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Atomic structure of anthrax PA pore elucidates toxin translocation

Jiansen Jiang^{1,2}, Bradley L. Pentelute³, R. John Collier⁴, and Z. Hong Zhou^{1,2,‡}

¹Department of Microbiology, Immunology and Molecular Genetics, University of California, Los Angeles, California 90095, USA.

²California NanoSystems Institute, University of California, Los Angeles, California 90095, USA.

³Department of Chemistry, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139, USA.

⁴Department of Microbiology and Immunobiology, Harvard Medical School, Boston, Massachusetts 02115, USA.

Summary

Anthrax toxin, comprising protective antigen (PA), lethal factor (LF) and edema factor (EF), is the major virulence factor of *Bacillus anthracis*, an agent that causes high mortality in human and animals. PA forms oligomeric prepores that undergo conversion to membrane-spanning pores by endosomal acidification, and these pores translocate the enzymes LF and EF into the cytosol of target cells¹. PA is not only a vaccine component and therapeutic target for anthrax infections but also an excellent model system for understanding the mechanism of protein translocation. Based on biochemical and electrophysiological results, researchers have proposed that a Φ -clamp composed of Phe427 residues of PA catalyzes protein translocation via a charge-state dependent Brownian ratchet^{2–9}. Although atomic structures of PA prepores are available^{10–14}, how PA senses low pH, converts to active pore and translocates LF and EF are not well defined without an atomic model of the PA pore. Here, by cryo electron microscopy (cryoEM) with direct electron counting, we have determined the PA pore structure at 2.9-Å resolution. The structure reveals the long-sought-after catalytic Φ -clamp and the membrane-spanning translocation channel, and supports the Brownian ratchet model for protein translocation. Comparisons of four structures reveal conformational changes in prepore to pore conversion that support a multi-step mechanism by which low-pH is sensed and the membrane-spanning channel is formed.

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‡ Corresponding author. Z. Hong Zhou Hong.Zhou@ucla.edu Phone: 1-310-983-1033.

Online Content Any additional Methods, Extended Data display items and Source Data are available in the online version of the paper; references unique to these sections appear only in the online paper.

Author Contributions J.J. designed and performed the experiments, analyzed data, and wrote the paper; Z.H.Z. initialized and supervised the research, analyzed data, and wrote the paper. B.L.P. and R.J.C analyzed data and wrote the paper.

Author Information 3D cryoEM density maps of anthrax PA pore have been deposited in the Electron Microscopy Data Bank under the accession numbers EMD-6224 (intact PA pore) and EMD-6225 (lacking the membrane-spanning β barrel). The coordinates of atomic model of anthrax PA pore have been deposited in the Protein Data Bank under the accession number 3J9C.

The authors declare no competing financial interests.

Triggering conversion from PA prepore to pore by *in vitro* acidification leads to rapid and irreversible aggregation. Attempts to prevent aggregation by screening detergents have largely failed¹⁵. By low pH treatment of PA prepores directly on EM grids containing a thin layer of continuous carbon film, we obtained dispersed particles of PA pore without aggregation (Extended Data Fig. 1). We then acquired drift-corrected cryoEM images (Extended Data Fig. 1b–d) and reconstructed a map at an overall resolution of 2.9 Å using 60,455 particles (Fig. 1, Extended Data Fig. 2, Supplementary Video 1). The resolution for most regions of the cryoEM map is ~2.8 Å (Extended Data Fig. 2c). Our map reveals rich high-resolution structural features, including amino acid side chains and 14 chelated Ca²⁺ ions (Extended Data Fig. 1e–h) and has allowed unambiguous *de novo* atomic modeling (Extended Data Table 1) and detailed structure and function analyses.

The overall structure of the PA pore has a "flower-on-a-stem" architecture, including corolla, calyx, and stem from top to bottom (Fig. 1b, 2a, Supplementary Video 1). Each PA protomer is divided into four domains in the PA prepore¹⁰, named 1', 2, 3, and 4. In the PA pore, domains 1', 3, and 4 form the corolla and domain 2 forms the calyx and the stem; therefore we designate the parts of domain 2 corresponding to the calyx and the stem as 2c (residues 259–274 and 354–487) and 2s (residues 275–353), respectively (Fig. 2b, c). Domains 1' and 2c form a compact structure responsible for substrate protein binding and intake (Fig. 2). Domain 2s is an extended β hairpin (2β_{2s} and 2β_{3s}), seven copies of which assemble to form a membrane-spanning 14-stranded β barrel 105 Å in length and 27 Å (from C_α to C_α) in diameter (Fig. 2). Domain 3 is located peripherally and has close contact with domains 1' and 2c (Fig. 2b). The cryoEM density of domain 4 is weak and has the lowest resolution among all domains (inset of Fig. 1b and Extended Data Fig. 2c), likely due to its flexibility from minimal contact with the other domains. Rigid-body fitting of domain 4 of the PA prepore crystal structure to the cryoEM map shows domain 4 shifts ~4 Å towards the central axis in the PA pore (Extended Data Fig. 3).

The translocation channel of the PA pore has a funnel shape and can be divided into four parts based on diameter: mouth, Φ-clamp, throat, and tube (Fig. 3a, b). Its surface is negatively charged and mainly hydrophilic, but hydrophobic patches are seen at the α-clamp¹⁴ of the mouth, near the Φ-clamp, and at the middle of the tube (Fig. 3a, c). As proposed based on the PA prepore structure¹⁰, the negatively charged surface promotes passage of cations, and the hydrophilic surface facilitates passage of substrate proteins/polypeptides.

The mouth has a 30-Å opening and inner diameters varying down to 20 Å (Fig. 3a), and can accommodate protein secondary structure elements, but not folded domains such as LF_N. The Φ-clamp below the mouth becomes the bottleneck of the entire channel with a solvent-excluded inner diameter of only 6 Å (Fig. 3a, b), which is smaller than protein secondary structure elements and therefore may only allow passage of fully unfolded polypeptides. Underneath the Φ-clamp are the throat, which is an enlarged (~18 Å) bulb-shaped chamber, and the tube formed by the 14-stranded β barrel with inner diameters in the range of 12–18 Å and rich in Ser and Thr residues in its middle region (Fig. 3a, b). The large diameter (>12 Å) of the throat and tube can accommodate an α helix which may be formed by polypeptides after passing the Φ-clamp (Fig. 3a). The diameter and the hydrophilic property

of this part of channel are similar to those of the exit tunnel of the ribosome, which is proposed to translocate α helices^{17,18}.

The mouth and the tube are the only two openings of the channel, and are accessible to the endosomal and cytosolic compartments, respectively (Fig. 3a). The rest of the channel is "water-tight" and without holes permeable to small molecules. The substrate protein blocks the small hole of the Φ -clamp prior to or during translocation². Thus the Φ -clamp may act as a gate separating the endosomal and cytosolic compartments. The differences of proton concentration (pH) and electrical potential (ψ) across the Φ -clamp (Fig. 3a) may serve as the primary driving force for substrate protein translocation³.

In contrast to the hydrophilic inner surface of the β barrel, its outer surface is largely hydrophobic (Fig. 3c, d). The Phe residues (313, 314, and 324) form two aromatic belts on opposite sides of the lipid bilayer (Fig. 3d), which may stabilize membrane insertion^{19,20}. The cryoEM map shows that the transmembrane region is surrounded by a cloud of disordered densities, which we interpret as bound detergent molecules that were added during sample preparation (Fig. 3e). Surprisingly, an additional hydrophobic surface in the middle of the β barrel containing a cluster of hydrophobic residues (I289, V332, I334, L338, and L340 from each protomer) was also bound with detergent molecules (Fig. 3d–f). However, the function of this additional hydrophobic region is unknown.

The overall architecture of the PA pore is similar to those of bacterial toxins α -hemolysin and *Vibrio cholerae* cytolysin (VCC) in their membrane-inserted conformations (Extended Data Fig. 4) despite differences in functions^{21,22}. The β barrel of the PA pore is twice long as compared to the other two toxins and may facilitate LF and EF translocation by entropically stabilizing an α helix within the confinement of the cylindrical channel^{9,17}. This longer barrel may also be necessary to accommodate its receptor situated between domain 4 and the host membrane¹¹ (Extended Data Fig. 4). Except for this length difference, the 14-stranded β barrels of the PA pore, α -hemolysin, and VCC share geometries, such as diameter, twist of β strands, and pitch length (Extended Data Fig. 4).

In the PA pore, the seven $2\beta_{10}$ – $2\beta_{11}$ loops of the heptamer converge to form an iris with a 6-Å hole bounded by a symmetric arrangement of the seven Phe427 residues (Fig. 4). By contrast, these loops in the PA prepore do not engage each other^{10,12}, but circumscribe a 30-Å central hole with adjacent Phe427 residues spaced ~14 Å apart (Extended Data Fig. 5a). In the PA pore, each $2\beta_{10}$ – $2\beta_{11}$ loop is stabilized by close interactions with the $2\beta_7$ – $2\beta_8$ loop of the same protomer and the $2\beta_{10}$ – $2\beta_{11}$ loops of its two neighboring protomers. The hydrogen bonds between Asn399 and Ser428 and between Asn399 and Lys397' of an adjacent protomer form a chain of interactions, which give rise to a ring of loops at the iris inside the PA pore (Extended Data Fig. 5b). Consistently, Ser428 mutant abolished protein translocation, and the Lys397 or Asn399 mutation resulted in a dominant negative effect^{23–25}. In the PA homologues found in *Clostridium* species, Lys397 and Asp426 are both replaced by uncharged Gln²⁶, implying that this chain of interactions might be formed differently in these homologues.

The aryl plane of Phe427 from each protomer is parallel to the central axis of the PA pore. Neighboring Phe427 residues interact with each other by aromatic CH- π interaction in a tilted T-shaped configuration²⁷ and possibly by hydrophobic interaction, thus forming the Φ -clamp (Fig. 4, Extended Data Fig. 5). The integrity of the Φ -clamp is required for catalysis of protein translocation, as mutation of even a single Phe427 residue of the Φ -clamp severely reduced translocation efficiency and disrupted the seal against cation passage⁵. Because the PA pore translocates polypeptides with various side chains, the Φ -clamp may act like an elastic "O-ring" changing its size and shape to allow an unhindered passage of different amino acid residues while maintaining a good seal during translocation. In the recently reported structure of bacterial amyloid secretion channel CsgG, eight Phe residues form a Φ -clamp, which differs from that in the PA pore in having a 9.5-Å hole and facilitating protein translocation in an ungated manner²⁸.

