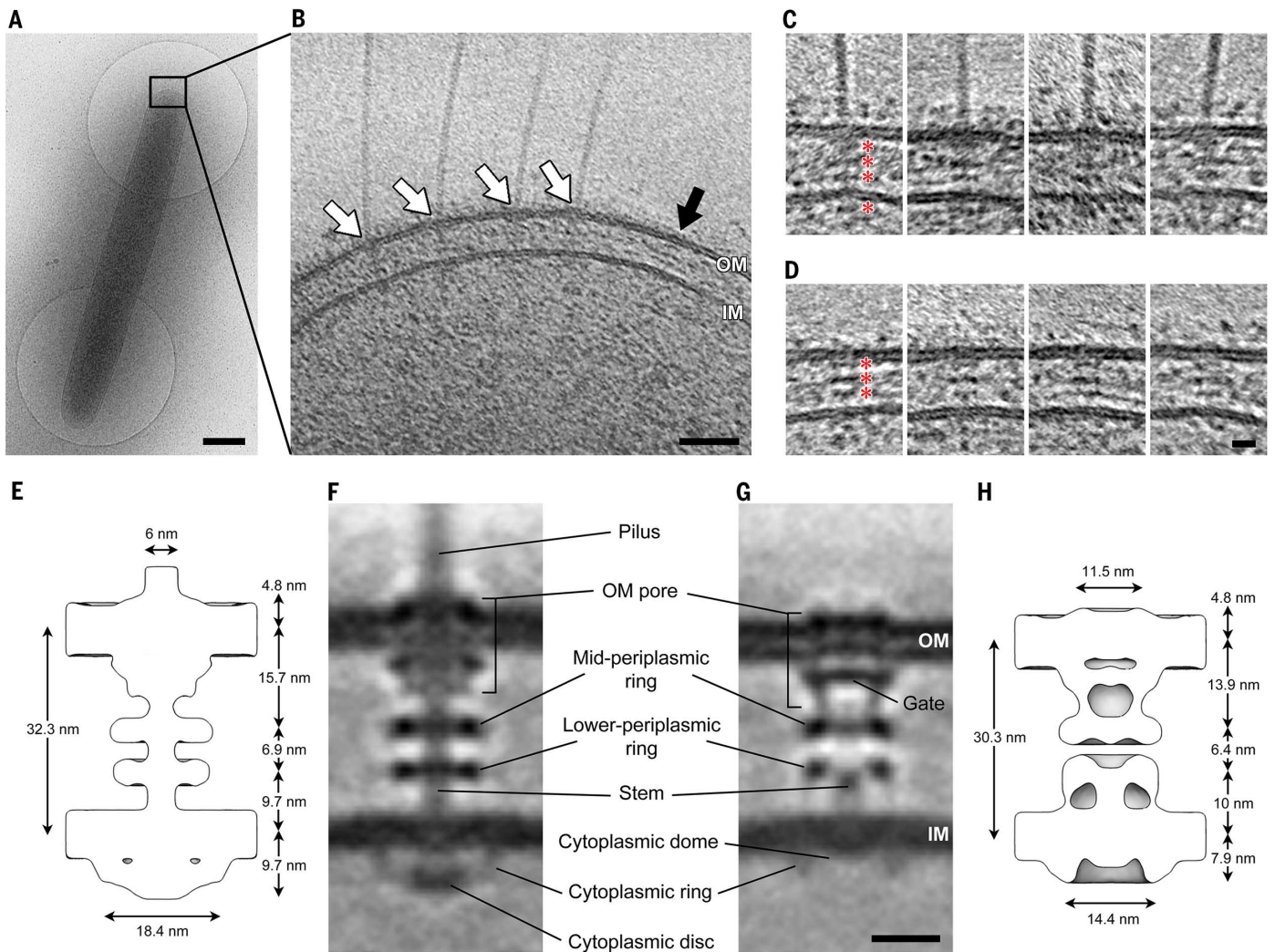


Fig. 1. Visualizing the T4PM in intact *M. xanthus* cells. (A) A frozen-hydrated *M. xanthus* cell on an EM grid. (B) Slice through tomogram of the cell pole. White arrows, pilated T4PM basal bodies; black arrow, an empty T4PM basal body. Outer and inner membranes (OM and IM) are indicated. (C and D) Examples of slices through subtomograms containing pilated and empty T4PM basal body structures, respectively. Red asterisks denote periplasmic and cytoplasmic den-

