Two Distinct Conformations in 34 FliF Subunits Generate Three Different Symmetries within the Flagellar MS-Ring

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ABSTRACT The bacterial flagellum is a protein nanomachine essential for bacterial motility. The flagellar basal body contains several ring structures. The MS-ring is embedded in the cytoplasmic membrane and is formed at the earliest stage of flagellar formation to serve as the base for flagellar assembly as well as a housing for the flagellar protein export gate complex. The MS-ring is formed by FliF, which has two transmembrane helices and a large periplasmic region. A recent electron cryomicroscopy (cryoEM) study of the MS-ring formed by overexpressed FliF revealed a symmetry mismatch between the S-ring and inner part of the M-ring. However, the actual symmetry relation in the native MS-ring and positions of missing domains remain obscure. Here, we show the structure of the M-ring by combining cryoEM and X-ray crystallography. The crystal structure of the N-terminal half of the periplasmic region of FliF showed that it consists of two domains (D1 and D2) resembling PrgK D1/PrgH D2 and PrgK D2/PrgH D3 of the injectisome. CryoEM analysis revealed that the inner part of the M-ring shows a gear wheel-like density with the inner ring of C23 symmetry surrounded by cogs with C11 symmetry, to which 34 copies of FliFD1–D2 fitted well. We propose that FliFD1–D2 adopts two distinct orientations in the M-ring relative to the rest of FliF, with 23 chains forming the wheel and 11 chains forming the cogs, and the 34 chains come together to form the S-ring with C34 symmetry for multiple functions of the MS-ring.

IMPORTANCE The bacterial flagellum is a motility organelle formed by tens of thousands of protein molecules. At the earliest stage of flagellar assembly, a transmembrane protein, FliF, forms the MS-ring in the cytoplasmic membrane as the base for flagellar assembly. Here, we solved the crystal structure of a FliF fragment. Electron cryomicroscopy (cryoEM) structural analysis of the MS-ring showed that the M-ring and S-ring have different rotational symmetries. By docking the crystal structure of the FliF fragment into the cryoEM density map of the entire MS-ring, we built a model of the whole periplasmic region of FliF and proposed that FliF adopts two distinct conformations to generate three distinct C11, C23, and C34 symmetries within the MS-ring for its multiple functions.

KEYWORDS bacterial flagellar motor, rotor, MS-ring, type III secretion

The bacterial flagellum is a filamentous organelle for locomotion in many bacterial species. The flagellar filament is rotated by a motor embedded in the cell membrane and functions as a screw to thrust the cell. The motor consists of the rotor and several stator units (1, 2). The stator unit is a transmembrane complex of the MotA and...
MotB family proteins. The rotor is composed of the MS-ring and the C-ring (Fig. 1A). The MS-ring is a transmembrane ring assembly of a single protein, FliF, and has a two-tier ring structure; the S-ring is an upper ring located in the periplasm, and the M-ring is a lower ring embedded in the inner membrane (3). FliF is a 60-kDa protein with a large periplasmic region between two transmembrane helices (Fig. 1B) (4). The C-ring is a cytoplasmic cup-like structure composed of FliG, FliM, and FliN proteins. The torque is generated by the interaction between FliG and MotA and is transmitted from the rotor to the flagellar filament through a drive shaft called the rod, followed by a universal joint called the hook (Fig. 1A).

The MS-ring provides a base for the assembly of the flagellar structure. The C-ring is attached to the cytoplasmic surface of the MS-ring through the interaction between FliG and FliF (5–7). The rod construction begins in the central hole of the MS-ring at its periplasmic side, and thus, the proximal end of the rod is inserted in the MS-ring. The filamentous part of the flagellum, including the rod, hook, and filament, is termed the flagellar axial structure. The component proteins of the axial structure are translocated across the inner membrane via the flagellar protein export apparatus, which is a member of the type III secretion (T3S) family (8, 9). The MS-ring holds a transmembrane region of the export apparatus in its central hole and therefore serves as housing for the export apparatus (Fig. 1A). The flagellar formation is thought to be initiated by the assembly of the type III export gate complex composed of FlhA, FlhB, FliP, FliQ, and FliR with the help of the FliO scaffold (10), followed by the recruitment of FliF through an interaction between FliF and FlhA (11) to form the MS-ring around the export gate complex (10). The export gate component proteins form partial gate complexes but cannot assemble into the complete export gate complex without FliF, indicating that the MS-ring stabilizes the export gate complex (10, 12).