Evidence from electrophysiological studies indicates that the proton gradient across the endosomal membrane is the primary driving force for unidirectional translocation of proteins through the PA pore³. Proton-driven transporters usually involve protonation-dependent conformational changes of two or more alternating gates. In the PA pore, however, there is only one gate, *i.e.* the Φ -clamp, and neither the Φ -clamp nor its nearby residues likely undergo a protonation-dependent conformational change. Indeed, the PA pore structure supports the charge-state dependent Brownian ratchet model²⁻⁹, proposed earlier. In this model a negative electrostatic barrier within the pore hinders the passage, by Brownian motion, of deprotonated acidic residues^{3,4,29}. The fact that acidic residues in the acidic environment of the endosome have a higher probability of being protonated, and thus free to pass the barrier, than ones in the neutral environment of the cytosol, leads necessarily to unidirectional movement of polypeptides across the barrier. Consistent with this model, the PA pore structure shows three acidic residues, Asp425, Asp426 and Glu398, to be near the Φ -clamp and proximal to the pore axis (Fig. 3b), generating a strong negative electrostatic barrier demarcating the endosomal and cytosolic compartments. In addition, the highly conserved acidic residue Asp425 located directly underneath the Φ -clamp is ideally positioned as a proton sink that may strip off protons from protonated acidic residues passing the Φ -clamp (Fig. 4c). Besides the charge barrier in the Φ -clamp, efficient protein translocation may require additional charged spots, such as the top region of the β barrel⁸.

A key question regarding the low-pH triggered conversion from prepore to pore is how PA senses pH. A notable conformational change during the conversion is that the $2\beta_{10}$ - $2\beta_{11}$ loop is flipped from one side to the other (Fig. 5a, Extended Data Fig. 6). Interestingly, this loop has different conformations in the two crystal structures of PA octameric prepores ($[\text{PA}_{63}]_8$, PDB ID: 3HVD; $[\text{PA}_{63}]_8[\text{LFN}]_4$, PDB ID: 3KWV) whereas the remaining parts of the structures are largely unchanged^{9,13,14}; *i.e.*, this loop is in the pore and prepore states in 3HVD (Extended Data Fig. 6b) and 3KWV, respectively. This comparison suggests that the $2\beta_{10}$ - $2\beta_{11}$ loop can switch between two conformations without affecting the overall structure. Mutation of the conserved Asn422 or Asp425 in this loop abolished the prepore to pore conversion at low pH in a dominant negative manner²⁵. Therefore we interpret this loop as a pH sensor. Surprisingly, this loop does not possess any His, a residue whose pKa of ~ 6 falls in the pH range of 5-7 where the prepore to pore conversion occurs. The

conformational change of this loop might result from destabilization and rearrangement of hydrogen bonds upon exposure to low pH, or alternatively from perturbed pKa of Asp residues (425 and 426) in this loop. We note that other parts of PA might impact pH sensing. Prior work has shown mutation of residues near or in the $2\beta_2$ – $2\beta_3$ loop, or at the domain 2/ domain 4 interface, or deletion of the $2\beta_2$ – $2\beta_3$ loop changed the pH threshold for conversion but did not abolish the low-pH sensitivity^{25,30}. These observations suggest that the involved residues might regulate pH sensing.

We find that the prepore to pore conversion involves three layers of conformational changes, all located within domain 2 (Fig. 5b). The first layer is the above-mentioned pH-sensing $2\beta_{10}$ – $2\beta_{11}$ loop. Low-pH triggered conformational change of this loop is relayed to the following layers sequentially as detailed below.

The second layer involves the $2\beta_7$ – $2\beta_8$, $2\beta_5$ – $2\beta_6$, and $2\beta_{12}$ – $2\beta_{13}$ loops, which are positioned in an anti-parallel manner (Fig. 5a). In the PA pore, the N-terminal half of the $2\beta_7$ – $2\beta_8$ loop moves downward to form a U-shaped turn tethered to the putative pH-sensing $2\beta_{10}$ – $2\beta_{11}$ loop via a hydrogen bond between Asn399 and Ser428 (Fig. 5a, Extended Data Fig. 5b, 6e). This new inter-loop interaction may relay the low-pH triggered conformational changes from the first layer to the second layer as supported by the dominant negative effect of Asn399 or Lys397 mutant^{24,25}, which breaks the hydrogen bond network connecting these two loops (Extended Data Fig. 5b). As a result of the downward pull from the $2\beta_7$ – $2\beta_8$ loop, all three loops of the second layer become straightened in the PA pore, with parts of the $2\beta_5$ – $2\beta_6$ and $2\beta_{12}$ – $2\beta_{13}$ loops forming strands to augment $2\beta_6$ and $2\beta_{12}$, respectively (Fig. 5a, Extended Data Fig. 6c, f). This conformational change is supported by the observation that mutating any of Pro379, Val455, and Asn458 of these loops led to defects in protein translocation²⁵. This conformational change leads to the convergence of domains 2c and formation of the Φ -clamp by a 15° rotation of domain 2c around the hinge residues Ile261 and Tyr456, with its distal edge moving for 12 Å towards the central axis of the PA pore (Fig. 5c, Supplementary Video 2). Indeed, tethering domain 2 to domain 4 by receptor binding or stabilizing the interface between these two domains by a poly- γ -D-glutamate capsule impeded domain convergence and therefore decreased the pH threshold required for the prepore to pore conversion^{11,12,30}. Conversely, destabilizing the domain 2/domain 4 interface by point mutations increased the pH threshold³⁰.

The third layer, resulting from the second layer of conformational changes, involves release and refolding of the precursor of domain 2s (*i.e.*, $2\beta_2$, $2\beta_3$, $2\alpha_1$ and the membrane insertion loop), and leads to the formation of the membrane-spanning β barrel (Fig. 2, Fig. 5b, Extended Data Fig. 7). The rotation of domains 2c opens the pocket between domains 2c and 4 to release the precursors of domains 2s, which then come together to form the β barrel (see details in Extended Data Fig. 7b). The β barrel formation might be the result, rather than the cause, of the convergence of domains 2c, as supported by a cryoEM map obtained from a subset of particles that shows converged domain 2c in the pore state but lacks the β barrel (Extended Data Fig. 8). We note that the stability of the domain 2/domain 4 interface was found to be a rate-limiting barrier to the conversion³⁰ and the separation between domains 2c and 4 might also happen at an earlier stage of the conversion.

Taken together, these results suggest a multi-step mechanistic model of the low-pH triggered conversion from prepore to pore (Fig. 5d). First, the $2\beta_{10}$ – $2\beta_{11}$ loop as a pH sensor changes its conformation upon acidification (layer 1). Second, as a consequence, the $2\beta_7$ – $2\beta_8$, $2\beta_5$ – $2\beta_6$, and $2\beta_{12}$ – $2\beta_{13}$ loops become ordered (layer 2), resulting in the convergence of domains 2c and the formation of the Φ -clamp. Last, as a result of the separation of domain 2c from domain 4, the precursors of domains 2s are released and refold into a uniform β barrel (layer 3), which, as a final step, inserts into the endosomal membrane, ready for delivering the toxic enzymes into the cytosol.

In the PA pore, the convergence of domains 2c creates a large inter-protomer interface to stabilize the pore conformation (Extended Fig. 7a). Key to the formation of a functional PA pore is a newly identified interface between the $2\beta_9$ – $2\beta_{10}$ loop of one protomer and the triple-stranded β sheet ($2\beta_{12}$, $2\beta_6$, and $2\beta_7$) of its neighbor (Fig. 5e). The above-mentioned convergence of domains 2c brings them together to form a new binding interface primarily mediated by hydrogen bonds between the backbone of the loop and the side chains of Thr390, Thr393, and Asp451 of the β sheet (Fig. 5e). These interactions explain the dominant negative effect of Ser382 or Thr393 mutant, the loss of PA activity of the Cys mutant of Thr390 or Asp451, and the conservation of Thr390 and Thr393 in PA homologues²⁵. By contrast, these residues are accessible and free of interactions in the PA prepore, and thus might be potential drug targets for blocking the conversion to functional PA pores.

In summary, the 2.9-Å structure of anthrax PA pore suggests the $2\beta_{10}$ – $2\beta_{11}$ loop as a pH sensor to trigger conformational changes for prepore to pore conversion, supports a charge-state dependent Brownian ratchet model of proton-driven protein translocation, and can inform both efforts to engineer PA to target cancer cells and to design measures to block anthrax toxin entering cells^{14,21}. A more detailed mechanism of protein translocation through the PA pore awaits additional experimental evidence, such as structures of the PA pore in the act of polypeptide translocation.

METHODS

Preparation of PA prepores

PA₆₃ prepores were prepared following the described procedures³¹.

EM sample preparation and data acquisition

For negative-stain EM, 2 μ l of 0.1% polylysine solution (Polysciences) was first applied to a glow-discharged grid covered with carbon film and then removed by blotting with filter paper in 2 min. The polylysine treatment produced different orientations of particles on EM grids³², therefore overcoming the problem of preferred orientation for the PA pore. Immediately after removal of polylysine, 2 μ l of PA prepore (~50 μ g/ml) was applied to the grid and incubated for 1 min. The grid was then washed with the high pH buffer (50 mM HEPES, 50 mM NaCl, pH 8.0) twice followed by two washes with the low pH buffer (50 mM NaOAc, pH 5.0, 0.05% Igepal CA-630) to induce the conversion of prepore to pore. After removal of excess buffer, the grid was stained with 0.8% (w/v) uranyl formate.

For cryoEM, Quantifoil R1.2/1.3 holey grids were covered with a thin layer of continuous carbon film a day before use. The procedure for pore induction on cryoEM grids was carried out following the same procedure for the negative-stain EM except that the last step of staining was not used. About 1.5 μl of low pH buffer was left on the grids before they were transferred into an FEI Vitrobot Mark IV. The grids were then blotted and flash-frozen in liquid ethane in the Vitrobot at 100% humidity. The frozen grids were stored in liquid nitrogen before use.

Negative-stain EM micrographs were acquired with *Leginon* automation software^{33,34} and a TIETZ F415MP 16-megapixel CCD camera at 68,027 \times magnification in an FEI Tecnai F20 electron microscope operated at 200 kV. The micrographs were saved by 2 \times binning to yield a pixel size of 4.4 \AA .