sity layers. (E and H) Schematic envelopes of the subtomogram averages of wild-type pilated (E) and $\Delta pilB$ empty (H) T4PM basal bodies, showing their molecular dimensions. (F and G) Slices through subtomogram averages of wild-type pilated (F) and $\Delta pilB$ empty (G) T4PM basal bodies, with annotations of structural features. Scale bars, 500 nm (A), 50 nm (B), 5 nm [(C) and (D)], 10 nm [(F) and (G)].

solubilization used in the single-particle reconstructions fails to support this structure. By comparing the $\Delta pilP$ mutant structure to that of the wild-type empty basal body (Fig. 2, B2), and noting that the single-particle reconstructions contained no TsaP, we infer that PilQ forms the OM channel as well as part of the mid-periplasmic ring (Fig. 2, B3).

PilP

Because PilP is known to interact directly with PilQ (21, 26) and the $\Delta pilP$ structure shows a decrease in density of the mid-periplasmic ring (Fig. 2, B2), we hypothesized that PilP is part of the mid-periplasmic ring. To confirm this, we imaged a mutant in which PilP is fused to sfGFP, giving rise to an active fusion protein (fig. S2). Consistently, in this strain, additional densities

appeared at the periphery of the mid-periplasmic ring (Fig. 2, C1 to C5).

TsaP

Single-particle EM reconstructions have shown that TsaP forms a ring-like structure surrounding the PilQ channel, but only top views were obtained (17). To localize TsaP in the OM pore complex in three dimensions, we imaged a $\Delta tsaP$ mutant (Fig. 2, D1 and D3). In the $\Delta tsaP$ mutant, all other T4PM proteins accumulate and are incorporated into the T4PM basal body as in the wild type (fig. S2) (17). Difference maps pinpointed TsaP's location to the upper periplasmic region of the OM pore complex just beneath the OM in both the pilated and empty basal bodies [Fig. 2, D2 (white arrow), D4 (white arrow), and D5]. Lack of TsaP caused disengagement of the OM channel

from the mid-periplasmic ring in the empty basal body (Fig. 2, D3, black arrow; note that the extensive systematic differences in Fig. 2, D4, reveal a global shift of the periplasmic rings and IM), making the overall length of the complex similar to that of the pilated basal body. In a previous study, lack of TsaP resulted in accumulation of pilus fibers in the periplasm in *N. gonorrhoeae* and formation of fewer pili in *M. xanthus*, suggesting a role for TsaP in correct PilQ channel function (17). Consistently, during our imaging, we observed that in the $\Delta tsaP$ mutant, about one-tenth as many pili were found on cell poles relative to wild-type cells (table S1).

PilO and PilN

Because knocking out *pilO* disrupts the entire alignment complex and causes mislocalization of

