Native structure of flagellar MS ring is formed by 34 subunits with 23-fold and 11-fold subsymmetries

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Abstract

The bacterial flagellar MS ring is a transmembrane complex acting as the core of the flagellar motor. It not only acts as the template for rod and C ring assembly but also houses the type III protein export gate for assembly of the rod, hook and filament. The cytoplasmic C ring, involved in torque generation and rotation switch, is directly attached to the MS ring, and a symmetry mismatch between 26-fold MS ring and 34-fold C ring had been a long puzzle as to whether this would play some role in motor function. Although this puzzle seemed to have been resolved by the recent high-resolution structure of the MS ring with 33-fold symmetry with a variation from 32-fold to 35-fold because the C ring also shows a similar symmetry variation, it still remained ambiguous whether their symmetries are matched in the native motor structure. Here we show that the native MS ring structure formed by full-length FliF is 34-fold with no symmetry variation whereas the C ring has a small symmetry variation, indicating a flexibility in C ring assembly to generate small symmetry mismatches. We also show two conformations of FliF in part of its periplasmic region to form the 34-subunit ring with 23-fold and 11-fold subsymmetries in the inner and middle M ring, respectively, to accommodate the export gate at the center of the M ring.

Bacteria actively swim in liquid environments by rotating long, helical filamentous organelle called the flagellum. The bacterial flagellum is supramolecular motility machinery consisting of the basal body that acts as a bi-directional rotary motor, the filament that functions as a helical propeller, and the hook as a universal joint connecting the basal body and the filament to transmit motor torque to the filament\cite{1-3}. The basal body consists of the MS ring, C ring, LP ring and the rod (Fig. 1a). The MS ring is a transmembrane protein complex made of FliF and is the base for flagellar structure, assembly and function. The MS ring is not only the mounting platform for the C ring, which is the cytoplasmic part of the basal body formed by the switch proteins FliG, FliM and FliN and acts as a bi-directional rotor of the flagellar motor as well as the switch regulator of the rotation direction, but is also a housing for the type III protein export apparatus that exports flagellar axial proteins for their assembly at the distal growing end of the flagellum\cite{2-4}. The rod is a drive shaft of the motor, transmitting motor torque to the hook and filament and is a helical assembly of four different rod proteins, FlgB, FlgC, FlgF and FlgG. The LP ring is a bushing tightly embedded in the peptidoglycan layer as well as the outer membrane and surrounds the distal, thicker tubular part of the rod.
formed mainly by FlgG\textsuperscript{5,6} to stabilize the high speed rotation of the motor\textsuperscript{1-3}. Motor torque is generated by cyclic interactions and dissociation of FliG at the top end of the C ring with the cytoplasmic domain of the transmembrane stator complex formed by MotA and MotB, which also acts as a proton channel to couple proton influx across the membrane with torque generation\textsuperscript{7}, where multiple number of stator units surround the rotor to become active depending on the external load\textsuperscript{8}.

For a long time until recently, the MS ring had been believed to have 26-fold rotational symmetry whereas the C ring has 34-fold symmetry with some minor variations based on biochemical and structural studies\textsuperscript{9-13}. This symmetry mismatch between the two rotor complexes had been a focus of debates on its possible role in the C ring assembly, torque generation and stepping rotation and the basis for the mechanistic understanding of torque generation mechanism\textsuperscript{11,14,15}. This puzzle was sort of resolved by the recent high-resolution structure of the MS ring with 33-fold symmetry with a variation from 32 to 35 (ref. \textsuperscript{16}) because the C ring also shows a similar symmetry variation peaked around 34- or 35-fold\textsuperscript{12,17}. However, it still remained ambiguous whether their symmetries are actually matched in the native motor structure, because the distributions of the symmetry variations are clearly different between the two rings, with a peak around 34- or 35-fold for the C ring\textsuperscript{12,17} and 33-fold for the MS ring\textsuperscript{16}. Also, the MS ring structures were analyzed for those formed by overexpressed FliF with some C-terminal truncations\textsuperscript{16}, which may have produced the observed symmetry variation.

There is also another interesting symmetry mismatch between the MS ring and the rod, which assembles directly on the MS ring while having a helical tubular structure with 11 protofilaments, with a helical symmetry of about 5.5 subunits per one turn of helix. The core structure of the type III protein export gate composed of FliP, FliQ and FliR\textsuperscript{18} is likely to be located inside the MS ring with a helical nature in their assembly to be directly connected with the proximal part of the rod formed by FlgB, FlgC and FlgF via an adapter protein FliE\textsuperscript{19}. However, it remains unknown how the symmetry mismatch between the protein export gate-rod and the MS ring is resolved to make flagellar assembly possible by the MS ring as the assembly template.

Here we report electron cryomicroscopy (cryoEM) structural analyses of the flagellar basal body and the MS ring formed by full-length FliF from \textit{Salmonella enterica} serovar Typhimurium (hereafter referred to \textit{Salmonella}) and that the symmetry of the native MS ring is 34-fold with no variation. Our own symmetry analysis of the C ring of the basal body showed a variation with a peak at 34-fold in agreement with the previous observation\textsuperscript{12},
indicating that there are still small symmetry mismatches between the two rings possibly
generated by some flexibility in the initiation of C ring assembly by FliG around the MS ring. 
The MS ring structure also showed 23-fold and 11-fold symmetries in the inner and middle 
parts of the M ring, respectively, possibly to accommodate the type III protein export gate to 
assemble at the center of the MS ring. The structure of the S ring and cylindrical collar in the 
upper part of the MS ring is now resolved at high resolution, and mutational analyses show 
that conserved residues of FliF are responsible not only for MS ring formation by stabilizing 
intersubunit interactions but also for the assembly of the type III protein export apparatus and 
the rod within the MS ring. Thus, it is now clear that the MS ring and C ring are tightly bound 
to each other to act as a rotor unit of the flagellar motor to generate and transmit torque to the 
rod, hook and filament for bacterial motility.