Frozen-hydrated cryoEM grids were loaded into an FEI Titan Krios electron microscope operated at 300 kV for automated image acquisition with *Leginon*. CryoEM micrographs were recorded on a Gatan K2 Summit direct electron detection camera using the electron counting mode at 22,500 \times nominal magnification (calibrated pixel size of 1.28 \AA on the sample level) and defocus values ranging from -1.8 to -5.1 μm . The dose rate on the camera was set to ~ 8 $\text{e}^-/\text{pixel}/\text{s}$. The total exposure time was 8 s and fractionated into 32 frames of subimages with 0.25 s exposure time for each frame. Frame images were aligned and averaged for correction of beam-induced drift using the GPU-accelerated program from Yifan Cheng's lab³⁵. The average images from all frames were used for defocus determination and particle picking, and those from the first 16 frames (corresponding to ~ 20 $\text{e}^-/\text{\AA}^2$ total dose on sample) were used for 2D and 3D image classification. In total, 12,416 micrographs were taken in a continuous session. The best 7,062 micrographs were selected for the following in-depth data processing.

Image processing

For negative-stain EM single particle reconstruction, 140,775 particles were picked from 1,115 negative-stain EM micrographs using the *batchboxer* program of *EMAN*³⁶. Particles were windowed out in 96 \times 96 pixels. The defocus value of each micrograph was determined by *CTFFIND*³⁷ and particles were corrected for contrast transfer function (CTF) by phase-flipping with the corresponding defocus and astigmatism values using *Bsoft*³⁸. An initial model was generated using the *startcsym* program of *EMAN*. The refinement was then performed with 7-fold symmetry using *EMAN*.

For cryoEM single particle reconstruction, 21,200 particles (256 \times 256 pixels) were initially picked by hand from 1,928 micrographs and subjected to auto-refinement by *RELION*^{39,40} using the negative-stain EM map obtained above as the initial model. The resulting cryoEM map (~ 4 \AA resolution) was used to generate projections that were then served as templates to pick 259,719 particles from all of the 7,062 micrographs using the *batchboxer* of *EMAN*. In our procedure, the defocus values of the micrographs were determined by *CTFFIND* and particles were corrected for CTF by phase-flipping using *Bsoft*. The particles were processed with 2D and 3D classifications using the recommended procedures of *RELION* (http://www2.mrc-lmb.cam.ac.uk/relion/index.php/Recommended_procedures). 2D class averages and 3D class reconstructions were inspected and those without high-resolution and

interpretable features were considered as "bad" classes. Particles contributing to the bad classes were discarded. The remaining 60,455 particles were selected for the final structure refinement. The C7 symmetry was applied throughout the 3D classification and 3D auto-refinement.

To maximize usable signals from the frame images acquired with the K2 Summit camera, we employed the resolution and dose-dependent model of radiation damage recently introduced in *RELION-1.3* in the following steps⁴¹. First, the particle images averaged from all 32 frames with whole-image drift correction were used for a preliminary 3D auto-refinement. Second, particle images from individual frames were used to calculate translational alignments for the particle-based drift correction. A running average of seven frames, a standard deviation of one pixel, and fitting of linear tracks through the translations for all running averages were used for the optimal translational alignment following the suggested protocol of *RELION*. Last, particle images from frame 3 to frame 27 (~30 e⁻/Å² total dose on sample) were translated using the above optimal alignment and weighted with different B-factors as estimated from the single-frame reconstructions to generate optimal "shiny" average images. Application of this procedure to the above selected particles yielded 60,455 "shiny" particles.

These "shiny" particles were then subjected to 3D auto-refinement in *RELION* to generate the final cryoEM map. *RELION* post-processing with a soft auto-mask⁴² estimated a resolution of 2.9 Å by the gold-standard FSC at 0.143 criterion and a B-factor of -95 Å², and the post-processing without any mask reported a resolution of 3.3 Å. The accuracies of rotation and translation reported by *RELION* 3D auto-refinement were 1.67° and 1.01 Å. For visualization and atomic model building, the cryoEM map was sharpened and low-pass filtered by *RELION* post-processing using the above-mentioned B-factor and resolution. Local resolution was calculated by *ResMap*⁴³ using the two cryoEM maps independently refined from halves of data.

3D classification and auto-refinement also identified a subset of 21,632 particles that led to a cryoEM map at 3.6 Å resolution which is in the pore state but lacks the density corresponding to the 14-stranded β barrel, *i.e.* only has the corolla and calyx of the PA pore.

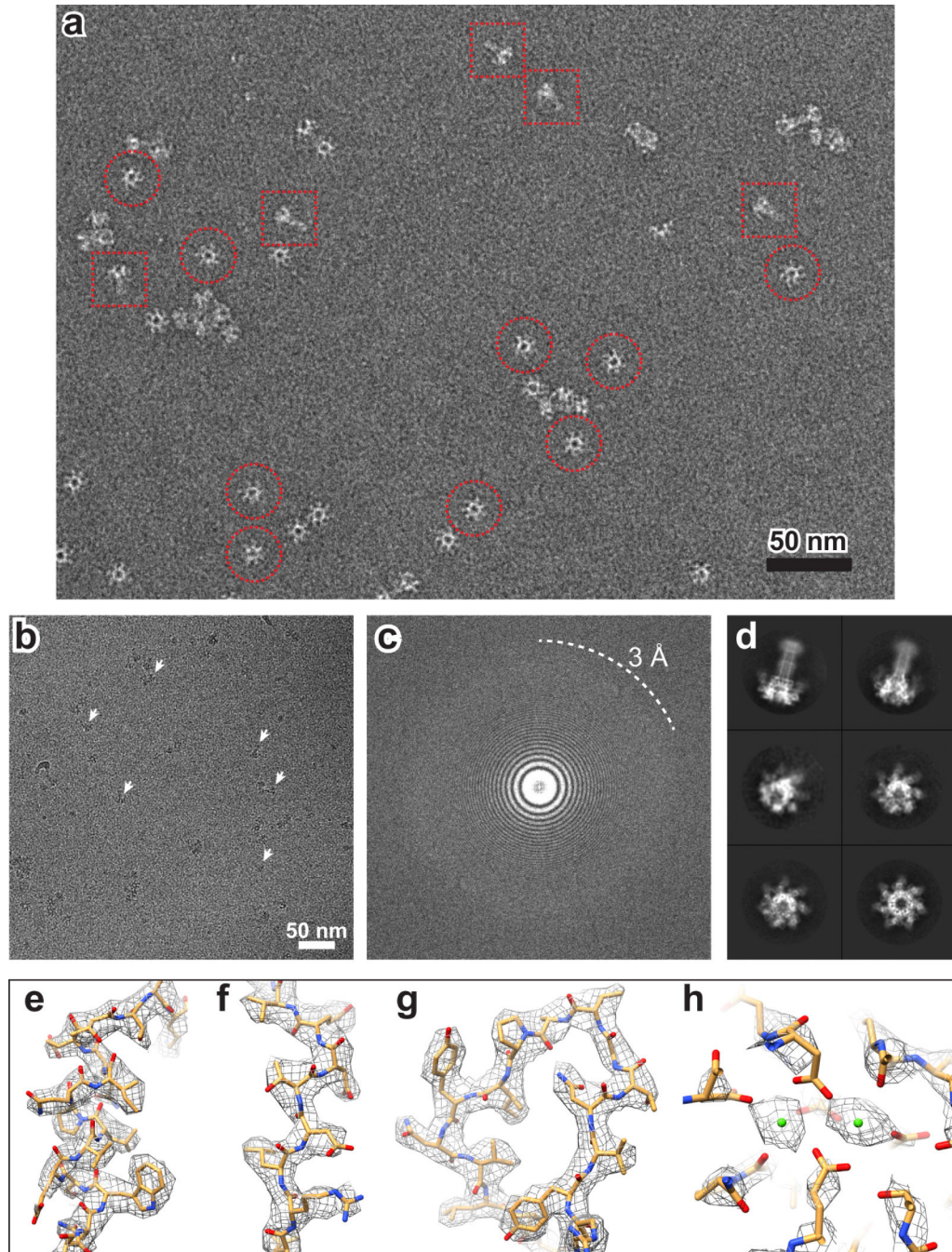
Atomic model building and refinement

De novo atomic model building of the PA pore except its domain 4 was carried out on the cryoEM map at 2.9 Å resolution using *Coot*⁴⁴. The coarse model was then refined using *PHENIX* in a pseudo-crystallographic manner⁴⁵. Note this procedure only improved the atomic model and did not modify the cryoEM map. Briefly, the cryoEM map was put into an artificial crystal lattice to calculate its structure factor using the *em_map_to_hkl.inp* utility program in *CNS*⁴⁶. The amplitudes and phases of the structural factor were used as pseudo-experimental diffraction data for model refinement in *PHENIX*. The restraints of Ramachandran, secondary structure, and non-crystallographic symmetry were used in the refinement.

The cryoEM maps and atomic models were visualized using *UCSF Chimera*⁴⁷ or *PyMOL*⁴⁸. The available crystal structures of PA₈₃ monomer (e.g. PDB ID: 1ACC and 3TEW) and