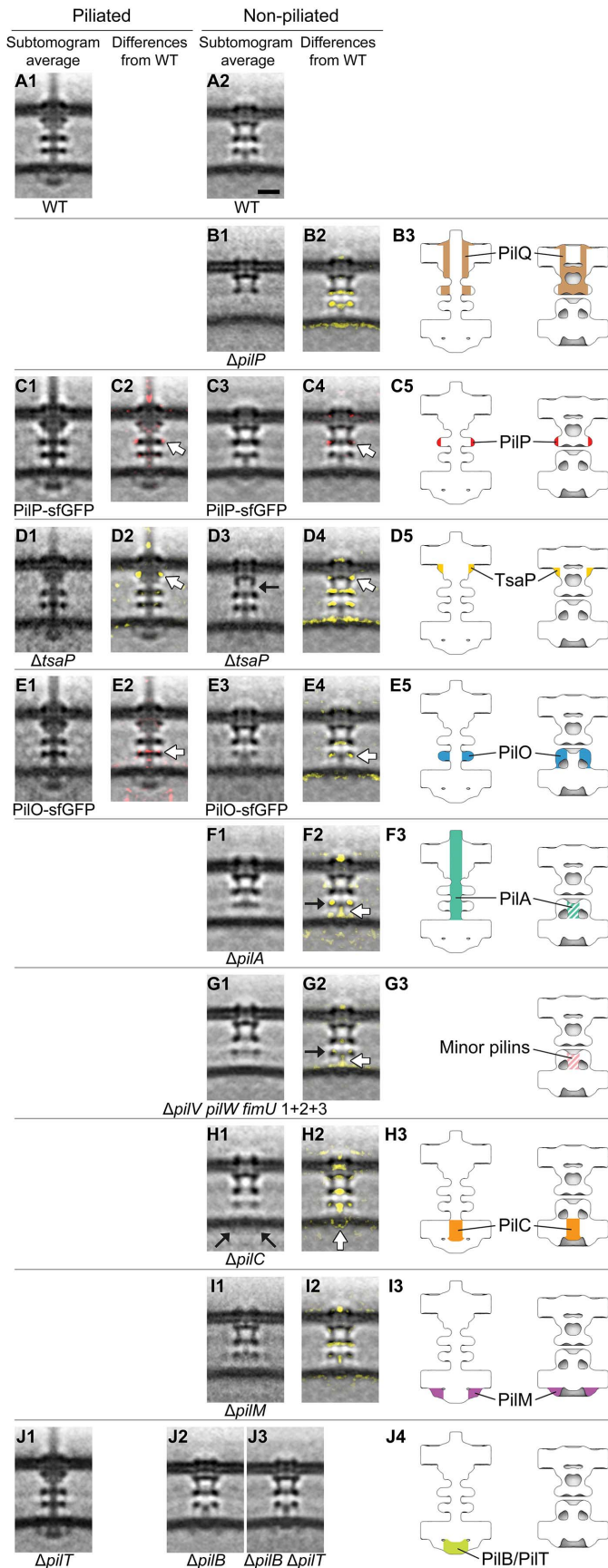


Fig. 2. Mapping T4PM components. First and third columns: Central slices of subtomogram averages of piliated and empty T4PM basal bodies, respectively, from different *M. xanthus* strains. Second and fourth columns: Differences in the T4PM mutant structures versus the wild type (red and yellow colors respectively denote addition and omission of densities, with opacities of 10%, 20%, 30%, 40%, and 50% corresponding to density differences of 1, 1.5, 2, 2.5, and 3 standard deviations, respectively, overlaid on the wild-type subtomogram averages). White arrows indicate the component locations identified by the difference maps. Fifth column: Schematic representations of piliated (left) and empty (right) T4PM basal bodies showing each identified component location. Scale bar in A2, 10 nm (applies to columns 1 through 4).