CryoEM structural analysis of the basal body

We isolated the flagellar basal body from a Salmonella mutant strain HK1003 [flgEΔ(9-20) 
ΔclpP::Cm deletion mutant], in which the number of the basal body was increased by the 
deletion of the ClpP protease, and analyzed the structure by cryoEM image analysis. Most 
of the isolated basal bodies are missing the LP ring because the LP ring easily dissociates 
from the rod in the absence of the hook (Fig. 1b, Extended Data Fig. 1a). This structure is 
called a rivet because of its shape, and this partial basal body structure is advantageous for 
visualizing the symmetry of the MS ring in the end-on view because the LP ring normally 
above the MS ring disturbs the visualization of the MS ring image. The two-dimensional (2D) 
class average images of its end-on views thus allowed the determination of the MS ring 
symmetry (Fig. 1c, Extended Data Fig. 1b). Of the 34,896 particle images of the basal body 
that were picked up from the 1,578 cryoEM micrographs, 23,104 particle images were near 
end-on views, separated into four classes with slight differences in the orientation, where each 
of the four class images clearly showed the 34-fold rotational symmetry (Fig. 1c). So the MS 
ring structure in the native flagellar basal body is composed of 34 FliF subunits.

We also analyzed the symmetry of the C ring of the native flagellar basal body. Although 
the symmetry of the C ring in the basal body has already been studied by cryoEM image 
analysis, the previous study used an indirect method in which side view images of the C ring 
were subjected to multireference alignment against projections of three-dimensional (3D) 
models with symmetries from 32- to 36-fold to sort the particle images into different 
rotational symmetries, and this may have produced some errors by misalignment and 
possible deformation of the C ring from a perfect cylinder. We therefore used only end-on
views of the C ring that show blobs of subunits and analyzed the rotational symmetry (Extended Data Fig. 1c). Since the flagellar motor can switch the rotation between the counter-clockwise (CCW) and clockwise (CW) direction, we looked at both CCW and CW locked C ring structures. As shown in Fig. 1d, the C ring symmetry has a certain range of variation from 32- to 35-fold for CCW and 33- to 36-fold for CW, but about two thirds are 34-fold symmetry with rather minor populations of non 34-fold symmetries (CCW: 32-fold, 2%; 33-fold, 20%; 34-fold, 66%; 35-fold, 13%, 36-fold, 0%; CW: 32-fold, 0%; 33-fold, 13%; 34-fold, 68%; 35-fold, 20%; 36-fold, 1%). This is in marked contrast to much higher populations of non 34-fold symmetries (32-fold, 2%; 33-fold, 17%; 34-fold, 42%; 35-fold, 30%; 36-fold, 7%) estimated in the previous study. Thus, although the symmetry mismatch between the MS ring and C ring is present, the probabilities of C ring assembly around the MS ring with extra or less FliG subunits to the FliF subunits is small, and the number of extra or less FliG subunits is mostly up to one, indicating that the flexibility in the template driven C ring assembly by FliG around the 34 FliF-subunit MS ring that causes a symmetry mismatch between these two rings is not so high.

**CryoEM structural analysis of the MS ring formed by overexpressed FliF**

We overexpressed full-length *Salmonella* FliF in *E. coli* and isolated the MS ring from the membrane fraction by sucrose gradient purification (Extended Data Fig. 2a,b). Although the ring-shaped particles were clearly observed in negative stained EM images (Extended Data Fig. 2c), it was difficult to determinate the structure of the MS ring by cryoEM image analysis because most of the particles were attached to the edge of carbon holes. To alleviate such awkward characteristics of the MS ring, we optimized the purification procedure using LMNG as a detergent, which drastically improved the particle dispersion and density in the holes and allowed high quality image data collection in sufficient particle numbers. A total of 339,861 particles were extracted from 1,589 micrographs and were analyzed. Representative 2D class averages showed homogeneous ring particles with clear 34-fold symmetry (Fig. 2a, Extended Data Fig. 3 and 4). We did not observe any other symmetries in the 2D class averages, indicating that the MS ring formed by full length FliF consists of 34 subunits and shows no variation in the ring stoichiometry, just as we observed for the MS ring in the native basal body structure as described above.

We then carried out 3D image reconstruction of the MS ring, but the resolution did not extend beyond 10 Å without rotational symmetry. We therefore performed iterative 3D refinement with 34-fold rotational symmetry. This process dramatically improved the
resolution of the 3D map to 3.7 Å resolution (Extended Data Fig. 3) and enabled us to
construct the atomic model of the S ring and collar (Fig. 2b,c). Unfortunately, however, a
large part of the M ring was somehow disordered in this particular construct.

The S-ring and collar is formed by residues 228–438 of Salmonella FliF (Fig. 2d), which
correspond to the latter half of the periplasmic region. The monomeric structure consists of
two structural regions: a globular domain with αββαβ motif that is known as a ring building
motif (RBM) (residues 228–270 and 382–438 designated as RBM3 in Fig. 2b); and a long,
extended up-and-down β structure (residues 271–381) consisting of a set of antiparallel chains
and a set of antiparallel β strands, β3 and β4, forming a β hairpin with invisible 51 residues
possibly forming a flexible loop (Fig. 2b). The chain connecting β3 and β4 at the top of the
collar (residues 305–354) was not modeled due to the poor density (Fig. 2b–d). The S ring is
made up of 34 RBM3 domains horizontally packed with their major axis oriented in the radial
direction to form a ring with a diameter of 24 nm. The long β hairpins in the upper part of the
collar are vertically lined up to form a 68-stranded cylindrical β-barrel structure. The two
extended antiparallel chains connecting the RBM3 domain and the vertical β hairpin are
inclined about 30 degrees from the ring axis. The model was nearly identical to the

corresponding part of the recent MS ring structure with 34-fold symmetry

The overall structure of the S-ring and collar resembles the SpoIIIAG structure despite
the low sequence identity (<15%) (Extended Data Fig. 5a,b). The unique large β-barrel
structure composed of vertically arranged β-strands is present in the SpoIIIAG ring. The ring
arrangement of the RBM3 domains is also similar to that of the SpoIIIAG, although the RBM
domain of the SpoIIIAG ring is tilted about 14 degrees compared with the nearly horizontal
orientation of that of the S ring (Extended Data Fig. 5a,b). The ring formation through the
RBM domain is commonly found in the T3SS injectisome, and the domain arrangement of
the S ring is similar to those in the T3SS ring except for the D2 domain of PrgH