The proteins of the flagellar basal body structure are homologous to the component proteins of the T3S injectisome, a needle-like organelle of pathogenic bacteria that delivers effector proteins into host cells for infection and pathogenicity (13). The flagellar basal body proteins also show similarities to the component proteins of the sporulation-essential channel (14, 15), which connects the mother cell and the forespore in Bacillus subtilis. FliF shares homology with the inner membrane ring components of the T3S injectisome SctJ (also named as EscJ in enteropathogenic Escherichia coli [EPEC] and PrgK in Salmonella SPI-1) and SctD (EscD in EPEC and PrgH in SPI-1) and the Bacillus sporulation channel components SpoIIAH and SpoIIAG (14–18).
sequence analysis revealed that the periplasmic region of FliF has three ring-building motifs (RBMs) (15), namely, RBM1, RBM2, and RBM3 (Fig. 1B), which are conserved in SctJ/SctD, SctJ/SctD/SpoIIIAH, and SpoIIIAG, respectively. X-ray crystallography and electron cryomicroscopy (cryoEM) analyses of the Salmonella SPI-I injectisome have revealed that PrgK and PrgH form a concentric double ring with 24-fold rotational symmetry in the periplasmic region of the basal body (18–21). The crystal structures of SpoIIAF and SpoIIAH have shown a structural similarity to PrgK (16, 17, 22). The high-resolution cryoEM structure of SpoIIIAG has revealed that SpoIIIAG forms a 30-fold symmetrical ring with a unique cylindrical β-barrel structure (23).

Recently, partial structures of the periplasmic region of the MS-ring have been determined at 2.6- to 3.3-Å resolution by cryoEM image analysis of purified MS-rings formed by recombinant FliF with some C-terminal truncations. (24). These structures showed a variation in their subunit stoichiometry and revealed that the S-ring shows C32 to C35 symmetry and consists of the C-terminal half of the periplasmic region of FliF, including RBM3. The S-ring comprises a globular domain forming a flat ring and an extended chain, including long antiparallel β-strands forming a cylindrical collar above the ring. The overall S-ring structure is similar to that of SpoIIIAG, albeit their symmetries are different. In addition, 21 or 22 copies of RBM2 form a ring in the inner part of the M-ring, and this is surrounded by 9 or 10 globular densities composed of RBM1 and RBM2. However, a few copies of RBM2 and more than 20 copies of RBM1 are missing in these structures, and the variation in the subunit stoichiometry is likely to be an artifact due to C-terminal truncations of FliF because the MS-ring in the flagellar basal body, as well as that formed by full-length recombinant FliF, showed only 34-fold rotational symmetry (25). Moreover, the cryoEM analysis of the purified flagellar basal body showed a C23 symmetry at the inner part of the M-ring (25). Therefore, the roles and functions of the RBM domains in the MS-ring formation and entire flagellar assembly remain obscure.

Here, we report the crystal structure of Aa-FliF58–213, which corresponds to the N-terminal half of the periplasmic region of FliF from Aquifex aeolicus, at 2.3-Å resolution. FliF58–213 is composed of two domains (D1 and D2), and they show structural similarity to the corresponding domains of SctJ, SctD, and SpoIIIAG. We constructed an atomic model for the inner part of the M-ring by combining the crystal structure with low-resolution cryoEM maps of the MS-ring with the help of structural similarity to the homologous injectisome proteins. The model indicates that FliF subunits adopt two distinct conformations in the M-ring structure to generate multiple symmetries within the ring. We also built a structural model of the entire periplasmic region of FliF by combining with the S-ring model determined by high-resolution cryoEM analysis (25). These results provide the structural basis of the flagellar assembly mechanism and the evolutionary relation to the T3S injectisome and sporulation channel.

RESULTS

CryoEM analysis of the Salmonella MS-ring complex. We expressed and purified the Salmonella MS-ring with Flig for single-particle cryoEM image analysis (see Fig. S1 in the supplemental material). Reconstruction of the MS-Flig ring complex without imposing any symmetries yielded an 8.6-Å resolution map (Fig. 1C and D). The map showed that the purified MS-ring consists of the following five regions: (i) the S-ring, (ii) the inner ring of the M-ring, (iii) the middle region of the M-ring with C11 symmetry, (iv) the outermost region of the M-ring, and (v) the innermost plug region of the M-ring (Fig. 1C and D, Fig. 2A). The maps of the outermost region and the innermost plug region of the M-ring show a blurred density, which may include the transmembrane region of FliF with detergent molecules. The overall view of the map is similar to that of the MS-ring map reported recently (24), although the rotational symmetry of each region is different. Unfortunately, the density corresponding to the Flig ring was invisible.

Crystal structure of a FliF fragment corresponding to RBM1 and RBM2. Partial atomic models of the MS-ring corresponding to the S-ring (RBM3) and the inner part of
the M-ring (RBM2inner) have already been determined based on high-resolution cryoEM maps (24). The middle part of the M-ring (RBM2outer) has also been modeled using medium-resolution cryoEM maps (24). However, RBM1 and a few copies of RBM2 remain unknown and are required for understanding their roles in the M-ring function. To build the atomic model of RBM1 and RBM2, we determined the crystal structure of a FliF fragment containing these two motifs. We initially tried to crystallize various fragments of *Salmonella* FliF, but no crystal was obtained. Therefore, we prepared FliF fragments from a thermophilic bacterium, *A. aeolicus*, and determined the crystal structure of a fragment consisting of residues 58 to 213 (Aa-FliF58–213) at 2.3-Å resolution (Fig. 2B; see Fig. S2A to E in the supplemental material; see Table S1 in the supplemental material). The crystal belongs to the space group of \( \text{H}3 \) and contains two essentially identical molecules in an asymmetric unit (Fig. S2E) (the root mean square deviation for \( \text{C}_\alpha \)).