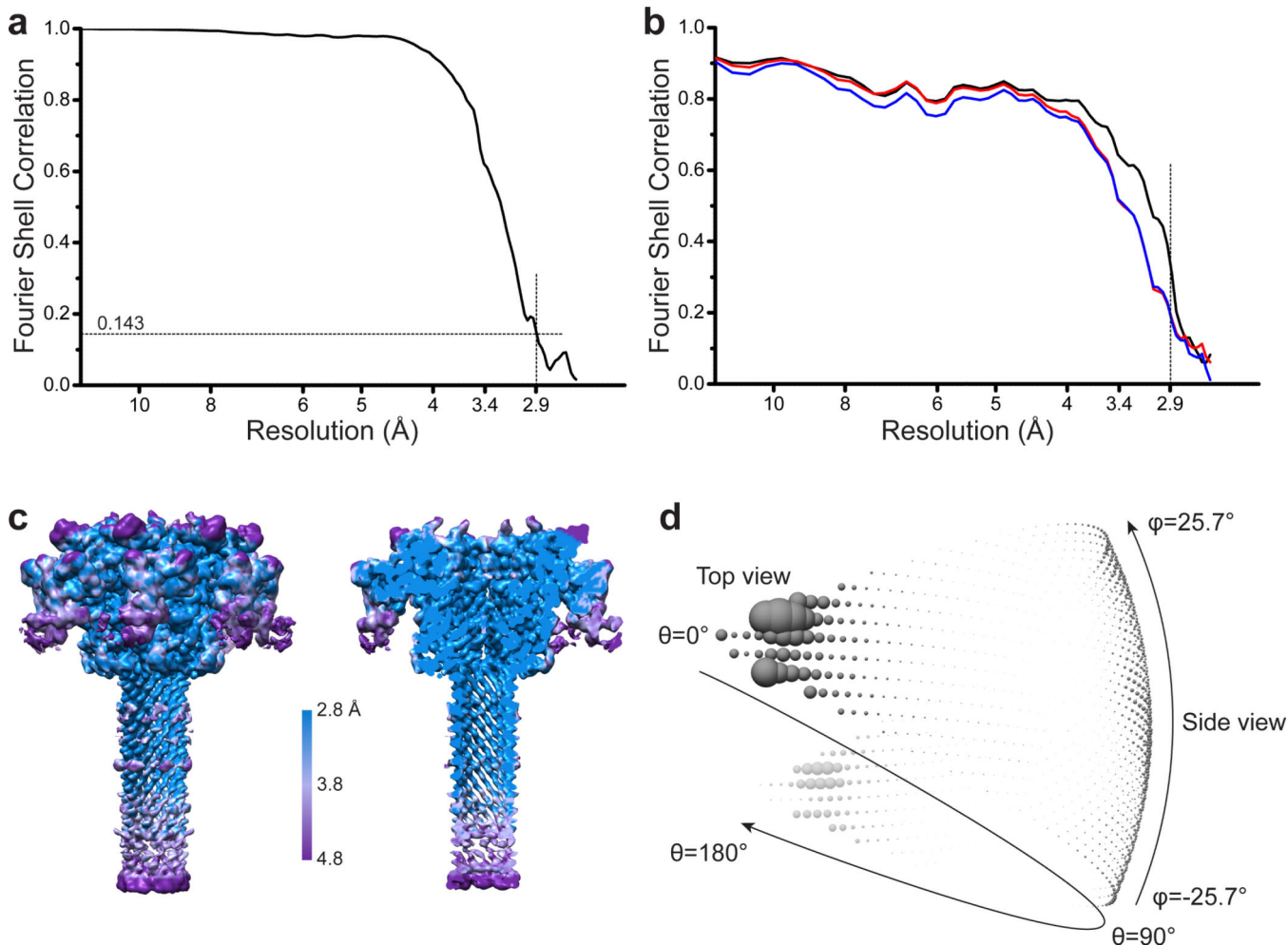
PA₆₃ heptameric prepore (PDB ID: 1TZO) are highly similar to each other, therefore 3TEW, which has the highest resolution among them, was used for structural comparison with our cryoEM structure of PA pore in most situations and 1TZO was used when the inter-protomer interaction or domain movement were considered.

Extended Data



Extended Data Figure 1. Negative-stain and cryoEM of the PA pore

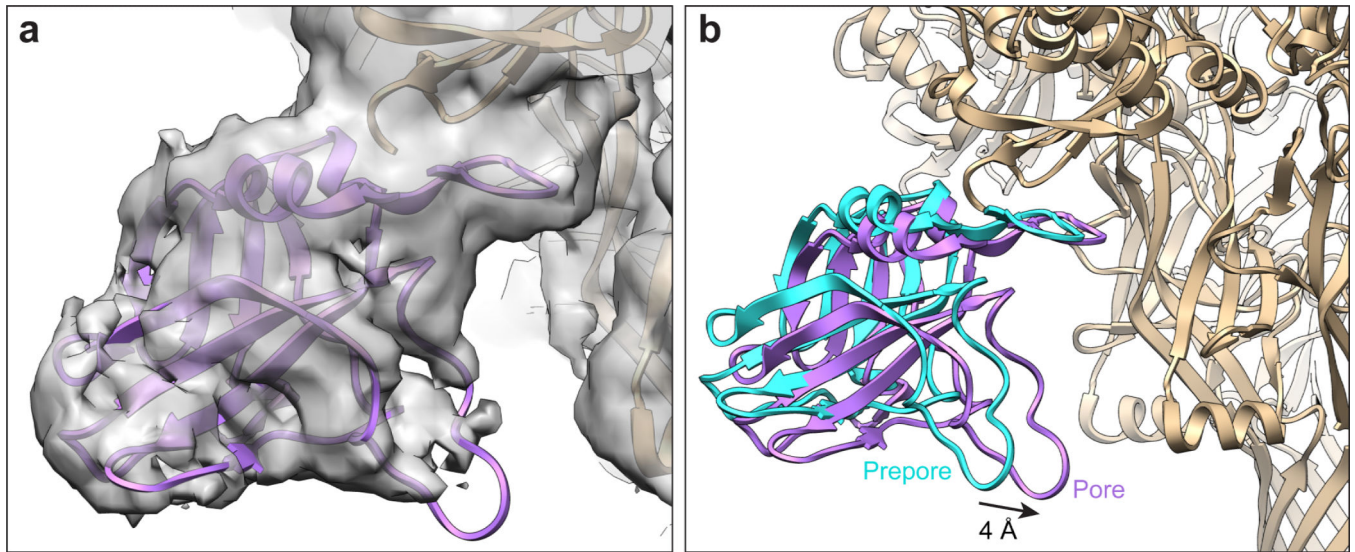
a, Negative-stain EM micrograph of PA pore particles. Some representative top-view and side-view particles are selected with circles and squares, respectively. **b**, A full-size drift-corrected cryoEM micrograph of 3710×3710 pixels of PA pore particles acquired from a Gatan K2 Summit direct electron detection camera, and at 300 kV accelerating voltage, $-2.7 \mu\text{m}$ defocus, and a total dose of $39 \text{ e}^-/\text{\AA}^2$. Some representative side-view particles of PA pore are indicated by arrows. **c**, Powerspectrum of the cryoEM micrograph in **(b)**. **d**, Representative cryoEM 2D class averages of particles at different orientations. **e-h**, Superimposition of representative regions of the cryoEM map (mesh) with the atomic model (stick), including α helix (**e**), β strand (**f**), loop (**g**), and Ca^{2+} ions (green spheres in **h**).



Extended Data Figure 2. Resolution estimation of the cryoEM single particle reconstruction of the PA pore

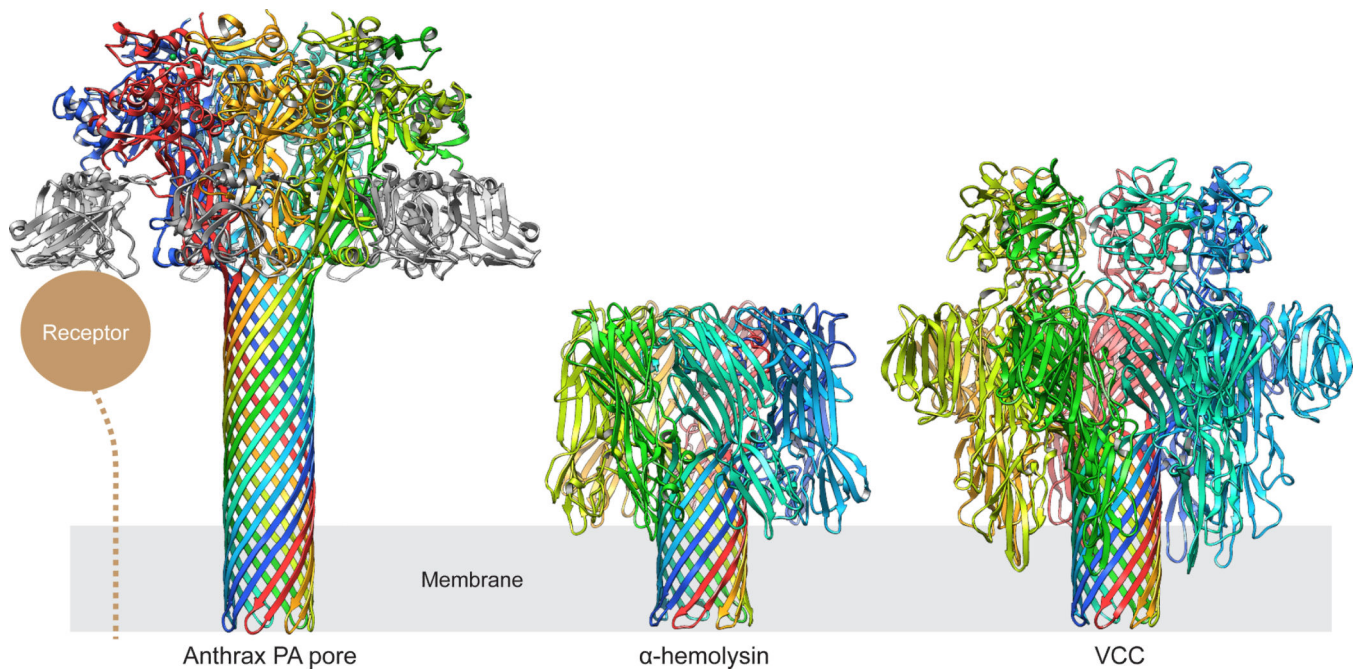
a, “Gold-standard” Fourier shell correlation (FSC) between two independently refined maps with an auto-mask that was corrected by phase randomization. The resolution was estimated by the gold-standard FSC at 0.143 criterion⁴². **b**, FSC of the final atomic model versus the final cryoEM map (black); of a model refined in the first of the two independent maps used for the gold-standard FSC versus the same map (red) and versus the second independent map (blue). **c**, Surface view and cut-through view of the unsharpened cryoEM map colored

by local resolution estimated by ResMap⁴³. **d**, Euler angle distribution of all particles used for the final 3D reconstruction.