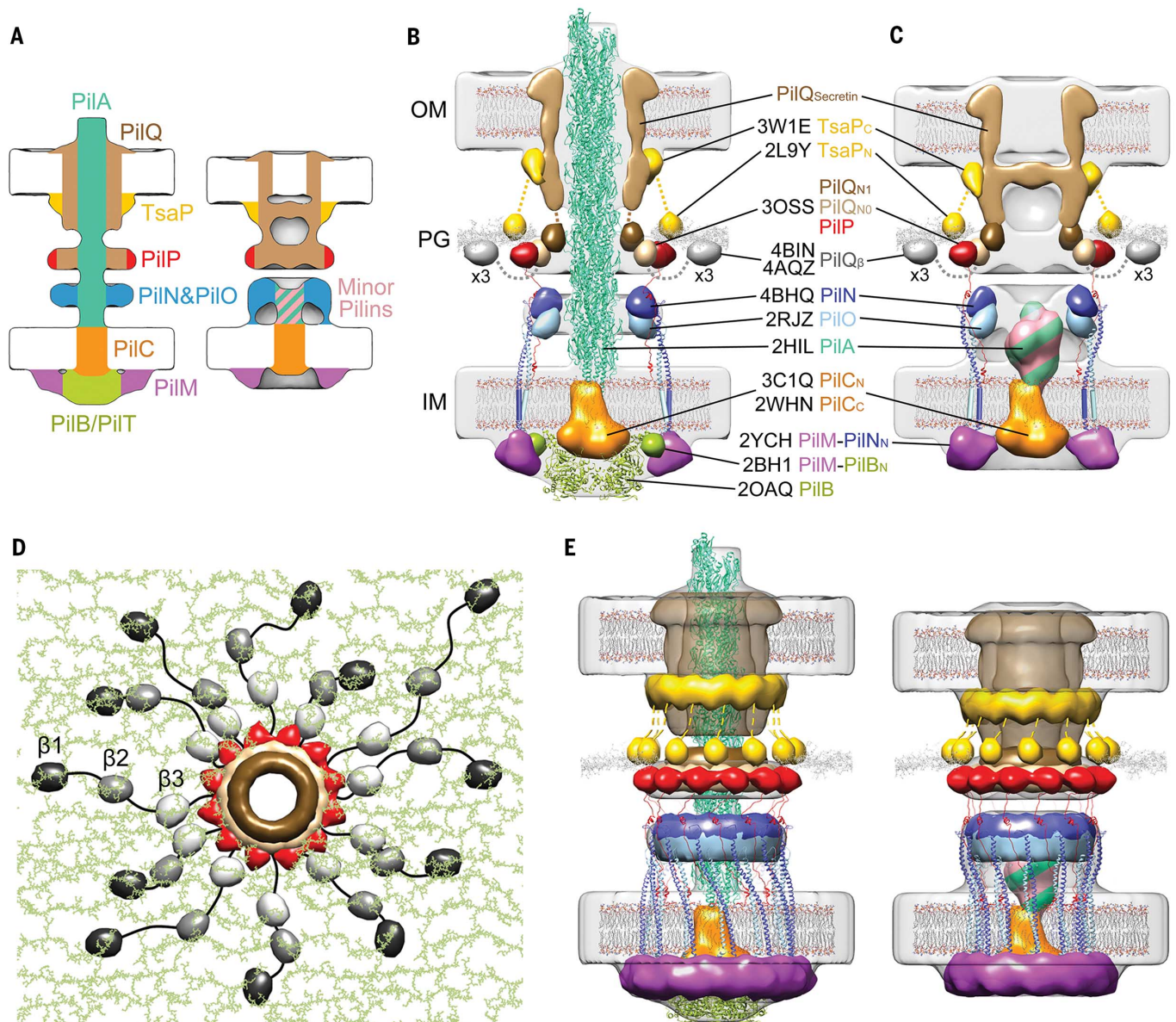


Fig. 3. Architectural models of the T4PM. (A) Summary schematics showing the component locations identified in the pilated and empty T4PM basal body structures. (B and C) Central slices of the architectural models of pilated and empty T4PM basal bodies, respectively, in which atomic models of T4PM components are placed in the *in vivo* envelopes according to the component maps in (A) and previously reported constraints and filtered to 3-nm resolution. (The process of how each component was placed is detailed in Movie 1 and the supplementary materials.) Models of each component are colored as in (A), with the transmembrane segments of PiIN and PiIO shown as cylinders; “x3” indicates three AMIN domains per PiIQ monomer, only one of which is shown.