**FIG 2** Structure of RBM1 and RBM2 of FliF. (A) The cryoEM surface map of the inner ring and the middle region of the M-ring with the central blurred density. The top view, the side view, and the vertical section are shown in the left, the middle, and the right, respectively. The scale bar is 50 nm. (B) \( \text{Ca} \) ribbon drawing of the crystal structure of Aa-FliF58–213. The model is color coded from blue to red from the N to the C terminus. The structure consists of two domains, namely, D1 (residues 58 to 122) and D2 (residues 123 to 213). (C to F) Comparison of the domain structures of Aa-FliF with those of PrgH and PrgK. (G) The structure model of the inner ring (cyan) and the middle region (magenta) of the M-ring. The top view, the side view, and the vertical section are shown in the left, the middle, and the right, respectively. (H) Superimposition of the D1 to D2 structure (pink) on the cryoEM map of the middle region of the M-ring after applying the 11-fold symmetry averaging (gray mesh). (I) 8.6-Å density maps of the middle region calculated from the D1 to D2 structure model.
The structure of Aa-FliF58-213 consists of two distinct domains corresponding to RBM1 (D1; Pro58 to Ser122) and RBM2 (D2; Arg123 to Asp213) (Fig. 1B, Fig. 2B). The D1 domain (FliFD1) is a compact globular domain composed of two α-helices (α1 and α2) and five β-strands (β1 to β5). β1, β2, and β3 form a core β sheet flanked by the two α-helices and another β-sheet made up of β4 and β5. The D2 domain (FliFD2) is more elongated and consists of two α-helices (α3 and α4) and four β-strands (β6 to β8 and β9). An intramolecular disulfide bridge is formed between C147 and C182 at a pole of FliF2, but it is not essential for the FliF function, as discussed later. Aa-FliF58-213 forms a dimer related by a pseudo-2-fold symmetry in the crystal, and β4 and β5 form an intersubunit β sheet with β1 of the dimer mate (Fig. S2E and F). However, the dimer is a crystal-packing artifact because FliF forms a ring structure, and the 2-fold symmetrical dimer cannot be fitted in the cryoEM density, as shown later. The overall structure of FliF2 is similar to the recently reported cryoEM structures of RBM2inner and RBM2outer (24) (Fig. S2G).

FliFD1 and FliFD2 structurally resemble the components of the T3S injectisome.

Amino acid sequence analysis showed that FliF has structural motifs conserved in SctD (PrgK in SPI-1) and SctJ (PrgH in SPI-1) of the T3S injectisome (15). The periplasmic region of PrgK is composed of two domains, namely, D1 (PrgKD1) and D2 (PrgKD2), and that of PrgH is composed of three domains, namely, D2 (PrgHD2), D3 (PrgHD3), and D4 (PrgHD4) (see Fig. S3A and B in the supplemental material). They form a concentric double ring with C24 symmetry (the inner PrgK ring and the outer PrgH ring) (Fig. S3C). We found that FliF2 shows structural similarity to both PrgKD1 and PrgHD2 (Fig. 2C and D, Fig. S3D), although the N-terminal α-helix of FliFD1 (α1) is longer than the corresponding helix of PrgH2. FliFD2 resembles both PrgKD2 and PrgHD3, which adopt a common RBM fold (Fig. 2E and F, Fig. S3D). These structural similarities suggest that the D1-D2 region of FliF (FliF2) also forms a concentric double-ring structure in a way similar to the PrgK-PrgH ring of the T3S injectisome.

Structure of the inner ring and the middle region of the M-ring. Aa-FliF58-213 shows an amino acid sequence identity of 44% with the corresponding region of Salmonella FliF (St-FliF60-213) without any insertions and deletions (see Fig. S4A in the supplemental material). Thus, we made a homology model of St-FliF60-213 based on the structure of Aa-FliF58-213 and used this model to construct the M-ring structure. We first built a tentative FliF ring model by superimposing FliFD1 and FliFD2 to the PrgK and PrgH rings independently (Fig. S3D to F). The tentative ring model contains a total of 48 FliFD1-D2 subunits, as follows: 24 subunits form the inner ring (fitted to the PrgK ring), and the remaining 24 subunits form the surrounding ring (fitted to the PrgH ring). The inner and outer diameters of the inner ring of the 24-mer model are comparable to those of the cryoEM density for the inner M-ring with C23 symmetry in the native Salmonella basal body (25), suggesting that the tentative 24-mer model has a similar subunit arrangement to the inner M-ring. Therefore, we produced a 23-mer model by applying a C23 symmetry operation to a subunit of the 24-mer inner ring model and then fit it into the cryoEM density with C23 symmetry (EMD-30613) (25) (Fig. S3G). The domain arrangement is similar to those in the recently reported RBM2inner ring models (24), although their symmetries are different, namely, C21 and C22. To further examine the correctness of the subunit arrangement in the inner ring, we replaced some residues of FliF at the subunit interface with cysteine and examined oligomerization by SDS-PAGE, followed by immunoblotting (see Fig. S5 in the supplemental material). The H156C/S200C mutant produced higher-order oligomers by intermolecular disulfide bonds, and this result supports the subunit arrangement in the inner ring model because H156 and S200 are close enough to form a disulfide bond in the ring model.