The chain folding arrangement of FliF in the collar is also similar to those seen in the
secretin rings of the type III secretion system (T3SS) of bacterial pathogens. Unlike them,
however, only a small number of hydrogen bonds are formed between these two chains in the
flagellar MS ring because the two antiparallel chains are twisted and apart from each other
toward the bottom of the cylindrical β-barrel structure, even though the phi-psi angles of each
residue in these extend chain are mainly those of the β type conformation. Residues 283–293
connecting to β3 of the vertical β hairpin is looping out to form a structure that looks like a
saucer to the cup upon assembly into the ring. The two antiparallel chains are swapped right
and left before connecting to the vertical β hairpin in the upper part of the collar (Extended
Mutation analysis of FliF for the MS ring formation

We examined whether the atomic model of the S ring represents the structure of the physiologically functional MS ring of the flagellar basal body by mutations of FliF followed by assays of cell motility, flagellar protein export and assembly of the basal body and MS ring. The subunit interface of the RBM3 domains in the S ring is mainly mediated by the hydrophobic interactions. Ile-252, Leu-253, and Val-266 are relatively well conserved between FliF and SpoIIIAG\(^{21}\) (Fig. 3a). Ile-107 and Val-120 of SpoIIIAG corresponds to Ile-252 and Val-266 of FliF, respectively, and the replacement of each residue by Arg inhibits SpoIIIAG ring formation. We therefore replaced Ile-252, Leu-253 and Val-266 by Ala or Arg and analyzed the effects of these mutations on cell motility in soft agar to test whether these three residues are involved in MS ring formation (Fig. 3b). These substitutions did not significantly affect the steady cellular level of FliF as judged by immunoblotting with polyclonal anti-FliF antibody (Fig. 3c). Wild-type FliF fully restored motility of a ΔfliF mutant. The V266A mutant variant complemented the ΔfliF mutant to the wild-type level, and the L253A mutant variant restored the motility to a considerable degree, but the I252A mutant variant did so only poorly. The I252R, L253R and V266R mutant variants did not complement the ΔfliF mutant at all (Fig. 3b). In agreement with this, the I252A and L253A mutants formed a few flagellar filaments but the I252R, L253R and V266R mutants did not at all (Fig. 3d).

The MS ring is also a housing for the flagellar type III protein export apparatus\(^{28,29}\). Therefore, we tested whether the poor flagellar formation by the fliF(I252A) and fliF(L253A) mutant strains compared with the wild-type and fliF(V266A) mutant strains is a consequence of their reduced protein export activity. We examined the secretion levels of the hook-capping protein FlgD, the hook protein FlgE and the filament protein FliC as representative export substrates and found that the reduction in their secretion levels was well correlated with the reduction in the levels of flagellar formation (Fig. 3d,e). These two mutants produced hook-basal bodies (Fig. 3d), indicating that these two mutations do not affect MS ring formation so significantly. Therefore, these results suggest that the I252A and L253A mutations affect the assembly of the type III protein export apparatus into the MS ring. The I252R, L253R and V266R mutations abolished flagellar protein export (Fig. 3e), explaining the complete loss of motility of these mutant strains (Fig. 3b).

The cytoplasmic face of the MS ring is the template for assembly of the C ring by the
switch proteins FliG, FliM and FliN. Because the C ring is also a housing of the cytoplasmic part of the type III protein export apparatus and the loss of C ring considerably reduces the flagellar protein export activity and thereby impairs the cell motility, we tested whether the FliF mutations we examined above also affect C ring formation to cause the reduction in cell motility. If the C ring is formed normally on the cytoplasmic face of the MS ring, FliG, FliM and FliN should be detected in the membrane fraction of the cells because the MS ring is a transmembrane complex. Only a very small amount of FliN was found in the membrane fraction of the ΔfliF mutant as expected, but large amounts of FliN were detected in the membrane fractions of all the fliF mutant cells just as observed for wild-type cells (Fig. 3c). This suggests that these FliF mutations do not affect C ring formation at all and that mutations of Ile-252, Leu-253 and Val-266 have direct effects on the assembly of the type III protein export gate within the MS ring.

To examine the effect of these FliF mutations on the stability of the MS ring, we purified the MS rings from E. coli cells overexpressing these mutant variants of FliF and observed them by negative staining EM. All the FliF mutations significantly affected the stability, and most of the ring-shaped particles of the size of the MS ring were not completely closed (Fig. 3f). Ile-252 is located at the interface between FliF subunits whereas Leu-253 and Val-266 are not, suggesting that Ile-252 is responsible for intermolecular interactions between FliF subunits whereas Leu-253 and Val-266 are involved in proper folding of each FliF subunit to form the S ring. To further confirm this, we carried out in vitro disulfide cross-linking experiments.

Since His-263 and Ala-388 are in close proximity to Ile-252 in the atomic model of the S ring (Fig. 3a), we replaced Ile-252, His-263 and Ala-388 by Cys and analyzed cell motility in soft agar plates. Cys mutation of each residue did not drastically affect cell motility. The H263C variant complemented the ΔfliF mutant to the wild-type level, and the I252C and A388C variants restored the motility to a considerable degree although not to the wild-type level (Fig. 3g). However, neither I252C/H263C nor I252C/A388C double mutation variant complemented the ΔfliF mutant to restore cell motility, suggesting that crosslinked FliF subunits in the MS ring may have impaired its function, such as insertion of the type III protein export gate into the MS ring. Intersubunit disulfide crosslinking of FliF by the double Cys mutations was confirmed by higher-order oligomeric species observed in the membrane fraction isolated from the fliF(I252C/H263C) mutant upon inducing disulfide crosslinking by adding iodine (Fig. 3i, red dots), confirming that the atomic model of the S ring we built here is a physiological one. Such oligomers were observed neither for the wild-type, I252C nor
H263C mutant (Fig. 3i). The appearance of an extra band of FliF monomer with a slightly faster mobility in the SDS-PAGE gel (Fig. 3i, blue dot) can be due to intramolecular disulfide crosslink, suggesting a significantly different conformation of FliF monomer before its assembly into the MS ring. It is also possible that the conformational change of FliF monomer by intramolecular disulfide crosslink disturbs the proper assembly of the MS ring. The replacement of Cys-263 by Tyr in this double Cys FliF mutant considerably restored the motility (Fig. 3h), suggesting a requirement of flexibility in the MS ring structure for its proper function.