PiIC, leaving only PiIQ and TsaP in the basal body (fig. S2) (21), we imaged a mutant with an active PiIO-sfGFP fusion protein to map the location of PiIO (fig. S2). In the pilated structure, we observed additional density in the lower periplasmic ring (Fig. 2, E1 and E2), but in the empty basal body structure, we observed decreased density in this ring (Fig. 2, E3 and E4). Both results suggest that PiIO localizes to the lower periplasmic ring

(Fig. 2, E5), because the sfGFP tag likely added density to the ring in the pilated form but perturbed the ring in the empty basal body. As with *pilO*, a $\Delta pilN$ mutant also disrupts the entire alignment complex and also causes mislocalization of PiIC, leaving only PiIQ and TsaP in the basal body (fig. S2) (21). So far, we have been unable to generate a functional PiIN protein fused to a tag. Therefore, the same method could not be

Note that the empty T4PM basal body is shown with five PiIA major pilin subunits in the short stem; however, the short stem likely also contains minor pilins. (D) Top view of the PiIP HR domains and the PiIQ NO and N1 domains in the architectural model [colored as in (B) and (C)], with PG model (colored green) as background; 36 AMIN (β) domain models from 12 PiIQ proteins are randomly placed on PG and connected by long flexible linkers (black) with lengths within 20 nm between $\beta 1$ and $\beta 2$ domains (70 residues), 12 nm between $\beta 2$ and $\beta 3$ domains (40 residues), and 12 nm between $\beta 3$ and NO domains (40 residues). (E) Overall architectural models of pilated (left) and empty (right) T4PM basal bodies. For clarity, the PiIQ AMIN domains displayed in (D) are not shown.

used to localize PiIN in the T4PM basal body. Nonetheless, the two structural homologs PiIO and PiIN interact directly (21, 30), likely forming heterodimers (30). Therefore, we assume that PiIN is also located in the lower periplasmic ring.

PiIA and minor pilins

In the wild-type pilated basal body structure, we observed a rod-like stem structure that passes up

from the IM through the lower periplasmic ring, mid-periplasmic ring, and OM pore. This stem has the same diameter as the PilA helical polymer (18) (~6 nm) and is directly connected to the pilus outside of the cell. Also, this long stem is missing in the empty basal body structure. These observations suggest that the stem is the part of the pilus fiber located in the periplasm and is associated with the basal body. In the empty basal body, a short stem is also present between the lower periplasmic ring and the IM. To investigate whether this remaining short stem is formed by PilA or by other T4PM components, we first imaged a $\Delta pilA$ mutant that lacks the major pilin PilA. As expected (45), no T4P formed in this mutant.

In agreement with the observation that the T4PM assembles in a $\Delta pilA$ mutant (fig. S2) (21), we were able to identify empty basal body structures in the cells and generate an average of the $\Delta pilA$ mutant, which clearly lacked any stem [Fig. 2, F1 and F2 (white arrow)]. Because the minor pilins in *P. aeruginosa* were recently suggested to form a complex that primes pilus assembly (46) and the four minor pseudopilins in the T2SS are thought to prime pseudopilus formation by the major pseudopilin (47, 48), we generated mutants lacking as many as nine of the 10 minor pilins encoded in the *M. xanthus* genome (fig. S5A). Lack of nine of the minor pilins abolished T4P-dependent motility, and T4P did not assemble; however, all 10 core components of the T4PM accumulated at wild-type levels in total cell extracts (fig. S5, B to D). Consistently, when we imaged the mutant deleted for nine of the minor pilin genes, we did not detect T4P (table S1). We did, however, detect empty basal body structures, and they clearly lacked the short stem [Fig. 2, G1 and G2 (white arrow)].

On the basis of these observations, we conclude that the extended stem is made of the major pilin PilA (Fig. 2, F3) and that the short stem is composed of an assembly-priming complex consisting of minor pilins and PilA (Fig. 2, F3 and G3). In the PilA and minor pilin mutants, the structure of the lower periplasmic ring was perturbed and the short stem was absent (Fig. 2, F2 and G2, black arrows). Because it is unlikely for PilA to withdraw its hydrophobic α helix from the IM or pilus fiber to participate in the lower periplasmic ring, the simultaneous changes of the short stem and the lower periplasmic ring suggest a structural/functional linkage between them (see below).