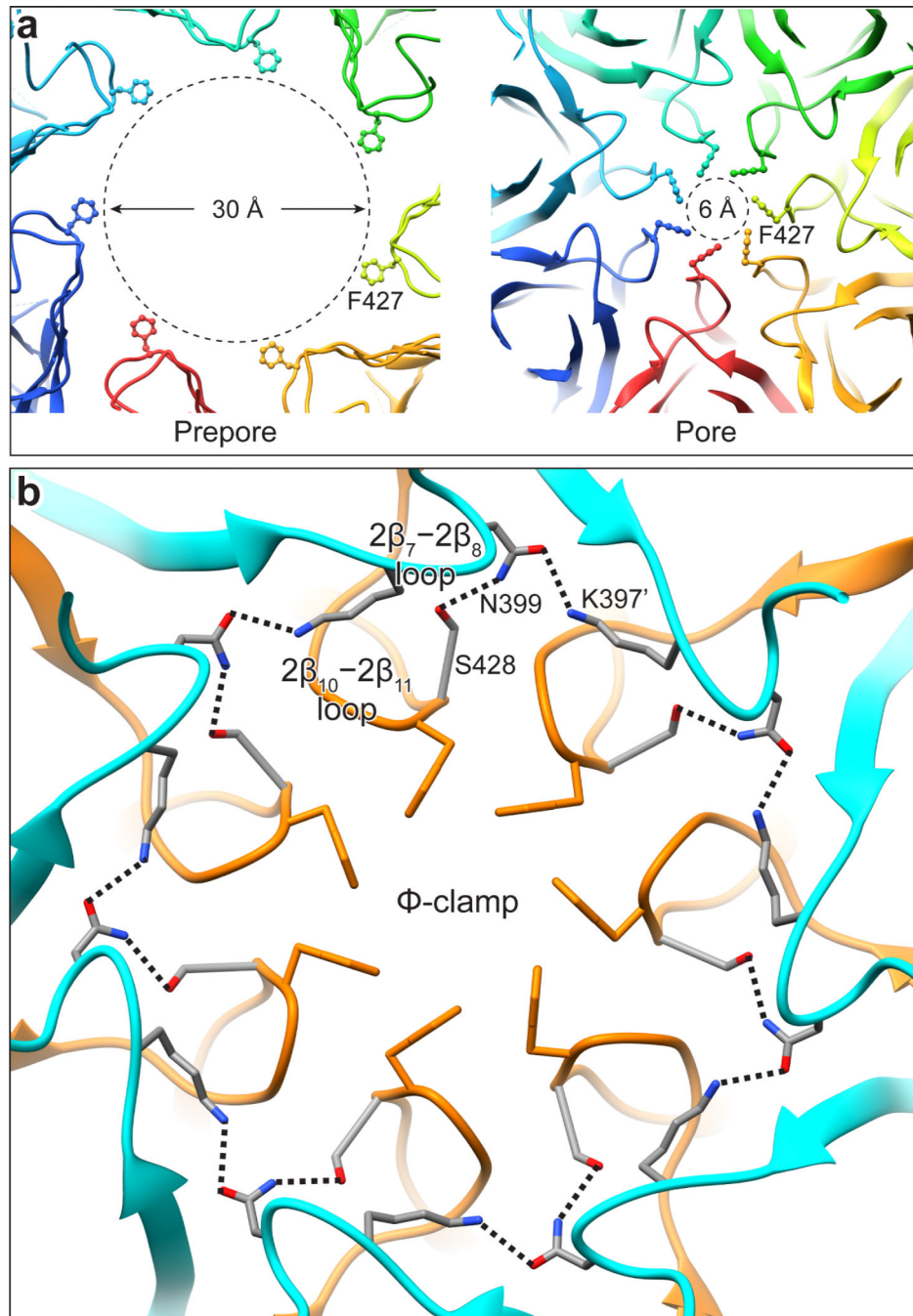
Extended Data Figure 3. Fitting of domain 4 into the cryoEM map of PA pore

a, The crystal structure of domain 4 (purple ribbons) from PA prepore (PDB ID: 1TZO) fits the cryoEM map (grey surface) of PA pore with a good agreement (cross-correlation coefficient: 0.91). **b**, Translocation of domain 4 during the prepore to pore conversion. Domain 4 is shifted inwards as a rigid-body for 4 Å from the prepore conformation (dark cyan) to the pore conformation (purple).



Extended Data Figure 4. Comparison of the structures of the PA pore, α-hemolysin (PDB ID: 7AHL), and *Vibrio cholerae* cytotoxin (PDB ID: 3O44)

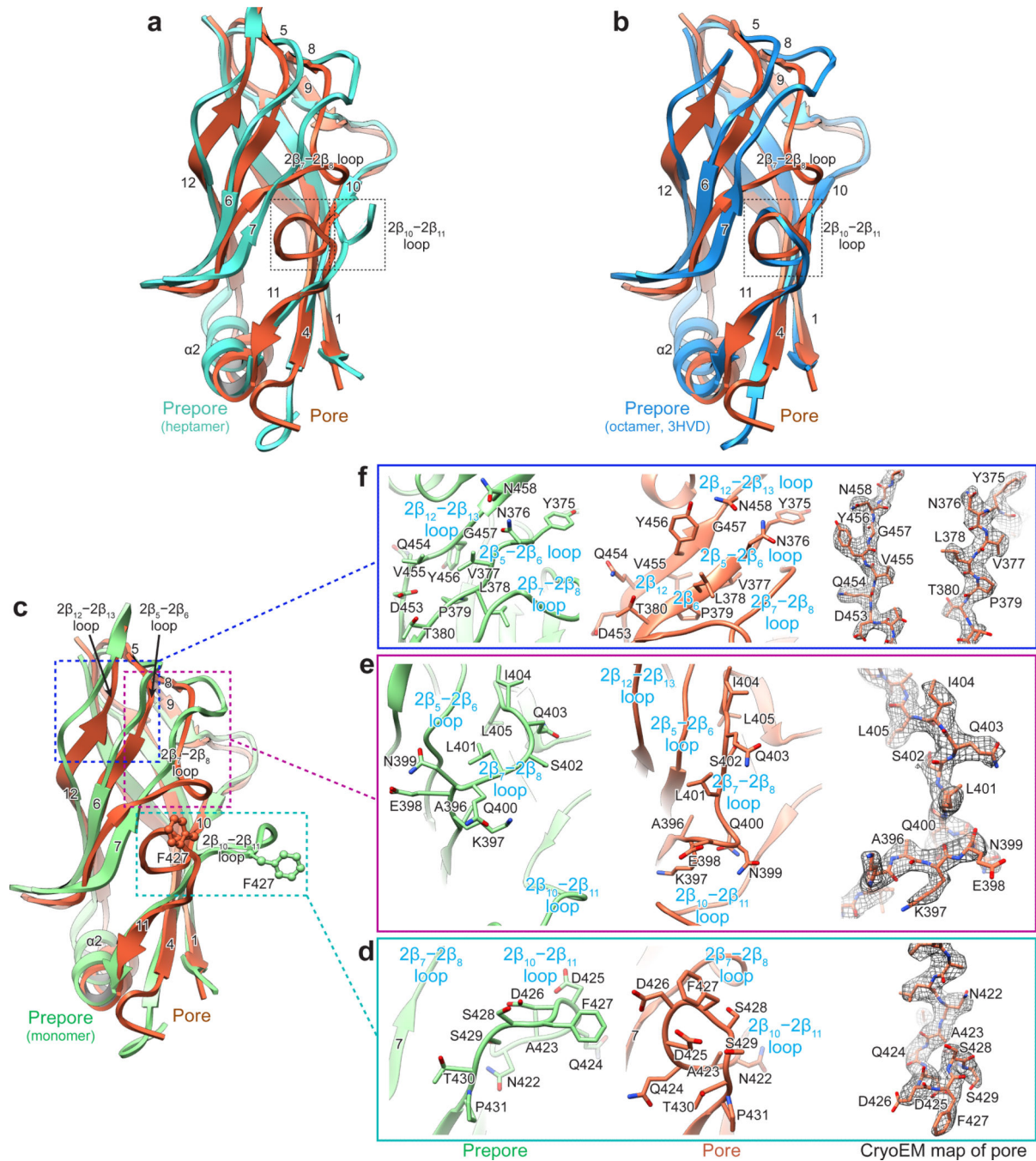
These three complexes are in their membrane-inserted forms. The approximate position of the lipid bilayer is illustrated with grey shade. The host cell receptor (TEM8 or CMG2) of PA that binds to domain 4 (grey ribbons) is schematically illustrated.



Extended Data Figure 5. Assembling of the Φ -clamp in the PA pore

a. Comparison of the Φ -clamp in the PA pore with its corresponding region in the PA prepore. The residues Asp426 and Phe427 missing in the crystal structure of the PA₆₃ prepore (PDB ID: 1TZO) are modeled based the crystal structure of the PA₈₃ monomer

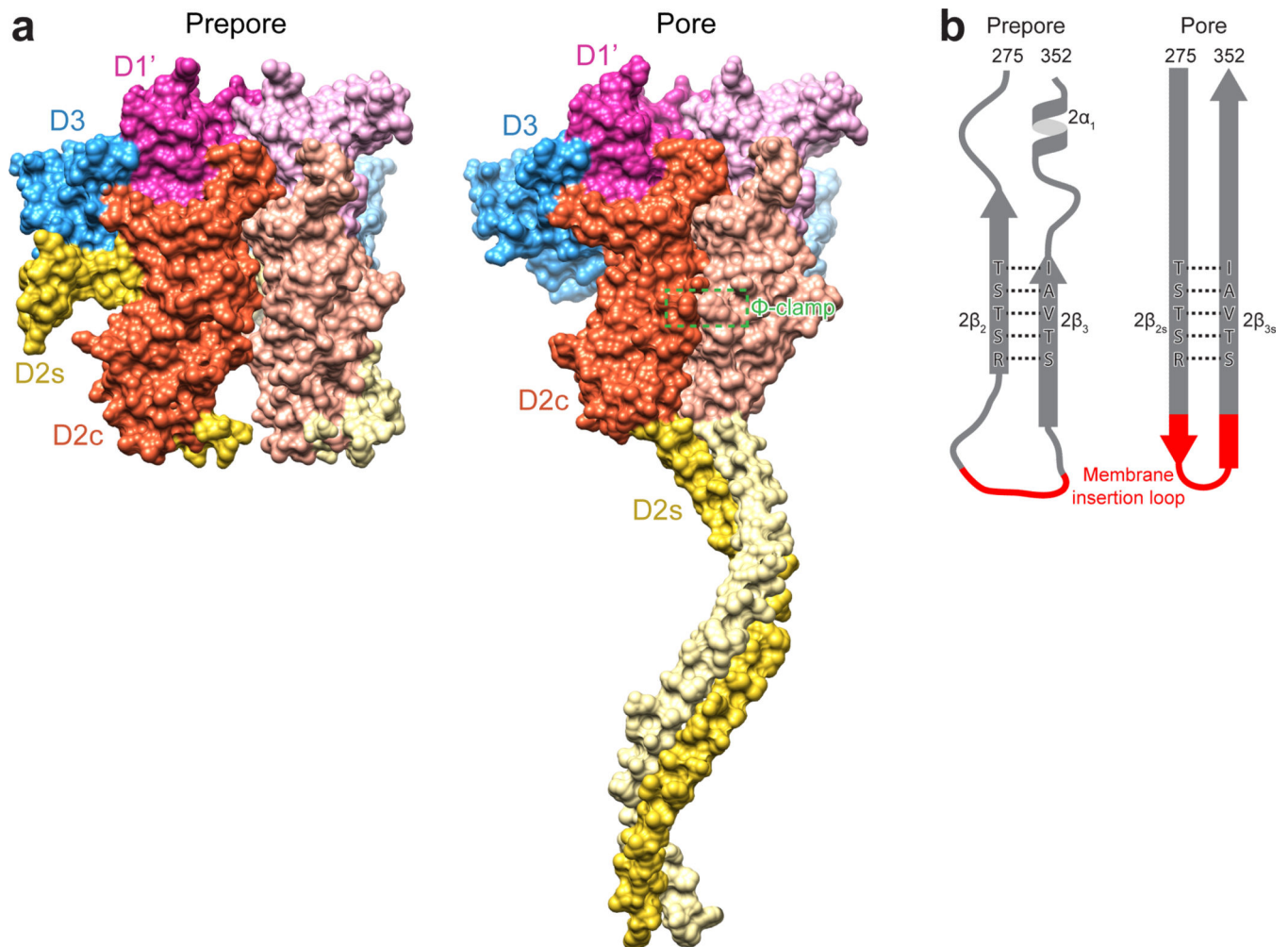
(PDB ID: 1ACC). **b**, Close-up top view of the Φ -clamp region in the PA pore. The $2\beta_7$ - $2\beta_8$ and $2\beta_{10}$ - $2\beta_{11}$ loops are colored in cyan and orange, respectively. The hydrogen bonds between Ser428, Asn399, and Lys397', which form a chain tethering the $2\beta_7$ - $2\beta_8$ and $2\beta_{10}$ - $2\beta_{11}$ loops together, are depicted with dashed lines.



