Snapsots of actin and tubulin folding inside the TRiC chaperonin

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The integrity of a cell’s proteome depends on correct folding of polypeptides by chaperonins. The chaperonin TCP-1 ring complex (TRiC) acts as obligate folder for >10% of cytosolic proteins, including the cytoskeletal proteins actin and tubulin. Although its architecture and how it recognizes folding substrates are emerging from structural studies, the subsequent fate of substrates inside the TRiC chamber is not defined. We trapped endogenous human TRiC with substrates (actin, tubulin) and cochaperone (PhLP2A) at different folding stages, for structure determination by cryo-EM. The already-folded regions of client proteins are anchored at the chamber wall, positioning unstructured regions toward the central space to achieve their native fold. Substrates engage with different sections of the chamber during the folding cycle, coupled to TRiC open-and-close transitions. Further, the cochaperone PhLP2A modulates folding, acting as a molecular strut between substrate and TRiC chamber. Our structural snapshots piece together an emerging model of client protein folding within TRiC.

The group II chaperonin TRiC (also called chaperonin containing tailess complex polypeptide 1, CCT) is essential for the folding and function of a growing list of proteins driving diverse cellular processes1–4. It is tasked to fold >10% of cytosolic proteins, particularly those with complex domain topology, such as the essential cytoskeletal proteins actin and tubulin. TRiC also suppresses the misfolding and aggregation of neurotoxic proteins, including huntingtin1 and α-synuclein. In addition, it may assist with the assembly of proteins into functional complexes, such as histone deacetylase and viral capsids2. The architecture of TRiC has been the subject of extensive X-ray crystallography and cryo-electron microscopy (cryo-EM) studies3–11. Eight paralogous subunits (CCT1–CCT8) assemble into a hexadecamer of two back-to-back rings, enclosing a folding chamber12. This architecture facilitates nascent protein recognition at the apical domain with a built-in lid, as well as binding and hydrolysis of ATP in the equatorial and intermediate domains to drive closure for open-and-close conformational transition13. The evolutionary divergence to eight paralogous subunits further allows TRiC to fine-tune substrate specificity for the various client proteins, through differential recognition modes involving the subunit apical domains14 and different rates of ATP binding and hydrolysis between subunits15–18. The TRiC subunits also contribute to positive cooperativity within each ring and negative cooperativity between rings, resulting in an ATP-dependent allosteric network19,20 that creates asymmetry in conformations21,22.

By contrast, events that follow entrapment and confinement of nascent proteins in the chamber interior to assist folding into their functional states12 remain inadequate, as atomic details of client protein interactions with the TRiC interior are lacking. The two canonical TRiC substrates, actin and tubulin, are abundant filament-forming cytoskeletal proteins21. Owing to their complex domain topology, these proteins depend on TRiC to achieve their native folds, and inherited mutations in actin and tubulin that disrupt TRiC engagement are associated with human disease (for example, congenital myopathies)24. TRiC-mediated folding of actin and tubulin also requires the assistance of two cochaperone classes, prefoldin (PFD) and phosducin-like proteins (PhLPs). PFD is tasked to bind and stabilize nascent polypeptides emerging from ribosomes, directing them to TRiC to increase efficiency in protein folding25. PhLPs seem to modulate activity in the context of a TRiC–substrate–cochaperone ternary complex26. To obtain insights into the folding of native substrates by TRiC, we set out to isolate endogenous human TRiC for cryo-EM structure determination. Our strategy aimed to entrap endogenous TRiC-bound substrates and cochaperones along the folding cycle, a feat not readily accomplished using reconstituted systems.