Internal structure of the MS ring with 23-fold and 11-fold subsymmetries

The 3D reconstruction of the basal body at about 6.8 Å resolution (Fig. 4a-c, Extended Data Fig. 6) from cryoEM image data shown in Fig. 1b revealed interesting features in the inner core of the M ring in addition to the 34-fold rotational symmetry of the S ring (Fig. 1c). The inner part of the M ring is a flat ring with clear 23-fold symmetry, and its central hole accommodates a helical assembly of many rod-shaped densities possibly representing α-helices, which looks very similar to the structure of the export gate complex formed by FliP, FliQ and FliR\textsuperscript{19}. As we stated earlier, cryoEM 3D reconstruction of the MS ring structure from an FliF overexpression construct had a large part of the M ring density disordered and therefore not showing its internal feature in detail (Extended Data Fig. 3), but the MS ring formed by FliF overexpressed from another construct produced the entire MS ring density that can be identified in the basal body (Fig. 4c,e, Extended Data Fig. 7). We identified an 11-fold symmetry density features just outside the inner M ring with 23-fold symmetry. Even 2D class average images showed the 11-fold symmetry just outside the collar (Fig. 4d, Extended Data Fig. 7). We therefore enforced 11-fold symmetry on the entire MS ring structure to see which parts of the MS ring have 11-fold symmetry (Fig. 4c). The outer part of the M ring was featureless but the middle part just outside the inner M ring with 23-fold symmetry showed a strong 11-fold symmetry feature as a hinge connecting to the outer M ring. These structural features indicate that the MS ring is formed by 34 FliF subunits with two distinct conformations. All the 34 subunits contribute their RBM3 in the latter half of the periplasmic region of FliF to the S ring and collar, but 23 copies form the inner M ring and 11 copies form the hinge with their RBM1 and RBM2 in the former half of the periplasmic region (Fig. 2 and 4c).

CryoEM structural analysis of the export gate complex formed by FliP, FliQ and FliR overexpressed in E. coli has revealed a helical nature of multi-subunit assembly of the
FliP$_5$FliQ$_4$FliR (10 subunits) and FliP$_5$FliQ$_4$FliR$_1$FlhB$_1$ (11 subunits) complexes$^{19,30}$, and the helical parameter is similar to those of the flagellar axial structures, such as the rod, hook and filament, with about 5.5 subunits per turn of the 1-start helix, which can also be regarded as a tubular structure with 11 protofilaments. However, the 11-fold symmetry feature of the internal part of the M ring is not the one directly associated with the export gate at the center of the M ring. It is the 23-fold symmetry internal core ring of the M ring that forms a hole of the right size to accommodate the export gate at the center of the M ring over a symmetry mismatch.

**Discussion**

The long-term puzzle on the role of symmetry mismatch between the MS ring with 26-fold rotational symmetry and the C ring with 34-fold symmetry$^{9-13}$ had produced much debate on its role in the C ring assembly, torque generation and stepping rotation$^{11,14,15}$. This puzzle was sort of resolved by the recent high-resolution structure of the MS ring with 33-fold symmetry with a variation from 32- to 35-fold$^{16}$ because the C ring also shows a similar symmetry variation around 34- or 35-fold$^{12,17}$. However, it was still not clear whether the symmetries of these two rings are actually matched in the native motor structure, because the distributions of the symmetry variations are clearly different between the two rings, with a peak around 34- or 35-fold for the C ring$^{12,17}$ and around 33-fold for the MS ring$^{16}$.

By cryoEM structural analyses of the flagellar basal body isolated from Salmonella cells and the MS ring formed by overexpressed full-length FliF of Salmonella, we determined the rotational symmetry of the native MS ring structure to be 34-fold with no variation (Fig. 1). The recent high-resolution structures of the MS ring with 33-fold symmetry with some variation would therefore be an artifact, possibly produced by the C-terminal truncations of FliF$^{16}$. We also carried out the symmetry analysis of the basal body C ring in a more direct manner than the previous study$^{12}$ by looking at enface views of the ring revealing subunit blobs (Extended Data Fig. 1c) and showed a symmetry variation to have a peak at 34-fold with much less populations of non 34-fold symmetry rings. So, the symmetries of the two rings are matched in the majority of the flagellar basal body albeit there are still minor populations having symmetry mismatches of one more or one less subunit in the C ring. The C ring assembly is thought to be initiated by co-folding of the C-terminal chain of FliF extending out on the circumference of the MS ring in the cytoplasm and the N-terminal domain of the FliG$^{31,32}$. Also, mutation studies of Salmonella flagella have identified two types of FliF-FliG fusion proteins that are functional in motility$^{33}$. These results predicted a
rather strict 1:1 stoichiometry of FliF and FliG in the C ring assembly around the MS ring. However, our present structural analysis clearly shows symmetry mismatches between the two rings albeit rather small, suggesting a certain level of flexibility in FliG ring assembly around the MS ring to initiate the entire C ring assembly together with the other switch proteins FliM and FliN.

The structure of the S ring and cylindrical collar in the upper part of the MS ring is now resolved at high resolution (Fig. 2), and mutation analyses based on the atomic model show that conserved hydrophobic residues of FliF located at the intersubunit interface are responsible not only for MS ring formation by stabilizing intersubunit interactions but also for the assembly of the type III protein export apparatus and the rod within the MS ring (Fig. 3). Interestingly, The structures of the basal body and MS ring also showed 23-fold and 11-fold symmetries in the inner part of the M ring and a density feature at the center that looks exactly like the protein export gate complex formed by FliP, FliQ and FliR19 (Fig. 4). Contributing only 23 copies of RBM1-2 of total 34 to the formation of the inner core M ring would possibly be to form the central hole of an appropriate size for the protein export gate to efficiently assemble at the center of the MS ring. Thus, it is now clear that FliF is folded into two distinct conformations to build the MS ring structure with different symmetries in different parts for its multiple functions. This was also suggested by the recent high-resolution structures albeit their detailed features were unfortunately non-physiological ones possibly due to the C-terminal truncations of FliF16. The folding and assembly of 34 FliF subunits forming the multi-symmetry structure of the native MS ring will be described elsewhere. It is at least confirmed now that the MS ring and C ring are tightly bound to each other to act as a rotor unit of the flagellar motor to generate and transmit torque to the rod, hook and filament for bacterial motility.

Methods

Bacterial strains, plasmids and media

Bacterial strains and plasmids used in this study are listed in Table 1. L-broth and soft tryptone agar plates were prepared as described previously28,34. Ampicillin was added at a final concentration of 100 µg/ml.