The surrounding ring subunits of the tentative ring model overlapped with the cryoEM density of the middle region of the M-ring, but its symmetry is C11. Thus, we omitted 13 protomers and rearranged the remaining 11 protomers of FliFD1-D2 in the tentative surrounding ring by applying C11 symmetry (Fig. S3G). Then, we modified the orientation of each domain by fitting it into the cryoEM density with 11-fold
that the structures of mera protein can form the MS-ring albeit not at the dots were less than those of the FliF chimera labeled with GFP, although the number and intensity of the cell surface, suggesting that expressed expressing FliF cent protein (GFP) to its C terminus (Fig. S4E and F), indicating that the disul bridge between C147 and C182 in the D2 domain of mutant (Fig. 3F). These results indicate that the i-loop is important for the FlhA assem-

Functional analysis of a FliF chimera. To obtain functional evidence that supports our ring model based on the Aa-FliF58-213 structure, we constructed a FliF chimera protein with residues 58 to 214 of Salmonella FliF (St-FliF) replaced by residues 56 to 212 of A. aeolicus FliF (Aa-FliF) (Fig. S4B) and expressed it in Salmonella ΔfliF cells. The deletion of fli completely abolished motility in soft agar (Fig. S4C) because the flagellum was not produced. Expression of Aa-FliF did not complement the motility defect of the Salmonella ΔfliF mutant, whereas the expression of the FliF chimera restored the motility to a signifi-

The “i-loop” in FliF is required for assembly of the flagellar export gate complex. In our M-ring model, a characteristic protruding loop connecting β6 and β7 and containing βi (residues 157 to 170 in Aa-FliF and 159 to 172 in St-FliF) (Fig. 3A), termed the i-loop, surrounds the hole of the inner ring (Fig. 3B). To investigate the role of the i-loop in flagellar formation, we deleted residues 161 to 170 of St-FliF (FliFΔi-loop) to see the phenotype. This deletion completely abolished cell motility and flagellation (Fig. 3C and D). In contrast, fluorescent dots were found in Salmonella ΔfliF cells expressing FliFΔi-loop labeled with GFP at a similar level to that of the cells expressing wild-type St-FliF-GFP (Fig. 3E). The M-ring is essential for the assembly of the flagellar protein export gate complex composed of FlhA, FlhB, FliP, FliQ, and FliR (11). FlhA forms the homo-nommer in the complex, and the deletion of fli completely abolishes the assembly of FlhA (11). Because the i-loop is adjacent to A174 and S175 of FliF, whose deletions are partially suppressed by extragenic suppressor mutations in the N-terminal transmembrane domain of FlhA (see Fig. S6 in the supplemental material), we analyzed the effect of i-loop deletion on the assembly of the FlhA ring. FlhA labeled with yellow fluorescent protein (YFP) formed fluorescent dots in the cell expressing wild-type FliF, and the analysis of the fluorescence intensity revealed that FlhA forms nonamer (12) (Fig. 3F), but no clear fluorescent dot was observed in the i-loop deletion mutant (Fig. 3F). These results indicate that the i-loop is important for the FlhA assembly within the MS-ring.

Structure of the periplasmic region of the MS-ring. We integrated the inner and middle region of the M-ring model with the C34 S-ring model (PDB ID: 6SD4) and constructed the periplasmic structure model of the MS-ring (Fig. 4A to G). The MS-ring is composed of the 34 FliF subunits in the native Salmonella basal body. FliFΔ1-2 adopts two distinct structures in the M-ring. The inner M-ring is formed by 23 copies of FliFΔ1-2 (Fig. 4A) and is surrounded by 11 copies of FliFΔ1-2 in a distinct orientation from those in the inner ring (Fig. 4B). These chains form the two conformationally distinct FliF subunit groups, forming the inner and middle region of the M-ring with 23- and 11-fold symmetry, respectively, and are joined to form the S-ring and collar with
34-fold symmetry (Fig. 4C to H). The distance between the C terminus of the S-ring and the density of the outermost region of the M-ring are about 30 Å. Therefore, the 13 residues following the C terminus of the S-ring to the N terminus of FliF TM2 should be adopting an extended conformation.