Results

Tagging and purifying endogenous TRiC–substrate complexes. We chose to add a purification tag to the CCT5 subunit of endogenous TRiC/CCT. Of all eight subunits, CCT5 was chosen owing to previous success in the expression and purification of recombinant CCT5 with a His-tag at the carboxy terminus27. Using CRISPR–Cas9 knock-in, we inserted a C-terminal 3xFLAG tag into the CCT5 genomic locus of human HEK293T cells and isolated endogenous TRiC using FLAG-affinity chromatography28 (Extended Data Fig. 1). A ~50-kDa protein was found to co-purify with TRiC.
To reveal proteins that co-purify with TRiC complexes, we prepared two sets of samples for in-solution digest, analysed by liquid chromatography (LC) with tandem mass spectrometry (MS) (LC–MS/MS). This includes one set from the CCT5-FLAG cell line and one set from wild-type untransfected HEK293T cells as a reference (Supplementary Data 1). Tubulin and other TRiC-associated proteins were enriched in the CCT5-FLAG samples as compared with the wild-type reference samples, with tubulin constituting the most abundant peptides detected beyond TRiC subunits29 (Extended Data Fig. 2). Other peptides that were present in samples from the CCT5-FLAG cell line, but not the reference samples, are derived from proteins including: PCNA interacting partner (PARPBP), target of rapamycin complex subunit LST8 (MLST8), estradiol 17-beta-dehydrogenase 8 (HSD17B8), actin-related protein 2/3 complex subunit 1B (ARPC1B), cullin-associated NEDD8-dissociated protein 1 (CAND1), cyclic AMP-dependent transcription factor ATF-7 (ATF-7), neural precursor cell expressed developmentally down-regulated protein 8 (NEDD8), denticleless protein homolog (DTL), tubulin-folding cofactor B (TBCB), regulator of MON1-CCZ1 complex (C18orf8), and phosducin-like protein PhLP2A (also known as PDCL3).

| Table 1 | Cryo-EM data collection, refinement and validation statistics |
|----------------|-----------------|-----------------|-----------------|-----------------|
| Data collection and processing | Closed, consensus (EMD-12605) (PDB 7NVL) | Closed, tubulin-bound (EMD-12607) (PDB 7NVN) | Closed, Actin/PhLP2A-bound (EMD-12606) (PDB 7NVM) | Open (EMD-12608) (PDB 7NVO) |
| Magnification | 81,000 | 81,000 | 81,000 | 81,000 |
| Voltage (kV) | 300 | 300 | 300 | 300 |
| Electron exposure (e⁻/Å²) | 62 | 62 | 62 | 62 |
| Defocus range (μm) | 0.75 to 2.5 | 0.75 to 2.5 | 0.75 to 2.5 | 0.75 to 2.5 |
| Pixel size (Å) | 1.06 | 1.06 | 1.06 | 1.06 |
| Symmetry imposed | C₃ | C₁ | C₁ | C₁ |
| Initial particle images (no.) | 3,856,544 | 3,856,544 | 3,856,544 | 3,856,544 |
| Final particle images (no.) | 316,195 | 93,758 | 63,082 | 50,405 |
| Map resolution (Å) | 2.5 | 3.0 | 3.1 | 3.5 |
| FSC threshold | 0.143 | 0.143 | 0.143 | 0.143 |
| Map resolution range (Å) | 2.5–4.4 | 2.8–4.8 | 2.9–6.6 | 3.2–9.9 |
| Map sharpening B factor (Å²) | -63.0 | -65.3 | -57.7 | -70.2 |
| Refinement | | | | |
| Initial model used (PDB code) | 6KS6 | 6KS6 | 6KS6 | 6NRA |
| Model-to-map resolution (Å) | 2.64 | 3.05 | 3.27 | 3.51 |
| FSC threshold | 0.5 | 0.5 | 0.5 | 0.5 |
| Model-to-map correlation (Phenix) | 0.73 | 0.86 | 0.74 | 0.81 |
| Model composition | | | | |
| Non-hydrogen atoms | 66,477 | 69,139 | 71,009 | 34,246 |
| Protein residues | 8626 | 8970 | 9219 | 4480 |
| Ligands | 48 | 48 | 48 | 44 |
| B factors (Å²) | | | | |
| Protein | 72.62 | 72.55 | 70.77 | 64.35 |
| Ligand | 171.45 | 171.45 | 158.28 | 136.15 |
| R.m.s. deviations | | | | |
| Bond lengths (Å) | 0.007 | 0.004 | 0.008 | 0.005 |
| Bond angles (°) | 0.827 | 0.607 | 0.863 | 0.816 |
| Validation | | | | |
| MolProbity score | 1.64 | 1.77 | 1.85 | 2.07 |
| Clash score | 4.63 | 6.80 | 7.32 | 6.81 |
| Poor rotamers (%) | 1.73 | 1.75 | 1.55 | 1.87 |
| Ramachandran plot | | | | |
| Favored (%) | 96.57 | 96.69 | 95.6 | 92.01 |
| Allowed (%) | 3.33 | 3.22 | 4.29 | 7.81 |
| Disallowed (%) | 0.09 | 0.09 | 0.11 | 0.18 |