Extended Data Figure 6. Conformational changes of domain 2c between the PA prepore and the PA pore

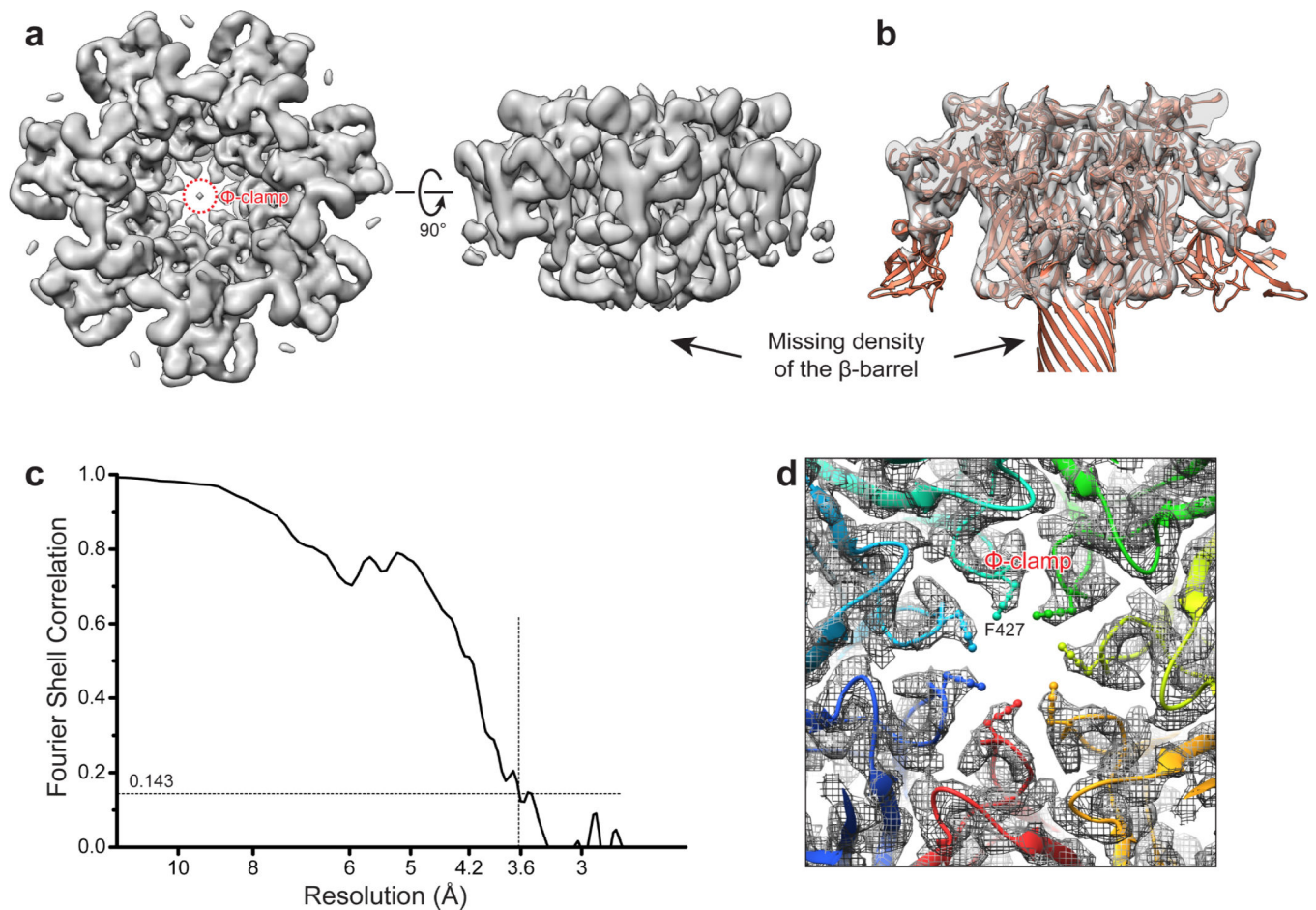
a, Superimposition of domains 2c of the PA pore and the crystal structure of the PA heptameric prepore (PDB ID: 1TZO). The dashed boxes highlight the conformational

difference of the $2\beta_{10}$ – $2\beta_{11}$ loop between these two structures. Residues Asp426 and Phe427 are not solved in the crystal structure of PA prepore due to their flexibility. **b**, Superimposition of domains 2c of the PA pore and the PA octameric prepore (PDB ID: 3HVD). Note the similarity of the $2\beta_{10}$ – $2\beta_{11}$ loop (dashed box) between these two structures. **c**, Superimposition of domains 2c of the PA pore and the PA prepore (monomer; PDB ID: 3TEW). **d–f**, Close-up views of the rearrangements of the $2\beta_{10}$ – $2\beta_{11}$ loop (**d**), the $2\beta_7$ – $2\beta_8$ loop (**e**), and the $2\beta_5$ – $2\beta_6$ and $2\beta_{12}$ – $2\beta_{13}$ loops (**f**) during the conversion from prepore to pore. Val377 and Leu378 of the $2\beta_5$ – $2\beta_6$ loop and Val455 and Tyr456 of the $2\beta_{12}$ – $2\beta_{13}$ loop are flipped upside down to obtain the β strand conformation in the PA pore, leading to an extension of $2\beta_6$ and $2\beta_{12}$ (**f**). CryoEM densities (mesh) corresponding to these loops are displayed to the right of **d–f** with atomic models superimposed, showing unambiguous atomic modeling. The superimpositions of the cryoEM map and the atomic model are shown in views different from the left panels for clarity. The conformational changes of these loops result in a more compact domain 2c with a decrease of 567 \AA^2 in its surface area.



Extended Data Figure 7. Compact assembly of the PA pore

a, Surface views of two neighboring protomers (the same domains in the two protomers are in different shades of the same colors) of the PA pore and the PA prepore (PDB ID: 1TZO) visualized from inside PA heptamer. Domains 1', 2c, 2s, and 3 are colored differently and domain 4 is not shown. The inter-protomer interface in the PA prepore is largely formed by domains 1' and 3 and domain 2 only contributes to this interface by its membrane insertion loop and C-terminal region ($2\beta_{13}$, $2\beta_{14}$, and $2\alpha_3$). In the PA pore, the convergence of domain 2c creates an inter-protomer interface without any gap, with an increase of interface area on domain 2c from $1,247 \text{ \AA}^2$ of the PA prepore to $2,106 \text{ \AA}^2$ of the PA pore as calculated using the PISA⁴⁹ (<http://www.ebi.ac.uk/pdbe/pisa/>). Additionally, formation of the β barrel also leads to extensive contacts, creating a new interface area of $1,195 \text{ \AA}^2$ between two protomers. **b**, Schematic of the conversion of $2\beta_2$, $2\beta_3$, $2\alpha_1$, and the membrane insertion loop of the PA prepore to $2\beta_{2s}$ and $2\beta_{3s}$ of the PA pore. Hydrogen bonds between $2\beta_2$ and $2\beta_3$, which are depicted with dashed lines, are maintained during the conversion. By contrast, $2\alpha_1$ and the connecting loops have to be fully unfolded and converted into β strands that collectively assemble the β barrel of the PA pore. Although the detailed events of β barrel formation are not yet clear, it is likely that assembly starts from the top in a zipper-like manner. A favorable scenario is that the convergence of domains 2c would place the top ends of the β strands close to each other to form a short β barrel, which could extend by pulling more residues together via formation of ordered hydrogen bonds until it reaches the bottom end. It is less favorable that the assembly starts from other regions because disordered hydrogen bonds and hydrophobic interactions could generate enormous non-productive, possibly irreversible pairings between strands.



Extended Data Figure 8. CryoEM single particle reconstruction of a subset of particles lacking the 14-stranded β barrel

a, Surface views of the cryoEM map lacking the β barrel. The map is unsharpened and has a resolution of 3.6 Å. **b**, Cross-section side view of superimposition of the unsharpened cryoEM map with the atomic model of the intact PA pore, showing the cryoEM map has the same conformation as the PA pore except for the absence of the 14-stranded β barrel. **c**, “Gold-standard” FSC (with an auto-mask that was corrected by phase randomization) between two independently refined maps. **d**, Top view of the Φ -clamp region of the cryoEM map (mesh) lacking the β barrel superimposed with the atomic model of the PA pore (ribbons), showing correct assembling of the Φ -clamp in the cryoEM map.