Purification of the basal body
Salmonella HK1003 [flgEΔ(9-20) ΔclpP::Cm] (CCW motor) and TM022 [flgEΔ(9-20) ΔclpP::Cm fliGΔPAA] (CW motor) mutant cells, in which the number of the basal body was increased by the deletion of the ClpP protease, were grown in 5 l of L-broth [1% (w/v) tryptone, 0.5% (w/v) yeast extract, 1% (w/v) NaCl] at 37°C until the cell had reached a late-logarithmic stage. The cells were harvested by centrifugation at 4,600 × g for 10 min. The pellets were suspended in 240 ml of 50 mM Tris-HCl buffer at pH 8.0 containing 0.5 M sucrose in an ice bath. EDTA and lysozyme were added to final concentrations of 10 mM and 0.1 mg·ml⁻¹, respectively, and the mixture was stirred for 30 min to convert cells to spheroplasts. Triton X-100 and MgSO₄ was added to the mixture to final concentrations of 1% and 10 mM, respectively. After stirring in the ice bath for 1 h, the mixture was centrifuged at 15,000 × g for 20 min to remove insoluble debris. An aliquot of 5 M NaOH was added to the supernatant to adjust pH to 10.5. The solution was centrifuged at 60,000 × g for 60 min, and the pellet was suspended in a buffer containing 10 mM Tris-HCl pH8.0, 5 mM EDTA and 1% Triton X-100. After repeating the above procedure twice, the basal body was purified by sucrose density gradient centrifugation at 68,000 × g for 14 h. Fractions with 20-50% sucrose were collected and centrifuged at 60,000 × g for 60 min. The pellet was suspended in Buffer S (25 mM Tris-HCl pH 8.0, 1 mM EDTA, 50 mM NaCl) containing 0.05% Triton X-100 and 0.05% LMNG or Buffer I (50 mM Tris-HCl pH 8.0, 50 mM NaCl, 25 mM imidazole, 0.05% Triton X-100 and 0.05% LMNG).

Overexpression of FliF and purification of the MS ring

E. coli BL21 (DE3) strain carrying either pKOT112 (ref. 36) (for data set 1) or pKOT105 (ref. 37) (for data set 2) was grown overnight in 20 ml of a cell culture medium containing 0.1 g tryptone, 0.05 g yeast extract, 0.05 g NaCl, 50 µg/ml of ampicillin and 30 µg/ml of chloramphenicol in a shaker at 37°C. A 20 ml of overnight culture was added to a 2 l of a cell culture medium containing 10 g tryptone, 5 g yeast extract, 5 g NaCl, 50 µg/ml of ampicillin and 30 µg/ml of chloramphenicol and the cells were grown in a 37°C orbital shaker (100 rpm) until the culture density reached an OD600 of 0.5~0.7. An aliquot of 0.5 M isopropyl β-D-1-thiogalactopyranoside (IPTG) was added to 2 L cell culture medium to a final concentration of 0.5 mM, and the growth was continued in a 30°C orbital shaker (100 rpm) for 4 h. The cells were collected by centrifugation at 4,600 × g for 10 min at 4°C. Harvested cells were resuspended in 40 ml of a French press buffer (50 mM Tris-HCl pH 8.0, 5 mM EDTA-NaOH, 50 mM NaCl) containing a protease inhibitor cocktail (Complete, EDTA-free) and were disrupted using a French press at a pressure level of 10,000 psi. After cell debris and
undisrupted cells were removed by centrifugation (20,000 × g, 20 min, 4°C), the crude membrane fraction was isolated by ultracentrifugation (90,000 × g, 60 min, 4°C). The pellet was solubilized in 40 ml of Alkaline buffer (50 mM CAPS-NaOH pH 11.0, 5 mM EDTA-NaOH, 50 mM NaCl, 1% TritonX-100) by mechanical shearing with a 1 ml plastic syringe with 20 gauge needle and was incubated at 4°C for 1 h. After insoluble material was removed by centrifugation (20,000 × g, 20 min, 4°C), solubilized proteins were collected by ultracentrifugation (90,000 × g, 60 min, 4°C). The pellet was resuspended in 3 ml of Buffer S containing 0.1% Triton X-100 by mechanical shearing as above, and the sample was loaded onto a 15–40% (w/w) continuous sucrose density gradient in Buffer C (10 mM Tris-HCl pH 8.0, 5 mM EDTA-NaOH, 1% TritonX-100) and spun by centrifugation in a swing rotor (49,100 × g, 13 h, 4°C). Density fractions with a volume of 700 µl each were collected by a gradient fractionator (BIOCOMP, NB, Canada) and a fraction collector and analyzed by SDS-PAGE for a band of FliF. Peak fractions were collected, and the MS ring was further concentrated by ultracentrifugation (90,000 × g, 60 min, 4°C). The pellet was resuspended in 30 µl of Buffer S containing 0.05% LMNG.

Sample vitrification and cryoEM data acquisition

For the basal body sample, Quantifoil Mo 200 mesh R0.6/1.0 holey carbon grids (Quantifoil) were glow discharged on a glass slide for 10 sec. A 2.5 µL aliquot of the sample solution was applied to the grid and blotted by a filter paper for 3 s × 2 times with 2 sec drain time at 100% humidity and 4°C. For the MS ring sample, Quantifoil Cu 200 mesh R0.6/1.0 holey carbon grids were glow discharged on a glass slide for 30 sec. A 2.6 µl aliquot of the sample solution was applied onto the grid and blotted by a filter paper for 7 sec at 100% humidity and 4°C, The grids were quickly frozen by rapidly plunging into liquid ethane using a Vitrobot Mark III quick freezing device (Thermo Fisher Scientific). The grids were inserted into a Titan Krios transmission electron microscope (Thermo Fisher Scientific) operated at 300 kV, with the cryo specimen stage cooled with liquid nitrogen. CryoEM images were recorded with a Falcon II 4k × 4k CMOS direct electron detector (Thermo Fisher Scientific) at a nominal magnification of ×75,000 for the FliF ring data 1, corresponding to an image pixel size of 1.07 Å for high-resolution image analysis, or ×59,000 for the FliF ring data 2 and the basal body, corresponding to an image pixel size of 1.4 Å (Extended Data Table 2), using the EPU software package (Thermo Fisher Scientific). Movie frames were recorded at a dose rate of 45 e-/pix/sec over an exposure time of 2 sec. The total accumulated dose of 90 e-/Å² was
fractionated into 7 frames. Image data sets of 1,589 (data set 1) and 806 micrographs (data set 2) were collected for the FliF ring, and 1,589 (for 2D classification) and 18,256 (for 3D reconstruction) micrographs for the basal body, using a defocus range between 1.0 and 3.0 µm.