During sample preparation. To reveal proteins that co-purify with TRiC complexes, we prepared two sets of samples for in-solution digest, analysed by liquid chromatography (LC) with tandem mass spectrometry (MS) (LC–MS/MS). This includes one set from the CCT5-FLAG cell line and one set from wild-type untransfected HEK293T cells as a reference (Supplementary Data 1). Tubulin and other TRiC-associated proteins were enriched in the CCT5-FLAG samples as compared with the wild-type reference samples, with tubulin constituting the most abundant peptides detected beyond TRiC subunits29 (Extended Data Fig. 2). Other peptides that were present in samples from the CCT5-FLAG cell line, but not the reference samples, are derived from proteins including: PCNA interacting partner (PARPBP), target of rapamycin complex subunit LST8 (MLST8), estradiol 17-beta-dehydrogenase 8 (HSD17B8), actin-related protein 2/3 complex subunit 1B (ARPC1B), cullin-associated NEDD8-dissociated protein 1 (CAND1), cyclic AMP-dependent transcription factor ATF-7 (ATF-7), neural precursor cell expressed developmentally down-regulated protein 8 (NEDD8), denticleless protein homolog (DTL), tubulin-folding cofactor B (TBCB), regulator of MON1-CCZ1 complex (C18orf8), and phosducin-like protein PhLP2A (also known as PDCL3).
In our TRiC–Nb18 complex, the Nb18 nanobody is bound at only two locations of TRiC (one per copy of CCT5; Fig. 1a,b). The binding interface consists of complementarity-determining region (CDR) 1, 2, and 3 loops of Nb18 and a hydrophobic patch at the CCT5 equatorial domain, proximal (~15 Å) to the ATP-binding site (Extended Data Fig. 3d). The Nb18–CCT5 interactions are mediated by Phe29, Arg53, and Trp101 from the CDR1, CDR2, and CDR3 loop regions, respectively (Extended Data Fig. 3e). The C7 epitope residues (for example Ile151, Met485, and Pro487) are not conserved among other CCT subunits, suggesting that Nb18 binding is CCT5-specific. In agreement, Nb18 did not bind to recombinant CCT4 in BLI experiments (Extended Data Fig. 3b, inset). CCT5–Nb18 crosslinks were also identified that confirmed the location of nanobody binding to CCT5 (Extended Data Fig. 3f). Our data revealed no detectable presence of homo-oligomeric CCT516 (which is present when CCT5 is produced by recombinant overexpression) in the endogenous isolation from native HEK293 cells, on two bases: first, in all steps of TRiC isolation and purification, the proportion of FLAG-tagged CCT5 remained stoichiometric with other CCT subunits; second, cryo-EM two-dimensional (2D) and three-dimensional (3D) classifications did not present any evidence of CCT5 homo-oligomeric complexes bound with more than two nanobodies per hexadecamer.

The classical TRiC features are illustrated in our structure in atomic details (Extended Data Fig. 6a), including conserved domain motifs and the ATP-binding sites that are occupied here by Mg\(^{2+}\)–ADP–AlF\(_3\) and a water molecule (Fig. 1d and Extended Data Fig. 6b). Importantly, our closed-TRiC model provides unprecedented clarity to intersubunit contacts within rings (cis) and across rings (trans) (Fig. 1f), through three salient features. First, intra-ring cis contacts are mediated by the N-/C-terminal β-strand of one subunit with the β1–β2 hairpin from the adjacent subunit, forming concerted four-stranded β-sheets within each of the two
In the CCT5-FLAG cell line, and 6 of the top 25 most enriched peptides from affinity LC–MS/MS analysis, many isoforms of tubulin were highly enriched we identified the additional density as tubulin (Fig. 2a, left). In our Data Fig. 4) were refined to generate a 3.0-Å reconstruction, where the largely disordered TBD is seen to extend into the cavity cen-