**DISCUSSION**

The crystal structure of RBM1 and RBM2 together with the cryoEM structure of the MS-ring, including a high-resolution structure of the S-ring (RBM3), revealed the periplasmic part of the MS-ring structure. Our cryoEM analysis of the MS-FliG ring complex demonstrated that the MS-ring in the basal body is made up of 34 FliF subunits and is consistent with results of the recent cryoEM studies of the MS-ring formed by full-length FliF and the MS-ring in the flagellar basal body (25). The S-ring with a cylindrical collar is shaped like a boater hat with C34 symmetry, whereas the inner part of the M-ring has a gear wheel-like structure consisting of the core C23 symmetry inner ring and the C11 symmetry cogs surrounding it. Thus, the three RBMs of FliF subunits adopt two distinct arrangements in the M-ring. FliF has two transmembrane helices and a periplasmic region (FliFP) between them. FliFP contains three globular RBM domains, namely, FliFD1, FliFD2, and FliFD3, and a long extended up-and-down structure (residues 268 to 382) including two antiparallel β-strands (Fig. 4H; see Fig. S7A in the supplemental material). The N-terminal half of FliFP containing FliFD1 and FliFD2 adopts the
following two distinct domain arrangements in the M-ring: the inner 23 copies form the inner ring and the remaining 11 copies surround the inner ring. The arrangement of FliFD2 in the inner ring resembles that of the D2 domain of the SctJ family proteins, such as PrgK and EscJ (Fig. S7C and D). The C-terminal half of FliFP containing FliFD3 and the extended structure forms the S-ring and collar. The structure of FliFD3 has a common RBM fold and resembles FliFD2 (Fig. S7A and C). FliFD3 forms a large ring structure with a similar domain arrangement to the inner ring as well as to the SctJ family protein rings (Fig. S7A and D). The long antiparallel $\beta$-strands in the extended structure are vertically lined up to form a cylindrical collar with a 68-stranded $\beta$-barrel structure. The overall structure of the S-ring resembles the SpoIIIAG structure (Fig. S7A and B), although the number of the subunits and therefore the symmetry are different.

Recently, Johnson et al. have solved multiple cryoEM structures of the MS-ring with a variation in subunit stoichiometry from 32 to 35 and found symmetry mismatch...
between the S-ring and M-ring (24). They mainly described the structure of the MS-ring with C33 symmetry, which our previous work suggests is not the native MS-ring structure in the basal body (25). But even in their model of the MS-ring formed by 34 copies of FliF, the inner M-ring is composed of 22 copies of RBM2 and another set of 10 copies composed of both RBM1 and RMB2 is surrounding it, and therefore, two copies of RBM2 are missing. Although the orientation of each domain in their model is similar to that of our model, the subunit stoichiometries and their azimuthal positions in the inner and middle region of the M-ring are different. Our model contains a complete set of RBM2 with 34 copies, as follows: 23 copies form the inner ring, and 11 copies form the outer surrounding densities of the middle region. Since we built our model based on the cryoEM map of the basal body and the MS-ring formed by 34 copies of full-length FliF, we believe that our model reflects the MS-ring structure of the native flagellum. The MS-FliG ring complex used for our cryoEM analysis of the MS-ring was prepared from the cells expressing not only FliF and FliG but also the export apparatus proteins. Therefore, the export apparatus proteins may be needed for FliF to form the 23-plus 11-subunit structure in the inner part of the M-ring even though they were dissociated from the MS-ring during purification.

The 8.6-Å cryoEM map of the MS-FliG ring complex showed blurred densities in the outer M-ring located below the periphery of the S-ring (Fig. 1C, colored yellow). This density is thought to be formed by the TM2 and C-terminal cytoplasmic region of FliF because the density disappears by truncation of these regions (4). The map also showed another unassigned blurred density below the central hole of the inner M-ring (Fig. 1C, colored gray) where the FlhA nonamer ring is supposed to be located in the basal body, as observed by electron cryotomography (26). However, the MS-FliG ring used to produce the map did not contain any export gate component proteins because they fell off during the purification process. The blurred density is close to the N terminus of FliF(A), of the inner M ring model, and the density of FliF(A) itself is also relatively poor in the map. Therefore, the N-terminal region of FliF, including TM1 and FliF(A) of the inner M-ring, may form a disordered aggregate in the purified MS-FliG ring. The export gate complex might be needed for the proper folding and arrangement of this region.

The flagellar protein export gate complex, which comprises FlhA, FlhB, and the FliP/Q/R complex, is expected to be accommodated in the central hole of the MS-ring (11). The FliP/Q/R complex is needed for the assembly of FlhA into the export gate complex (11). Deletion of the i-loop (residues 161 to 170) surrounding the hole of the inner ring affected the FlhA assembly but not the MS-ring formation (Fig. 3), suggesting that the i-loop interacts with the export gate complex. Recent cryoEM analysis of the injectisome proposed that the SpaP/Q/R complex, which corresponds to the flagellar FliP/Q/R complex, is stuck in the central hole of the SctJ ring, which corresponds to the inner ring of the M-ring (27–29). The SctJ subunits surround the middle of the bullet-shaped P/Q/R complex. In addition, cross-linking experiments have revealed that SctJ directly interacts with the P and R subunits of the complex (27). The inner diameter of the M-ring is comparable to that of the FliP/Q/R complex (27). Therefore, the i-loop of the inner M-ring may directly interact with the FliP/Q/R complex and accommodate it in the central hole. Thus, the FliP/Q/R complex may not be properly held in the inner M-ring without the i-loop, resulting in the collapse of the export gate complex. The inner M-ring shows a 23-fold rotational symmetry, whereas the FliP/Q/R complex is a helical assembly of five FliP, four FliQ, and one FliR molecules. The flexible i-loop would be needed to adjust the inner M-ring interface to the FliP/Q/R complex over the symmetry mismatch.