Symmetry analysis of the MS ring of the basal body

Many of the basal body particles in cryoEM images showed end-on views. The particle images were picked up using an in-house particle picking program applying a deep learning method based on YOLO neural network\(^3\). About 2,000 particles were manually picked from 200 micrographs and were used for a training of the neural network. In total 34,896 particles were extracted from 1,578 micrographs. 2D classifications were carried out using Relion-2.1 (ref. \(^3\)). After 2\(^\text{nd}\) 2D classification, class average images showing the rivet-like structure with the MS ring and the rod were used for rotational symmetry analysis of the MS ring. The outer part of the ring images was converted from Cartesian to polar coordinates, the autocorrelation function was calculated, and the rotational symmetry was analyzed by Fourier transformation (Extended Data Fig. 1b).

3D reconstruction of the basal body

The basal body in Buffer I was used for 3D structural analysis of the basal body. The particle images were picked up using an in-house particle picking program as mentioned above. About 2,000 particles were manually picked up from 200 micrographs and were used for training. In total 385,803 particles were extracted from 18,256 micrographs. 2D and 3D classifications and 3D reconstruction were carried out using Relion-2.1 (ref. \(^3\)) or Relion-3.0 (ref. \(^4\)). We used a 3D map of the basal body in a previous study\(^4\) as the initial 3D model. After subtracting the C-ring density, 275,548 good particle images were used to construct 3D images in five classes. The two best 3D classes were individually refined and then merged for the final 3D refinement. The final 3D map was reconstructed using 149,341 particles at a resolution of 6.8 Å.

Symmetry analysis of the C ring

CryoEM images of the wild-type (CCW) and CW mutant basal bodies were collected using a JEM-3200FSC electron cryomicroscope (JEOL) equipped with a liquid-nitrogen cooled specimen stage, an Ω-type energy filter and a field-emission electron gun, operated at an
accelerating voltage of 200 kV. The images were captured by an F415mp CCD camera (TVIPS) at a magnification of 88,800× corresponding to a pixel size of 1.69 Å, a defocus range of 1.0–2.5 μm and an electron dose of 40 e/Å². Defocus and astigmatism of the images were determined using CTFFIND3 (ref. 42). To estimate the symmetry of the C ring, end-on view images of the basal bodies were boxed out by BOXER 43. The C ring part of each end-on view image was converted from the Cartesian to polar coordinates, the autocorrelation function was calculated, and the rotational symmetry was analyzed by Fourier transformation (Extended Data Fig. 1c). The end-on view images with distinct rotational symmetries were also classified, aligned and averaged to show the C ring images with different symmetries in Extended Data Fig. 1c.

Image processing of the MS ring formed by overexpressed FliF

Image processing procedures of the MS ring are described in Extended Data Fig. 3 and 7. The movie frames were aligned to correct for beam-induced movement and drift by MotionCor2 (ref. 44), and the parameters of the contrast transfer function (CTF) were estimated by Gctf 45. For the FliF ring data set 1 with the outer M ring part disordered, in total 339,861 particle images were automatically picked up from 1,589 micrographs using Gautomatch (http://www.mrc-lmb.cam.ac.uk/kzhang/), and then 2D and 3D classifications were performed using Relion-2.1 (ref. 39) and 3.0 (ref. 40). Particle images from good 2D class average images were selected for the initial 3D model using CryoSPARC2 (ref. 46). In total, 99,560 particles from the best 3D class were subjected to ab initio reconstruction and hetero refinement, and finally 38,889 particles from the best hetero refinement model were subjected to non-uniform refinement with C34 rotational symmetry using Relion-2.1 (ref. 39). The final refinement yielded a 3D map with a global resolution of 3.70 Å and a B factor of -197.5 Å² according to 0.143 criterion of the Fourier shell correlation (FSC). The local resolution was estimated using CryoSPARC2 (ref. 46). The processing strategy is described in Extended Data Fig. 3, and the model refinement statistics in Extended Data Table 2.

For the FliF ring data set 2 with the M ring part well ordered, in total 156,459 particle images were automatically picked from 806 micrographs using Gautomatch (http://www.mrc-lmb.cam.ac.uk/kzhang/), and 2D and 3D classifications were performed using Relion-2.1 (ref. 39). Particle images from good 2D class average images were selected for the initial 3D model using CryoSPARC2 (ref. 46). In total, 18,883 particles from the best 3D class were subjected to 3D refinement with C1, C11 and C34 rotational symmetry using Relion-2.1 (ref. 39). The final refinement and postprocessing yielded a 3D map with a global
resolution and a B-factor of 12 Å and -35 Å\(^2\) for C1, 9.0 Å and -495 Å\(^2\) for C11 and 7.4 Å and
-450 Å\(^2\) for C34, according to 0.143 criterion of the FSC. The processing strategy is described
in Extended Data Fig. 7, and the model refinement statistics in Extended Data Table 2.

**Model building and refinement of the S ring structure**

The atomic model of the S ring was constructed using Coot\(^{47}\) and refined with real space
refinement based on the cryoEM map (EMD-30612) using Phenix\(^{48}\) under NCS constrains of
34-fold rotational symmetry and secondary structure restraints. The refinement statistics are
summarized in Extended Data Table 2.

**Site directed mutagenesis**

Site-directed mutagenesis was carried out using the QuickChange site-directed mutagenesis
method (Stratagene). All of the flif mutations were confirmed by DNA sequencing. DNA
sequencing reactions were carried out using BigDye v3.1 (Applied Biosystems) and then the
reaction mixtures were analyzed by a 3130 Genetic Analyzer (Applied Biosystems).

**Motility assays**

We transformed a *Salmonella* flif null mutant strain, TH12415 (Δflif), with pET22b-based
plasmids, and the resulting transformants were inoculated onto soft tryptone agar plates
containing 100 μg/ml ampicillin and incubated at 30°C.