PilC, PilM, PilB, and PilT

The T4PM components PilC, PilM, PilB, and PilT all have folded cytoplasmic domains and are therefore the candidates for the cytoplasmic ring, disc, and dome. Within the set of single-gene knockouts of these four proteins, only empty basal bodies were found on the $\Delta pilC$, $\Delta pilM$, and $\Delta pilB$ cells, as expected (Fig. 2, H1, I1, and J2), and only piliated T4PM basal bodies were found on the $\Delta pilT$ cells (Fig. 2, J1). In these four mutants, all the remaining T4PM proteins accumulated (fig. S2). Because the cytoplasmic ring was missing in the $\Delta pilM$ mutant (Fig. 2, I1) but appeared un-

perturbed in the $\Delta pilC$, $\Delta pilB$, and $\Delta pilT$ mutants, we conclude that the ring is composed of PilM (Fig. 2, I3). Because the ring and dome were retained in the $\Delta pilB$ mutant but not the disc, the disc must be PilB (Fig. 2, J2 and J4). The fact that the cytoplasmic disc had a size corresponding to that of a hexameric secretion or traffic ATPase (37–40) also strengthens this conclusion (see below). The cytoplasmic dome and disc were both missing in the $\Delta pilC$ mutant (Fig. 2, H2), but the cytoplasmic ring was still present (Fig. 2, H1, black arrows), which suggests that the dome is composed of PilC and that the PilB disc does not localize in its absence (Fig. 2, H3). Notably, the short stem structure was also missing in the $\Delta pilC$ structure, revealing that PilC is required to stabilize it.

With the identification of the ring as PilM, the dome as PilC, and the disc as PilB, it follows that (i) the PilM ring assembles in the absence of the PilC dome or PilB disc, (ii) the PilC dome and PilB disc both require the PilM ring, and (iii) the PilC dome assembles without the PilB disc, but the PilB disc requires the PilC dome. These interdependencies match previous observations (fig. S2) except in one regard: Earlier immunofluorescence experiments demonstrated that PilC can be incorporated into the T4PM independently of PilM (21) (fig. S2). Our observation that in the $\Delta pilM$ mutant all cytoplasmic densities and the short stem are missing in averages (Fig. 2, I2) clarifies that the PilM cytoplasmic ring is important for consistent incorporation of the other cytoplasmic proteins and formation of the short stem.

After the cytoplasmic ring, dome, and disc had been assigned to PilM, PilC, and PilB, respectively, no additional cytoplasmic density was available to interpret as PilT. The most likely explanation is that PilT occupies the same location as PilB. When generating subtomogram averages, because we could not tell whether any particular pilus was in the extension or retraction state, we averaged all identified particles. The fact that the $\Delta pilT$ mutant displayed more T4Ps than wild-type cells in our images (table S1) confirmed that the T4Ps were actively extending and retracting in our wild-type sample. Hence, the wild-type piliated structure was an average of both PilB- and PilT-bound states. The $\Delta pilT$ piliated structure (Fig. 2, J1), however, showed no clear addition or loss of cytoplasmic densities relative to the wild type (fig. S1, B and D), which suggests that the differences between PilB- and PilT-bound particles are not visible at this resolution. We conclude that both PilB and PilT form cytoplasmic disc structures (Fig. 2, J4) and bind to the basal body in a mutually exclusive manner.

To test this idea, we generated a $\Delta pilB \Delta pilT$ double mutant. As expected, this mutant lacked T4P-dependent motility and all other T4PM proteins accumulated as in the wild type (fig. S6). When we imaged the $\Delta pilB \Delta pilT$ double mutant (Fig. 2, J3), no clear differences were observed on its empty basal body relative to that of $\Delta pilB$ and wild-type strains (Fig. 2, A2 and J2), confirming our assignment of both PilB and PilT ATPases to the same cytoplasmic disc. The ATPase density was located ~2 nm away from the IM, preventing

any direct interaction between the ATPases and pilin subunits. We therefore conclude that PilC, which forms the cytoplasmic dome structure, lies between the ATPases and the stem and transduces force generated by ATP hydrolysis into pilus extension and retraction. In agreement with this model, the N-terminal cytoplasmic domain of PilC was recently shown to interact directly with PilB, and the C-terminal cytoplasmic domain was suggested to interact with PilT (49).