Extended Data Table 1

Data collection and structure refinement statistics.

Data Collection	
Particles	60,455
Pixel size (Å)	1.28
Defocus range (μm)	-1.8 to -5.1
Voltage (kV)	300

Total electron dose	39 e ⁻ /Å ²
Electron dose used in the final reconstruction	30 e ⁻ /Å ²
Refinement	
Resolution (Å)	2.9
Map sharpening B-factor (Å ²)	-95
Average B-factor (Å ²)	81
R.m.s deviations	
Bond lengths (Å ²)	0.009
Bond angles (°)	1.083

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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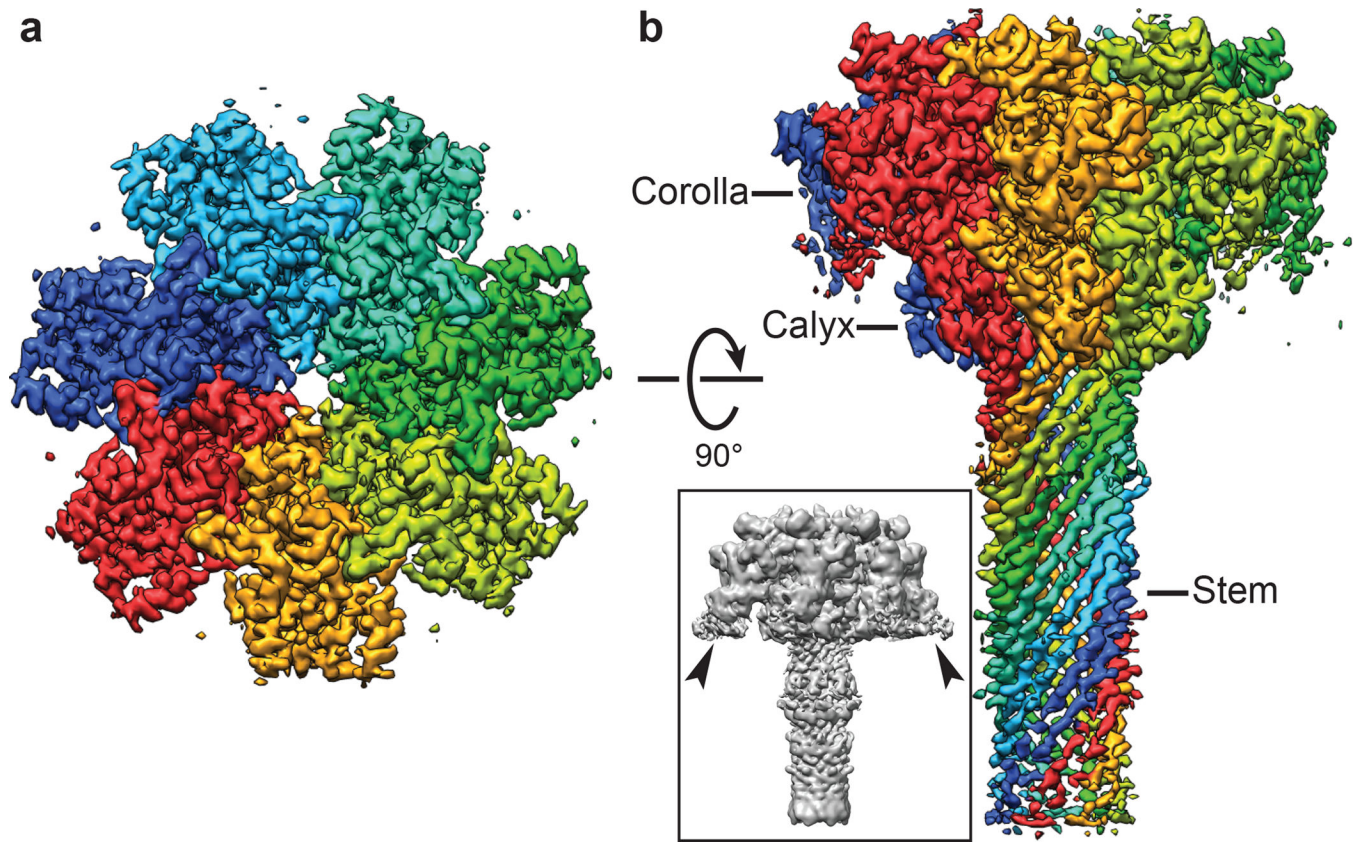


Figure 1. CryoEM reconstruction of the PA pore

a, b, Surface representations of the cryoEM map of the PA pore at 2.9 Å resolution as viewed from the top (**a**) and the side (**b**). Individual protomers of PA heptamer are color-coded. Inset of (**b**) shows the unsharpened cryoEM map in which the flexible domains 4 (arrowheads) are visible.

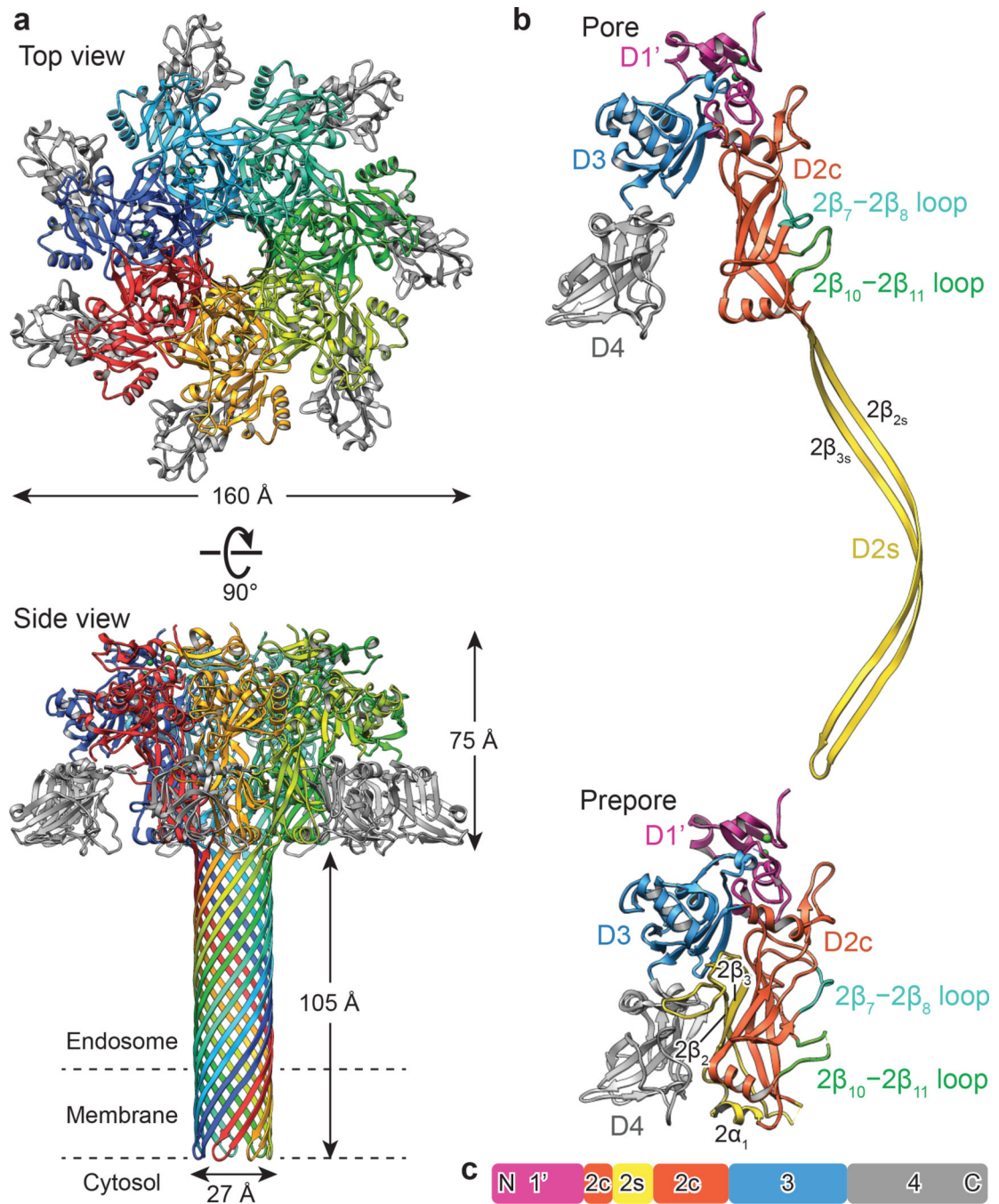


Figure 2. Atomic model of the PA pore

a, Top and side views of the atomic model of the PA pore shown as ribbons. Protomers are color-coded except for domain 4 (grey). **b**, Structural comparison of the protomers of the PA pore and prepore (PDB ID: 1TZO). The domains are colored differently according to **c**. **c**, domain organization of the PA protomer.