A conserved A-S(A)-V(I)-X-V(L/I) motif in RBM2 of FliF (ASVFL in residues 172 to 176 of Aa-FliF and ASVTV in residues 174 to 178 of St-FliF) is believed to be involved in the interaction with FlhA because the nonflagellation phenotype caused by deletion of residues A174 and S175 in Salmonella FliF was suppressed by mutations in and around the transmembrane region of FlhA (Fig. S6C) (30). The A-S(A)-V(I)-X-V(L/I) motif is in
strand β7 located at the interface to the neighboring subunit in the inner M-ring and is far from the inner membrane in the MS-ring model (Fig. 3A and B), even if the membrane is distorted around the basal body, as depicted in the model of the injectisome basal body (27). Therefore, A174 and S175 do not seem to directly interact with FlhA. The deletion of these two residues may change the size or shape of the central hole of the inner M-ring to cause trouble in accommodating the export gate complex. The suppressor mutations in FlhA may change the conformation of the export gate complex to be fitted into the central hole.

The structure of FlIFD2 resembles that of FlIFD3, PrgKD2, PrgHD3, SpoIIAF, and SpoIIAH and the globular domain of SpoIIAG (Fig. S7). They share an αβαβαβ motif known as a common RBM. They all form ring structures of different sizes by different numbers of subunits. FlIFD2 forms a 23-mer ring with an external diameter of about 16 nm. The external diameters of the PrgK 24-mer ring and the FlIFD3 34-mer ring are about 17 and 24 nm, respectively. They all share a similar domain orientation and similar neighboring subunit interaction surfaces to form the ring. The RBM domain of SpoIIAG forms a 30-mer ring with an external diameter of about 21 nm. The PrgH24 ring is composed of 24 subunits, and its external diameter is about 24 nm. Although the domain orientations of these two molecules in their rings are slightly different from that of the FlIFD3 ring, they also use common interfaces for the neighboring subunit interaction in the ring.

Our structural study has revealed the architecture of the periplasmic region of the MS-ring in the native basal body structure. However, the structure and symmetry of the transmembrane and cytoplasmic regions of the MS-ring remain unclear. The entire structure of the MS-ring is still required to fully understand the mechanism of flagellar assembly and the rotor function.

MATERIALS AND METHODS

Bacterial strains and plasmids. The bacterial strains and plasmids used in this study are listed in Table S2 in the supplemental material. The expression plasmids were constructed by PCR using DNA primers listed in Table S3 in the supplemental material. We cloned the full-length and various fragments of the flif gene from A. aeolicus (Fig. S2A). The FlIF fragments were designed based on the motif and the secondary structure prediction using PSIPRED (http://bioinf.cs.ucl.ac.uk/psipred/). L121M/L195M mutation in pNT32 and FlIFD161-170 mutation in pNT63 and pNT64 were introduced by the QuickChange site-directed mutagenesis (Agilent, CA, USA). The transformation of E. coli and Salmonella was performed by the heat shock method.

Expression and purification of the Salmonella MS-FlIF ring complex. A total of 15 ml of the over-expansion culture of Salmonella SJW1368 (ΔcheW-flhDII) cells harboring pMKM20001 (pTrc99CES/FlhB + FlhA + FlhC + FlhD + FlhF + FlhG + FlhH + G100 + Flk + HA-FLAG-His + FliE + Flg) was inoculated into 1.5 liters of fresh 2 × YT medium (1.6% [wt/vol] Bacto-tryptone, 1.0% [wt/vol] Bacto-yeast extract, and 0.5% [wt/vol] NaCl) containing 100 μg ml⁻¹ ampicillin, and cells were grown at 30°C until the density reached an optical density at 600 nm (OD600) of about 0.6. After 30 min of incubation at 4°C, the cells were grown at 16°C for 12 hours. Then, arabinose was added at a final concentration of 0.2% and incubated at 16°C for another 4 hours. Cells were harvested by centrifugation (6,400 × g, 10 min, and 4°C) and stored at −80°C. The cells were thawed, resuspended in 55 ml 50 mM Tris-HCl (pH 8.0), 50 mM NaCl, and 5 mM EDTA and disrupted by passage through a French pressure cell (FA-032; Central Scientific Commerce). The cell lysates were centrifuged (20,000 × g, 15 min, and 4°C) to remove undisrupted cells. The supernatants were ultra centrifuged (90,000 × g, 1 h, and 4°C). The harvested membranes were suspended in 40 ml of 50 mM N-cyclohexyl-3-aminopropanesulfonic acid (CAPS)-NaOH (pH 11.0), 50 mM NaCl, 5 mM EDTA, and 0.5% β-D-maltoside (DDM), followed by centrifugation (20,000 × g, 20 min, and 4°C), and finally ultracentrifugation (90,000 × g, 60 min, and 4°C). Pellets were resuspended in 25 mM Tris-HCl (pH 8.0), 50 mM NaCl, 1 mM EDTA, and 0.1% DDM and incubated at 4°C for 1 hour. The solution was loaded at a 15% to 40% (v/v/wt) sucrose density gradient in 25 mM Tris-HCl (pH 8.0), 50 mM NaCl, 1 mM EDTA, and 0.1% DDM. After ultracentrifugation (49,100 × g, 13 h, and 4°C), fractions containing FlIF and FliG were collected; diluted with 7 volumes of 25 mM Tris-HCl (pH 8.0), 50 mM NaCl, 1 mM EDTA, and 0.1% DDM; and ultracentrifuged (90,000 × g, 60 min, and 4°C). Pellets were resuspended in 30 μl of 25 mM Tris-HCl (pH 8.0), 50 mM NaCl, 1 mM EDTA, and 0.1% DDM.