Placing available component structures into the overall molecular envelope

Because atomic structures are available for homologs of >90% of the domains of the T4PM, we next sought to test the plausibility of our component maps by trying to place these structures into the overall molecular envelope in a way that would satisfy the maps and all known constraints, including domain sizes, structures, and connectivities (see supplementary materials, in particular figs. S7 to S11, and for details, and Movie 1 for a presentation of the process in three dimensions). Remarkably, in this process, the sizes and shapes of all domains analyzed fitted well into the molecular envelopes and resulted in a hypothetical working model of the piliated and nonpiliated basal bodies (fig. S12). Because the 2- to 4-nm resolution of the subtomogram averages was not high enough to reveal the orientation of each component within the EM envelope, we next generated models with most components filtered to 3-nm resolution (Fig. 3, B to E). The process of positioning and connecting the domains in the context of a full hypothetical structural model nonetheless revealed important relationships that rationalize numerous previous observations and suggest insights into T4PM assembly, structure, and function, as described below and shown in Movie 2.

Overall architecture and assembly

All T4PM components except the pilus and PilC appear to form rings. Each of the OM, peptidoglycan (PG), and IM cell envelope layers is negotiated or engaged by a proteinaceous ring, and there is an additional “floating” lower periplasmic ring. Each ring is linked to the rings above and below to create an integrated but flexible structure spanning the entire cell envelope. The subtomogram averages of assembled T4PM subcomplexes in different mutants provide snapshots of the assembly pathway and support the sequence [PilQ, TsaP] → [PilP, PilN, PilO] → [PilM, PilC, PilA, minor pilins] → [PilB, PilT] (fig. S13). The coiled-coil domains of PilN and PilO form a cage-like compartment above and within the IM. Although the exact number of subunit monomers in each ring remains unknown, the 1:1 connectivities between PilQ-PilP, PilP-PilN-PilO, and PilN-PilM suggest that all the rings have the same stoichiometry. We found that 12 copies of the ring components fit best into the EM density, in agreement with observations of secretin channels of T4PM and T2SS in multiple species (41, 42, 50). The entire basal body is therefore robustly anchored to the PG by ~12 TsaP N-terminal LysM domains and ~36 PilQ AMIN domains arranged irregularly (for

instance, as in Fig. 3D). PilQ in turn links to PilP, which links to the PilN-PilO ring, which links to PilM, which binds PilB (and PilT). Unless one of these known connectivities is only transitory in vivo, the only components free to rotate are PilC and potentially the pilus.

Pilus assembly and disassembly by a rotating PilC

The structure of the pilus can be thought of as either a 3-start left-handed helix or a 1-start or 4-start right-handed helix. Because no substantial pilus rotation has been observed during pilus retraction or extension, three different assembly structures or mechanisms can be imagined (18): (i) a fixed structure with three active sites adding pilins at each of the three sites needed to extend a 3-start helix; (ii) a fixed structure with four active sites adding pilins at each of the four sites needed to extend a 4-start helix; or (iii) a rotating structure with one or a few active sites adding pilins one at a time as it rotates around the axis of the fiber, extending the fiber as a 1-start helix (3, 51). As described above, the OM pore complex, alignment complex, and PilM-ATPase complexes are all directly or indirectly linked and anchored to the PG. The only T4PM component able to rotate is PilC. We found that the space inside the PilM-PilN-PilO “cage” can accommodate no more than one PilC dimer. It is unlikely that a PilC dimer would have three or four active sites for interacting with different PilA molecules; therefore, our models point to a 1-start assembly mechanism in which PilC rotates as it assembles the helical pilus fiber. It is also known that *N. gonorrhoeae* T4Ps extend and retract in increments smaller than the length of one helical turn (52), which is most easily explained by a mechanism with more than one step per turn, corresponding to the 1-start assembly mechanism. Another reason to favor models that involve rotation of some component such as PilC is that the homologous archaeal flagellar motor clearly begins to rotate its flagellar filament once this filament has been assembled (53).