**Fractionation of cell membrane.** *Salmonella* TH12415 cells harbouring an appropriate
plasmid were grown exponentially in 30 ml L-broth at 30°C with shaking. The cells were
harvested, resuspended in 3 ml PBS, and sonicated. After the cell debris was removed by
low-speed centrifugation, the cell lysates were ultracentrifuged (100,000 × g, 60 min, 4°C).
After carefully removing the soluble fractions, membranes were resuspended in 300 μl of
SDS-loading buffer and heated at 95°C for 3 min. After SDS-PAGE, immunoblotting with
polyclonal anti-FliF or anti-FliN antibody was carried out as described previously\(^{28}\). An ECL
prime immunoblotting detection kit (GE Healthcare) was used to detect target bands.
Chemiluminescence signals were detected by a Luminoimage analyzer, LAS-3000 (GE
Healthcare).

**Flagellar protein export assay.**
Details of sample preparations have been described previously\(^49\). Both whole cellular proteins and culture supernatants were normalized to a cell density of each culture to give a constant number of *Salmonella* cells. After SDS-PAGE, immunoblotting with polyclonal anti-FlgD, anti-FlgE or anti-FliC antibody was performed.

**in vivo disulfide crosslinking**

*Salmonella* TH12415 cells harboring pMMiF001, pMMiF008, pMMiF009 or pMMiF011 were exponentially grown in L-broth containing 100 μg/ml ampicillin at 30°C with shaking. Aliquots of the cultures containing a constant number of cells were centrifuged to harvest the cells. The membrane fraction was prepared as described previously\(^50\). Disulfide crosslinking were induced by adding iodine as described before\(^51\). After SDS-PAGE using 4–15% Mini-PROTEAN TGX Precast gels (Bio-Rad), immunoblotting with polyclonal anti-FliG antibody was carried out as described previously\(^28\).

**Data availability**

The cryoEM volume has been deposited in the Electron Microscopy Data Bank under accession code EMD-30612, EMD-30613, EMD-30360, EMD-30361, EMD-30363 and the atomic coordinates have been deposited in the Protein Data Bank under accession code 7D84. Other data are available from the corresponding author upon reasonable request.

**References**


37. Ueno, T., Oosawa, K. & Aizawa, S. M ring, S ring and proximal rod of the flagellar


Acknowledgements
We thank Kelly T. Hughes for his kind gift of a *Salmonella* fliF null mutant, Tomoko Yamamoto for a phage for *clpP* deletion, Hideyuki Matsunami for *Salmonella* HK1003 mutant strain and Noriyuki Takekawa for help preparing a figure. This research was supported in part by JSPS KAKENHI Grant Numbers 25000013 to K.N, 18K06155 and 20770083 to T.Miyata, 18K14639 to A.K, JP26293097 and JP19H03182 to T.Minamino, JP15H05593 and JP20K15749 to M.K., JP15H02386 to K.I., and MEXT KAKENHI Grant Numbers JP15H01640 to T.Minamino. This research was also supported by Platform Project for Supporting Drug Discovery and Life Science Research (BINDS) from AMED under Grant Number JP19am0101117 to K.N., by the Cyclic Innovation for Clinical Empowerment (CiCLE) Grant Number JP17pc0101020 from AMED to K.N. and by JEOL YOKOGUSHI Research Alliance Laboratories of Osaka University to K.N.

**Author contributions**

T.K. and K.N. conceived the project; A.K., T.K., T.Minamino and K.N. designed experiments; A.K. and T.Miyata prepared the sample for cryoEM; T.K. set up cryoEM imaging system including both hardware and software; A.K., T. Miyata, F.M. and T.K. collected and analyzed cryoEM image data; M.K. and T.Minamino performed genetic, biochemical and physiological experiments; A.K. and K.I. built the atomic model; All authors studied the atomic model; A.K., T.Miyata, T.Minamino, K.I, T.K. and K.N wrote the paper based on discussion with the other authors.

**Competing interests**

The authors declare no competing interests.

**Additional information**

Extended Data is available for this paper at

**Correspondence and requests for materials** should be addressed to T.K. and K.N.
Fig. 1 Schematic diagram and cryoEM image analysis of the flagellar basal body. a, Schematic diagram of the flagellar basal body. IM: inner membrane; PG: peptidoglycan layer; OM: outer membrane. b, Representative 2D class average images in the first round. The upper row images show end-on views of the MS ring with rod called rivet (see Extended Data Fig. 1a) except for the second from the right; in the lower row, the two panels on the left show the rivet with the C ring attached. c, Images showing the rotational symmetry of the S ring are extracted from the second round of 2D class average of the rivet class images and magnified. The rotational symmetries obtained by image analysis as shown in Extended Data Fig. 1 are indicated below, presenting direct evidence for the 34-fold symmetry of the native MS ring. d, The rotational symmetry distributions of the C ring: CCW in yellow is the wild-type motor in the CCW state (98 particles in total); and CW in blue is the CW-locked motor by FlIG ΔPAA mutation (75 particles in total).
Fig. 2 Structure of the S ring and collar of the MS ring formed by full-length FliF. a, Representative 2D class average images. b, Cα ribbon representation of the RBM3 domain model forming the S ring and collar in two orthogonal side views. c, CryoEM 3D image reconstruction of the S ring and collar (left) and Cα ribbon diagram of the atomic model (right) in end-on (upper) and side (lower) views. d, Domain organization of Salmonella FliF, indicating the region forming the S ring and collar. TM1 and TM2: transmembrane regions; numbers above the bars: residue numbers in the FliF sequence.
Fig. 3. Mutational analysis of conserved Ile-252, Leu-253 and Val-266. 

a, Molecular interface between FliF subunits. Ile-252, Leu-253 and Val-266 residues are located at an interface between FliF subunits. His-263 and Ala-388 are in close proximity to Ile-252. 

b, Motility assay of Salmonella TH12415 cells harboring pET3c (ΔfliF), pMMiF001 (WT), pMMiF002 (I252A), pMMiF003 (I252R), pMMiF004 (L253A), pMMiF005 (L253R), pMMiF006 (V266A) or pMMiF007 (V266R) in soft agar. The plate was incubated at 30 °C for 8 hours. 

c, Membrane localization of the MS ring protein FliF and the C ring protein FliN. The membrane fractions of the above transformants were prepared after sonication and ultracentrifugation. Then, the membrane fractions were subjected to SDS-PAGE and analyzed by immunoblotting with polyclonal anti-FliF or anti-FliN antibody. Positions of FliF and FliN are indicated by markers. 