Fig. 2 | TRiC–tubulin complex. a, Side slice map for closed TRiC, showing substrate density within one cavity of the chamber. b, Topology of native β-tubulin (PDB 6I2I). c, Overall model of TRiC–tubulin complex. d, Slice of the CCT8–CCT6–CCT3–CCT1 hemisphere from one ring, in contact with the substrate density (cyan). Inset, three orthogonal views showing region of β-tubulin built into the substrate density (cyan). Unmodeled TBD from tubulin (dark gray) is shown for reference. e, Top-down view of chamber interior, with tubulin density (cyan) adjacent to one hemisphere, and space in the interior that can accommodate the unmodelled TBD. Inset, at lower isosurface threshold, substrate density (gray surface) can accommodate the TBD (dark gray, taken from PDB 6I2I), which extends towards the C termini of CCT1 and CCT2.

rings. The intra-ring sheet formation is fully visualized in our model, and the consequence is a rigid septum mid-level between rings, acting as a barrier that constricts the interior into two largely separate cavities (one per ring) (Fig. 1c). Second, inter-ring interactions are largely maintained at the ring–ring interface by fitting the ‘plug’ (α4–α5 linker) of one subunit, into the ‘socket’ (α14–α15 linker) of the subunit in trans that it stacks with (Fig. 1f and Extended Data Fig. 7c). Third, our model unravels an ordered-coil preceding the N-terminal β-strand of each subunit, reaching out to the equivalent coil of not its stacked subunit, but to the +2 subunit in the anticlockwise direction (Fig. 1f and Extended Data Fig. 7b top). The exception to this N-terminal inter-ring network is CCT4/CCT4, where its N-terminus points to the solvent exterior.

TRiC–tubulin complex in the closed state. The closed-TRiC consensus map revealed additional density within the folding cavity (Fig. 1e and Extended Data Fig. 5), potentially representing a mixture of native nascent substrate proteins. To resolve these additional densities, we performed further 3D classification on the closed particles and simultaneously relaxed twofold symmetry19. Particles in the most abundant class (29.7% total closed particles; Extended Data Fig. 4) were refined to generate a 3.0-Å reconstruction, where we identified the additional density as tubulin (Fig. 2a, left). In our LC–MS/MS analysis, many isoforms of tubulin were highly enriched in the CCT5-FLAG cell line, and 6 of the top 25 most enriched peptides included different forms of β-tubulin (Supplementary Data 1 and Extended Data Fig. 2). Thus, the tubulin density likely reflects a mixture of different tubulin forms, including α-, β-, and γ-tubulin (sharing 31–41% sequence identity), although several isoforms of β-tubulin were among the most enriched peptides from affinity purification. We therefore built our model on the basis of tubulin β-2A chain (TUBB2A), the highest enriched tubulin peptide.

A TRiC–actin–cochaperone complex. Intriguingly, another symmetry-adjacent 3D class (20.0% of all closed particles; Extended Data Fig. 4) presented additional density within both ring cavities
During actin polymerization. As it attains tertiary structure, the subdomain 2 D-loop, a region undergoing disorder-to-order transition, becomes well defined in the cryo-EM density map. These subdomains interact with TRiC through residues that are near-identical among actin isoforms. Subdomain 1 is localized close to CCT3 and CCT4 apical domain (Fig. 3c and Extended Data Fig. 9a–c,g). CCT6, while subdomain 3 extensively contacts the hairpin termini of both ACTB and ACTG1 isoforms of non-muscle actin.

A native actin monomer folds into subdomains 1–4 with an inter-domain ATP-binding cleft. Our model of actin, positioned similarly to tubulin (Figs. 2a and 3a), depicts a partly folded monomer coordinated by specific interactions with the ‘CCT6 hemisphere’ of closed-TRiC state. Subdomains 1 and 3, known as the ‘barbed end’, are well defined in the cryo-EM density map. These subdomains interact with TRiC through residues that are near-identical among actin isoforms. Subdomain 1 is localized close to CCT3 and CCT6, while subdomain 3 extensively contacts the hairpin termini of CCT2, CCT7, and CCT8 (Fig. 3c and Extended Data Fig. 9a–c,g). By contrast, subdomains 2 and 4, known as the ‘pointed end’, are more disordered (Fig. 3b), making less contact to TRiC than the barbed end (Fig. 3c bottom inset). No density is observed for the subdomain 2 D-loop, a region undergoing disorder-to-order transition during actin polymerization. As it attains tertiary structure, the D-loop can extend into a groove formed in the CCT1 intermediate domain (Extended Data Fig. 9d,e). The observed cryo-EM density for subdomain 4 is weak (Fig. 3b), into which we built a C-α trace that revealed minimal TRiC contact with CCT4 apical domain (Fig. 3c and Extended Data Fig. 9f).