In our architectural model of the pilated basal body, the PilC dimer rests on top of (and is presumed to interact with) two opposing PilB subunits in the ATPase hexamer, and the six NTDs of PilB in the ATPase hexamer are clamped in place by interacting with every second PilM subunit in the cytoplasmic ring (fig. S12A). We therefore propose that ATP hydrolysis by the pairs of opposing PilB subunits that contact the PilC dimer causes PilB NTD movements that rotate the asymmetrical PilC dimer. We predict the rotation of PilC to have two consequences: (i) It “scoops” new PilA subunits one at a time out of the membrane and onto the base of the pilus, and (ii) PilC is transferred to the next pair of opposing PilB subunits. The transmembrane segments of PilC likely mate with the tapered tip of the helical pilus fiber in such a way as to extend and anchor the pilus to the basal body, thereby creating a complete binding pocket for the hydrophobic tail of the next PilA subunit to be incorporated. Once a new PilA subunit is incorporated, PilB drives PilC around a fraction of a turn, pushing the pilus up a frac-

tion of the length of a pilin subunit and recreating the binding pocket in the next available position.

One problem with this model is that the hexameric nature of PilB would predict elementary turns of 60° or 120°; neither would result in the exactly 3.6 subunits per turn needed to extend the pilus without any rotation. Perhaps the process does involve some slight rotations or Brownian motions of the proteins that would cause small slips in register. Pilus retraction would be accomplished by a switch from PilB to PilT. Thus, although the exact structure of the PilC dimer and its orientation in the T4PM remain uncertain, the architecture alone implies that both PilB and PilT act as ratchets biasing the assembly or disassembly process: While PilB holds the PilC-tip pocket on the next empty position of the pilus fiber until it is filled, and then quickly rotates to prevent subsequent dissociation, PilT holds the PilC-tip pocket on the last subunit of the tip until it dissociates back into the membrane, after which it quickly rotates to prevent reassociation.

The PilN-PilO and PilM rings likely sense pilus retraction signals and guide ATPase selection

Previous studies have shown that pilus retraction is induced by adding pilin-binding substrates (54) or pulling on the pilus directly (52). Recent studies with antibodies showed that tension induces conformational changes in the pilus itself (55), which could propagate into the basal body. Substrate binding may also induce similar conformational changes. Our results point to a model in which the alignment complex is an IM-crossing transmission module: Pilus retraction signals carried by the pilus itself into the basal body could be sensed in the periplasm by the PilN-PilO ring and then transmitted via the coiled-coil domains through the IM to modulate the conformation of the PilM ring, which in turn governs which ATPase is bound. This model rationalizes the recent report that conformational changes are required in the coiled-coil domains of PilN-PilO during the transition between T4P extension and retraction, which suggests that the alignment complex is not simply a static connector between IM and OM components, but instead plays a critical role in T4P dynamics (56).

The power of combined structural approaches to dissect complicated molecular machines

Solving the structures of large macromolecular machines is challenging. Traditional structural methods such as x-ray crystallography, nuclear magnetic resonance spectroscopy, and single-particle cryo-electron microscopy can deliver near-atomic resolution, but they all rely on purified samples. Many important biological structures such as flagellar motors (57), chemoreceptor arrays (58), and the T4PM studied here, however, may never be purifiable in a native state; we found that even the PilQ (fig. S4), PilC (fig. S10), and PilM-PilN-PilO (fig. S14) subcomplexes lose their native structure when purified. As a result, only structures of isolated subunits and small subcomplexes have

been determined. Cryo-electron tomography can reveal at least the architectures of these large machines in situ, as our work has shown. When there is sufficient additional information about the structures and connectivities of the components, working models can be built that provide a path toward structural understanding, where both the structural relationships and the new mechanistic insights they suggest can then be tested.

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SUPPLEMENTARY MATERIALS

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