Sample vitrification and cryoEM data acquisition. Copper 200 mesh R0.6/1.0 holey carbon grids (Quantifoil) were glow discharged on a glass slide for 30 s. A 2.6-μl aliquot of the sample solution was applied to the grid and blotted by filter paper for 7 s at 100% humidity and 4°C. The grid was frozen by rapid plunging into liquid ethane using a Vitrobot Mark III (Thermo Fisher Scientific, MA, USA). The grids were observed by a Titan Krios FEG transmission electron microscope (Thermo Fisher Scientific) operated at 300 kV with the cryospecimen stage cooled with liquid nitrogen. CryoEM images were recorded with a Falcon II 4k by 4k CMOS direct electron detector (Thermo Fisher Scientific) at a nominal

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Two Distinct Conformations of the MS-Ring Protein FliF

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magnification of ∼75,000 (image pixel size, 1.07 Å) using the EPU software package. The movie images were collected under a defocus range between 1.0 and 3.0 μm with an exposure time of 2 s at a dose rate of 45 e−·pix−1·s−1 (total accumulated exposure, 90 e−·Å−2). Each movie image was fractionated into 7 frames.

Image processing of cryoEM data. The movie frames were subsequently aligned to compensate for beam-induced motion using MotionCor2 (31), and the parameters for the contrast transfer function (CTF) were estimated using Gctf (32).

A total of 461,944 particle images of the MS-FliG ring complex were automatically picked from 6,961 micrographs using Gautomatch (https://www2.mrc-lmb.cam.ac.uk/research/locally-developed-software/zheng-software/kauto), and two-dimensional (2D) and three-dimensional (3D) classifications were performed using RELION-2.1 (33) or 3.0 (34).

Particles from good 2D classes were used for making the initial 3D model of the MS-FliG ring complex using cryoSPARC (35). A total of 53,522 particles from the best 3D class were subjected to 3D refinement, which produced a reconstruction with a resolution of 8.6 Å and a B-factor of −578 Å2 for the C1 symmetry model and with a resolution of 6.1 Å and a B-factor of −312 Å2 for the C11 symmetry model. We did not perform focused classification because the map resolution was not high and the total number of the particle images was not enough.

Expression and purification of Δe-FliF variants. E. coli BL21-CodonPlus(DE3)-RIPL carrying pNT30 to pNT37 was cultured in LB broth containing 50 μg ml−1 ampicillin at 37°C to an optical density at 660 nm of 0.6 to 0.8 and cooled on ice for about 30 min. A total of 0.5 mM isopropyl-β-D-thiogalactopyranoside (IPTG) was subsequently added to the culture, and the culture was prolonged for about 20 h at 16°C. Cells were collected by centrifugation (6,700 g) and suspended in Tris-Sodium Chloride (TN) buffer (50 mM Tris-HCl [pH 8.0] and 200 mM NaCl) containing cOmplete, EDTA-free reagent (Roche) and lysozyme (Wako, Japan). The cells were then disrupted by sonication and centrifuged at 20,000 × g for 10 min to remove cell debris. The supernatant was ultracentrifuged at 100,000 × g for 30 min. All the FliF fragments were successfully expressed (Fig. S2B), and FliF58-272,347-396 fragments were highly soluble (Fig. S2B). The soluble fraction was mixed with Ni-nitrilotriacetic acid (NTA) agarose (Qiagen, Germany) and then incubated on ice for 30 min with gentle mixing. The protein-bound agarose was washed with TN buffer containing 50 mM imidazole, and the proteins were subsequently eluted with TN buffer containing 400 mM imidazole. The protein was concentrated using an Amicon Ultra 10 K device (Merck Millipore, Germany), loaded on a size exclusion column (Superdex 75 10/300 GL; GE Healthcare, UK), and suspended with TN buffer. The peak fraction was collected and concentrated using an Amicon Ultra 10 K device. The expression and purity of the proteins were examined by SDS-PAGE.