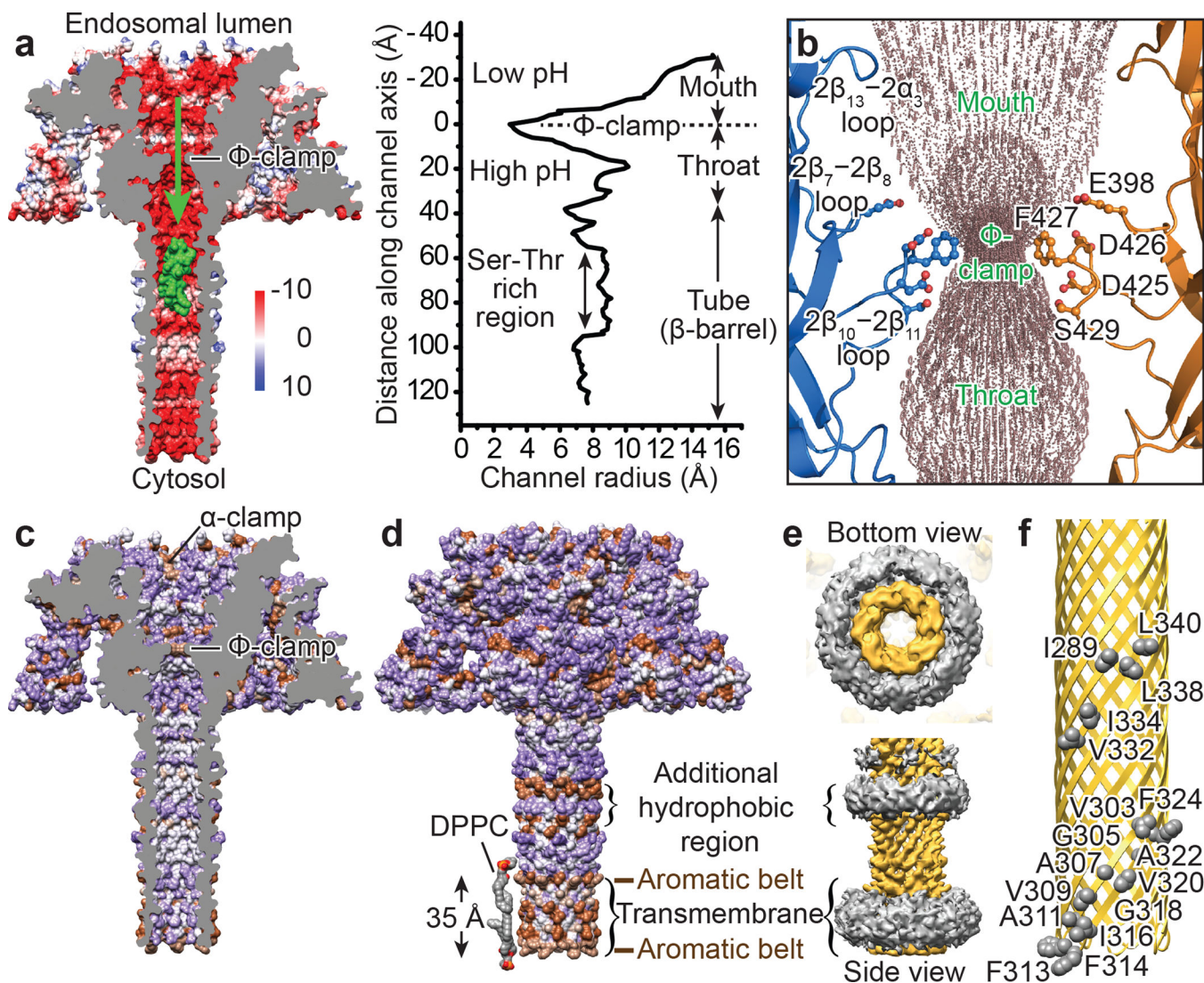


Figure 3. Translocation channel of the PA pore

a, Electrostatic surface (left) and channel radius profile (right; calculated with MOLE¹⁶) of the PA pore. An α helix (green surface model of residues 555–574 of LF; PDB ID: 1J7N) is modeled in the β barrel. The green arrow depicts the direction of protein translocation. **b**, The translocation channel (dots) running through the Φ -clamp. Two protomers of the PA pore are shown as ribbons. Residues E398, D425, D426, F427, and S429 (ball-and-stick) are exposed to the channel. **c**, **d**, Hydrophobicity surface of the PA pore (brown: hydrophobic; purple: hydrophilic; white: neutral). In **c**, the front half of the structure is removed to show the luminal surface. In **d**, two 1,2-dihexadecanoyl-sn-glycero-3-phosphocholine (DPPC) lipid molecules are modeled near the membrane insertion region. **e**, Bottom and side views of the segmented cryoEM map showing the β barrel (yellow) bound with disordered detergent molecules (grey). **f**, The 14-stranded β barrel (ribbons) and the hydrophobic residues (spheres) on its outer surface. The hydrophobic residues are depicted on different protomers for the ease of presentation.

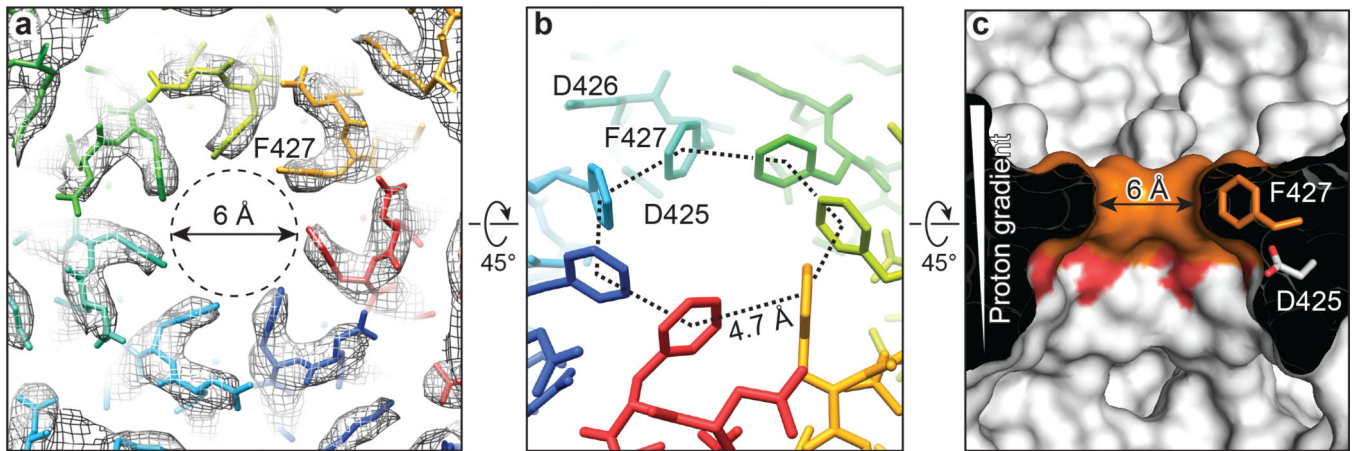


Figure 4. Structure of the Φ -clamp

a, Top view of the Φ -clamp region of the PA pore showing the cryoEM map (mesh) superimposed with the atomic model (stick). **b**, Tilted view of the Φ -clamp with seven protomers colored differently, showing the aromatic CH- π interaction. **c**, Cross-section side view of the translocation channel near the Φ -clamp region. The Φ -clamp (Phe427) and the conserved acidic residue Asp425 are colored in orange and red, respectively.

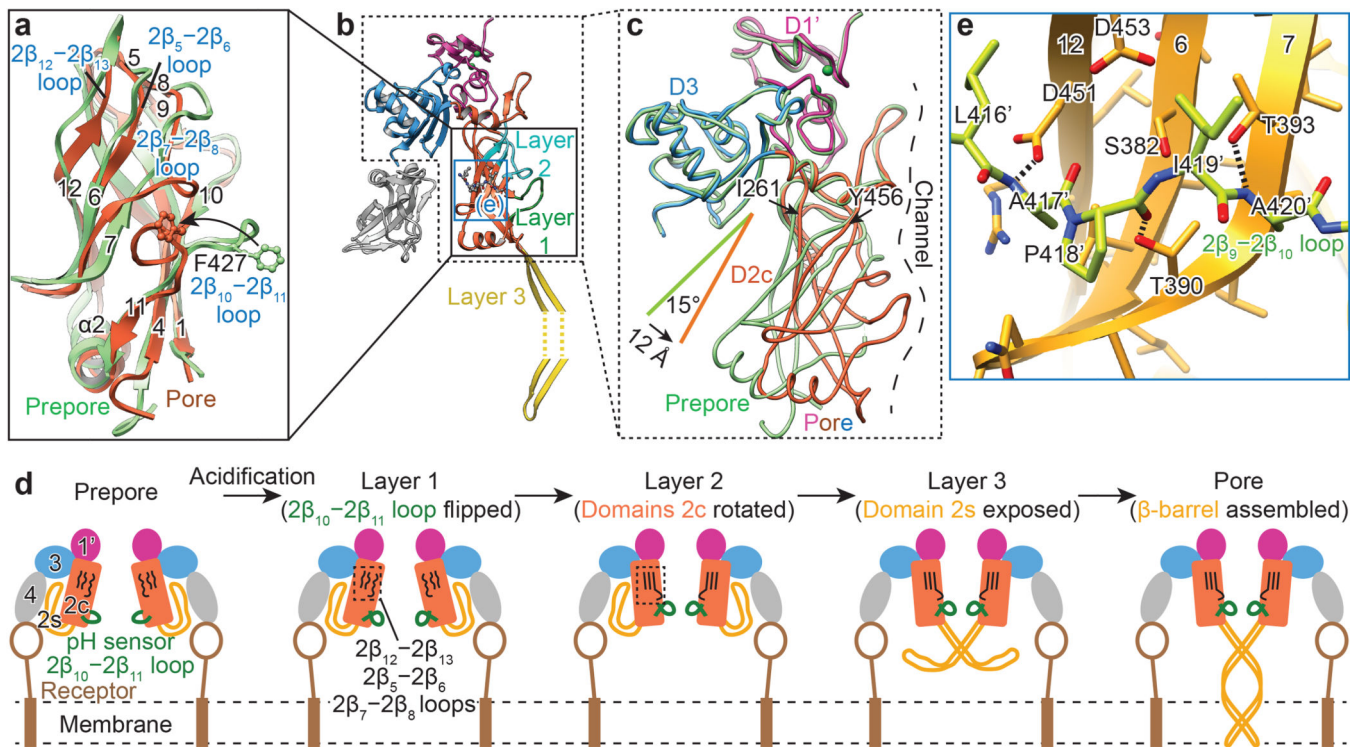


Figure 5. Conversion from prepore to pore

a, Superimposition of domains 2c of the PA pore and prepore (monomer; PDB ID: 3TEW).

Loops undergoing dramatic conformational changes during the prepore to pore conversion

are labeled (blue). **b**, Overview of the three layers of conformational changes during the

conversion. The interaction between the $2\beta_9$ - $2\beta_{10}$ loop and the β sheet (blue box) is shown

in panel **e**. **c**, Superimposition of the PA pore (pink, orange, and blue; colored by domain)

and prepore (light green; PDB ID: 1TZO). Domains 4 and 2s are removed for clarity.

Between the two structures, domain 2c has a C α rmsd=6.87 Å, whereas domains 1' and 3

remain unchanged with C α rmsd=0.54 Å and 0.58 Å, respectively. **d**, Illustration of the steps

of conformational changes during the prepore to pore conversion. For simplicity, only two

protomers are illustrated. **e**, Interface between the $2\beta_9$ - $2\beta_{10}$ loop (green) and the bent β sheet

(gold) of $2\beta_{12}$, $2\beta_6$, and $2\beta_7$ of the adjacent protomer in the PA pore. Hydrogen bonds

between the backbone of the loop and the side chains of the β sheet are depicted with dashed

lines.