Additional density remained beneath actin subdomains 1 and 3 (Fig. 3c). At low isosurface threshold, this is linked, by cylindrical density spanning the inter-ring septum, to the density in the trans cavity (Fig. 3a,c). We hypothesized that these connected densities could represent an actin-binding partner, for example one of the phosducin-like proteins (PhLPs). The human genome encodes phosducin (PDC) and four PhLPs that are grouped into subtype I (phosducin and PhLP1), subtype II (PhLP2A and PhLP2B), and subtype III (PhLP3). Among them, PhLP2A, PhLP2B, and PhLP3 have been shown to regulate TRiC-mediated folding of actin and tubulin. PhLP2A and the closely related PhLP2B were most consistent with the cryo-EM density (Supplementary Discussion). Importantly, PhLP2A, and not PhLP2B, was co-purified with our TRiC sample (Supplementary Fig. 5) and enriched in our LC–MS/MS analysis (Supplementary Data 1 and Extended Data Fig. 2). The PhLP2A sequence comprises an N-terminal helical domain with three predicted helices H1–H3, a central thioredoxin domain (TXND), and a charged C-terminus (Fig. 4a). We therefore identified PhLP2A helix H1, helix H3, and its preceding loop, in addition to TXND, and built their atomic models into the cryo-EM density (Fig. 3c and Supplementary Fig. 2).

The PhLP2A N-terminal helix H1, the most sequence-divergent region among PhLPs, fits into the cis ring density proximal to actin, anchoring the subdomain 1–3 interface (Figs. 3f and 4b and...
Supplementary Discussion). Consequently, actin subdomain 1 has rotated by nearly 40° away from the core, relative to native monomeric actin (Fig. 3d). Helix H1 in the cis ring is connected to the bulk of PhLP2A in the trans ring by weak cylindrical density that can accommodate helix H2. Although not modeled in our structure, helix H2 would traverse >60 Å longitudinally from the cis ring through the septum to the trans cavity, making minimal TRiC contacts (Figs. 3f and 4c). Upon reaching the level of the CCT2′ and CCT3′ apical domains in the trans cavity, helix H2 is followed by a loop region and helix H3, which traverses 35 Å latitudinally to the other hemisphere of this cavity close to CCT6, reaching the C-terminal TXND (Figs. 3f and 4d). This TXND fold, highly conserved across all PhLPs and superimposable with the PhLP2B TXND structure39, contacts TRiC through the CCT5 apical domain loops, and partly through the flanking subunits of CCT2′, CCT7′, and CCT4′ (Fig. 3f). The PhLP2A 22-aa C terminus was disordered loops, and partly through the flanking subunits of CCT2, CCT7, and CCT4 (Fig. 3f). These generated an open-state consensus map at 3.5-Å resolution (Fig. 6a), ranging locally from 3.5 Å (equatorial domains) to 9.0 Å (apical domains) (Extended Data Fig. 5). We built into the ordered map portion all equatorial domains up to the ATP-binding pockets (Extended Data Fig. 6c), which we again modeled with Mg2+-ADP-AlF6, although the ligand density for CCT4 and CCT5 subunits is less well featured. We then used an open-state map filtered to 4.0-Å resolution to model apical domains to the less-ordered map portion by molecular dynamics flexible fitting. This open TRiC conformation (Fig. 6b), previously seen with AMPPNP-bound yeast CCT, substrate-bound bovine CCT, and substrate MLST8-bound human CCT48, is the consequence of concerted rigid-body rearrangement of all (apical, intermediate, equatorial) domains, relative to closed-TRiC, to varying degrees among different subunits (Extended Data Fig. 7a). Such open-and-closed transitions impart major consequences on intersubunit as well as TRiC–substrate contacts.