The selenomethionine (SeMet) derivative FliF fragment was prepared from E. coli BL21-CodonPlus (DE3) RILX carrying pNT32B. The cells were cultured in SeMet minimal medium (0.1% [wt/vol] NH4Cl, 0.3% [wt/vol] Na2HPO4, 0.3% [wt/vol] NaH2PO4, 2% [wt/vol] glucose, 0.03% [wt/vol] MgSO4, 0.001% [wt/vol] Fe2(SO4)2, 0.001% [wt/vol] thiamine, and 0.005% [wt/vol] seleno-L-methionine). The proteins were purified in the same way as native FliF fragments.

Crystallization and X-ray data collection. Crystallization was carried out using the sitting-drop vapor-diffusion method. Crystallization drops were prepared by mixing 0.5 μl of about 10 to 30 mg ml−1 His-FliF58-213 with 0.5 μl of the reservoir solution. Initial screening was carried out using the screening kits Wizard classic I and II, cryo I and II (Emerald BioSystems, WA, USA), and crystal screen I and II (Hampton Research, CA, USA); and then the conditions were optimized. Crystals appeared within a week. Because proteins rapidly agglutinate at room temperature, we performed crystallization in a cold room (4°C). The best crystals were grown from the drop prepared by mixing 0.5 μl of 30 mg ml−1 protein solution with 0.5 μl of reservoir solution containing 100 mM Na/K phosphate (pH 6.2) and 2.5 M NaCl.

The X-ray diffraction data were collected at synchrotron beamlines BL41XU and BL2681 in SPring-8 (Harima, Japan) with the approval of the Japan Synchrotron Radiation Research Institute (JASRI) (proposal no. 2016A/B2541 and 2017A/B2588). The crystals were directly transferred into liquid nitrogen for cryoprotection no. 2016A/B2541 and 2017A/B2588). The crystals were directly transferred into liquid nitrogen for freezing. The diffraction data were collected under nitrogen gas flow at 100 K. The diffraction data were processed with MOSFLM (36) and scaled with AIMLESS (37). The diffraction data statistics are summarized in Table S1. The experimental phase was calculated using the SAD data of the selenomethionine derivative with the program PHENIX (38). The atomic model was built with Coot (39) and refined to 2.3 Å resolution with PHENIX (38) against the native crystal data of His-FliF58-213. The refinement statistics are summarized in Table S1.

In vivo disulfide cross-linking. Salmonella SJW1684 cells carrying pTH201 with/without mutations were cultured in LB broth containing 50 mg ml−1 ampicillin and 100 μM IPTG at 30°C, and a constant number of cells were collected by centrifugation (13,000 × g) and suspended with nonreducing SDS loading buffer, heated at 95°C for 5 min, and subjected to SDS-PAGE followed by immunoblotting using a polyclonal anti-FliF antibody.

Soft-agar plate assay for motility. A total of 2 μl of an overnight culture was spotted onto Terrific Broth (TB) soft-agar plates (1% [wt/vol] tryptone, 0.5% [wt/vol] NaCl, 0.25% [wt/vol] agar). The plates were incubated at 30°C for the appropriate time as described in the figure legends. The assay was performed at least three times to confirm the reproducibility of the results.

Observation of subcellular localization of FliF-GFP and FliA-YFP with fluorescence microscopy. The overnight cultures grown in LB broth were inoculated at a 100-fold dilution into TG broth (1% [wt/vol] tryptone, 0.5% [wt/vol] NaCl, and 1% [wt/vol] glycerol) containing 100 μM IPTG and cultured for 4 h at 30°C. Cells were harvested by centrifugation and resuspended in motility medium (10 mM potassium phosphate [pH 7], 0.1 mM EDTA, and 85 mM NaCl). Then, the cells were loaded between a coverslip and
a slide glass and incubated for 10 min to be attached to the coverslip surface. Unbound cells were washed away by the motility medium. The cells were observed by a fluorescence microscope (BX53 [Olympus, Japan] for GFP and BX50 [Olympus] for YFP) equipped with a 100-W high-pressure mercury lamp. Images were recorded using a digital charge-coupled-device (CCD) camera (Infinity2-1RM and Infinity Capture [Argo Corporation, Japan] for GFP and ORCA-Flash4.0 and High-speed Recording Software version 1.7.1.0 [Hamamatsu Photonics, Japan] for YFP).

**Data availability.** The atomic coordinate has been deposited in the Protein Data Bank (www.pdb.org) under accession code 7CIK. The cryoEM maps have been deposited in the Electron Microscopy Data Bank under accession codes EMD-30378 and EMD-30379. The coordinate data of the periplasmic region model of the MS-ring are available from the corresponding authors on request.

**SUPPLEMENTAL MATERIAL**

Supplemental material is available online only.

| FIG S1, TIF file | 1.8 MB |
| FIG S2, TIF file | 2.8 MB |
| FIG S3, TIF file | 2.6 MB |
| FIG S4, TIF file | 2.4 MB |
| FIG S5, TIF file | 0.8 MB |
| FIG S6, TIF file | 0.9 MB |
| FIG S7, TIF file | 1.5 MB |

**TABLE S1, DOCX file | 0.03 MB**

**TABLE S2, DOCX file | 0.04 MB**

**TABLE S3, DOCX file | 0.03 MB**

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