d, Immunoblotting analysis of FliF and FliN. Positions of FliF and FliN are indicated by markers. 

e, Western blots showing the expression levels of FliD, FliE, and FliC in the above transformants. 

f, Membrane fractions of the above transformants were subjected to SDS-PAGE and analyzed by immunoblotting with polyclonal anti-FliF or anti-FliN antibody. Positions of FliF and FliN are indicated by markers.
Electron micrographs of cells of TH12415 cells harboring pET3c (ΔfliF), pMMiF001 (WT), pMMiF002 (I252A), pMMiF003 (I252R), pMMiF004 (L253A) or pMMiF005 (L253R) and purified hook-basal bodies isolated from WT, I252A and L253A cells. e, Secretion assays. Whole cell proteins (Cell) and culture supernatant fractions (Sup) were prepared from the above strains. An 8 μl solution of each protein sample, which was normalized to an optical density of OD$_{600}$, was subjected to SDS-PAGE, followed by immunoblotting with polyclonal anti-FlgD (first row), anti-FlgE (second row) or anti-FliC (third row) antibody. f, Negative-stain EM images of the MS rings isolated from the six FliF mutants. g, Motility assay of _Salmonella_ TH12415 cells harboring pET3c (ΔfliF), pMMiF001 (WT), pMMiF008 (I252C), pMMiF009 (H263C), pMMiF010 (A388C), pMMiF011 (I252C/H263C) or pMMiF012 (I252C/A388C) in soft agar. The plate was incubated at 30 ºC for 7 hours. h, Isolation of pseudorevertants from the I252C/H263C mutant cells. Motility assay of _Salmonella_ TH12415 cells harboring pET3c (ΔfliF), pMMiF001 (WT), pMMiF008 (I252C), pMMiF011 (I252C/H263C) or pMMiF011SP-1 (I252C/H263Y) in soft agar. The plate was incubated at 30 ºC for 7.5 hours. i, Disulfide crosslinking of FliF–FliF interface. The membrane fractions were prepared from TH12415 cells expressing wild-type FliF FliF(I252C), FliF(H263C) or FliF(I252C/H263C), and disulfide crosslinking were induced by adding iodine. Then, each sample, which was normalized to an optical density of OD$_{600}$, was treated with N-ethylmaleimide, followed by non-reducing SDS-PAGE with a 4–15% gradient SDS-gel and finally immunoblotting with polyclonal anti-FliF antibody. Oligomeric and monomeric forms are indicated by red dots and an arrow, respectively. A blue dot presumably indicates FliF monomers with an intramolecular disulfide bond. Molecular mass markers are shown on the left.
Fig. 4 CryoEM structure of the 34-subunits MS ring with 23-fold and 11-fold subsymmetries. a, b, End-on views of the S ring (a) and inner part of the M ring (b) of the basal body, showing the 34-fold and 23-fold rotational symmetries, respectively. The dot- and rod-like densities at the center of the rings represent the structure of the type III protein export gate formed by FliP, FliQ and FliR. c, 3D reconstructions of the basal body (left) and the MS ring formed by full-length FliF (right) in side (upper) and end-on (lower) views. d, 2D class average image of the MS ring in end-on view, revealing the 11-fold rotational symmetry in the M ring. e, A central section of the MS ring density through the axis (blue) is well superimposed on that of the basal body density, indicating the structural identity of the MS ring formed by full-length FliF with that of the basal body.
Extended Data

Native structure of flagellar MS ring is formed by 34 subunits with 23-fold and 11-fold subsymmetries

Akihiro Kawamoto, Tomoko Miyata, Fumiaki Makino, Miki Kinoshita, Tohru Minamino, Katsumi Imada, Takayuki Kato and Keiichi Namba
Extended Data Fig. 1 CryoEM single particle 2D Image analyses of the basal body M ring and C ring. a, 2D class average images of the basal body in the first round with identification of the types of substructures schematically depicted below. b, The upper panel shows the second round 2D class average for the images of the rivet class (pink boxes in a). The rotational symmetry of the S ring was analyzed by image analysis described in the lower panels. c, Symmetry analysis of the C ring with end-on view images. The upper panel describes how the image analysis is done, and the lower panels show C ring images of different rotational symmetries. WT: the wild-type motor in the CCW state; and CW: the CW-locked motor by FliG ΔPAA mutation\textsuperscript{35}. The image presented here are averages of many particles for easy recognition of their symmetries, but the actual symmetry analysis was done with individual particle images used in a previous study\textsuperscript{13}. 
Extended Data Fig. 2 Purification of the MS ring formed by full-length FliF. a, The sucrose gradient for purification of the MS ring with fraction numbers on the right. b, SDS-PAGE band pattern of each fraction of the sucrose gradient shown in a. c, Negative-stain EM image of the purified MS rings.
Extended Data Fig. 3 CryoEM single particle 3D image analysis of the MS ring formed by FliF expressed from a plasmid pKOT112 (ref. 36) (data set 1).
Extended Data Fig. 4 CryoEM single particle 2D image analyses of the MS ring. Upper three panels show 2D class average images of the MS ring in end-on views with slightly different orientation, which all clearly indicate 34-fold rotational symmetry of the S ring and collar, as analyzed by converting the images into the polar coordinates as shown in the lower panel.
Extended Data Fig. 5 Structural comparison of the S ring and SpoIIIAG. a, the S ring; b, SpoIIIAG (PDB ID: 5WC3) in Cα ribbon representation with rainbow color according to the sequence from the N-terminus in blue to the C-terminus in red. c, Magnified views of part of FliF forming the collar above the S ring, as indicated by the square box in a.
Extended Data Fig. 6 CryoEM single particle 3D image analysis of the flagellar basal body focusing on the MS ring and rod.
Extended Data Fig. 7 CryoEM single particle 3D image analysis of the MS ring formed by FliF expressed from a plasmid pKOT105 (ref. 37) (data set 2).
### Extended Data Table 1. Strains and plasmids used in this study

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*pKOT112 was supposed to contain a sequence corresponding to a C-terminally truncated FliF fragment (Ser-1 to Asp-456) but the plasmid we received actually contained full-length FliF.*
Extended Data Table 2  CryoEM data collection, refinement and validation statistics

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Refinement

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