In open TRiC, a substantial portion of the N terminus is disordered in most subunits (Extended Data Fig. 7b bottom). Therefore, unlike closed-TRiC, the pre-strand coil is not available for inter-ring contacts and the N-terminal β-strand is missing from intra-ring sheet formation (which is now three-stranded). Additionally, the inter-ring plug-socket interactions seen in closed-TRiC (Extended Data Fig. 7c) have now disengaged in open TRiC with a rearranged interface (Extended Data Fig. 7d).

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Substrates bind differently to TRiC in the open state. Beyond the closed states, initial 2D classification revealed TRiC in the open state (open TRiC) for 10.7% of particles (Extended Data Fig. 4). These generated an open-state consensus map at 3.5-Å resolution (Fig. 6a), ranging locally from 3.5 Å (equatorial domains) to 9.0 Å (apical domains) (Extended Data Fig. 5). We built into the ordered map portion all equatorial domains up to the ATP-binding pockets (Extended Data Fig. 6c), which we again modeled with Mg2+-ADP-AlF6, although the ligand density for CCT4 and CCT5 subunits is less well featured. We then used an open-state map filtered to 4.0-Å resolution to model apical domains to the less-ordered map portion by molecular dynamics flexible fitting. This open TRiC conformation (Fig. 6b), previously seen with AMPPNP-bound yeast CCT, substrate-bound bovine CCT, and substrate MLST8-bound human CCT48, is the consequence of concerted rigid-body rearrangement of all (apical, intermediate, equatorial) domains, relative to closed-TRiC, to varying degrees among different subunits (Extended Data Fig. 7a). Such open-and-closed transitions impart major consequences on intersubunit as well as TRiC–substrate contacts.

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The rearranged intra-/intersubunit contacts in open TRiC indicate that the septum structure at the ring–ring interface has become more dynamic and less constricted between the two cavities when compared with closed-TRiC (Fig. 6c). Indeed, our open TRiC map revealed weak substrate density at the level of the equatorial domains, surrounded by the less rigid septum (Fig. 6d). This density is positioned >30 Å deeper into the chamber when compared with the actin and tubulin positions in closed-TRiC, occupying the same site as MLST8 substrate in the human open TRiC structure42. We reason that this density represents a summed average of different substrates sampled in the data set and cannot be attributed to a single substrate. This density is in close proximity with CCT7 subunits from both rings (Fig. 5d) while engaging more transiently the dynamic N and C termini of other subunits from both rings.

Discussion
Nearly two decades have elapsed since we caught the first glimpse of a type II chaperonin, initially through the archaeal thermosome, and subsequently the yeast and mammalian TRiC12. More recent structural studies have provided clarity on how the PFD cochaperone loaded with client polypeptides latches onto the TRiC apical domains25, how sequence motifs from diverse substrates are recognized by apical domains through specific multivalent subunit contacts14, and how differential rates of ATP binding and hydrolysis across TRiC subunits turbo-charge the chaperonin conformations34. Therefore, a mechanistic understanding has emerged: nascent unfolded polypeptides, such as actin and tubulin, are recognized at the surface-exposed apical domain of nucleotide-free TRiC43,44 and are released into the chamber by the ATP-induced lid formation45.

Fig. 5 | Crosslinking mass spectrometry analysis of TRiC-substrate interactions within the folding cavity. a, b. Crosslinks between TRiC and tubulin (a) and between TRiC and actin/PhLP2A (b) reported from literature are mapped onto the TRiC structure with bound proteins. Left, identified crosslinks compatible with our structural models are indicated in green lines; crosslinks that are not sterically compatible with our model are indicated in yellow lines and could represent other stages of the substrate folding cycle that are not captured in our structures. Spheres represent lysine residues that are crosslinked. Right, magnified views showing the atomic environment of 3 TRiC–tubulin, 1 actin-PhLP2A, and 1 TRiC-PhLP2A crosslinks that support our models.
The subsequent events underlying the substrates as they become folded inside the TRiC chamber remain less defined. Biophysical and structural characterization of substrate-bound TRiC is technically challenging, if one relies on co-expression or reconstitution of constituent proteins via a recombinant host. The alternative is to study endogenous TRiC samples, with early studies involving bovine and mouse testes; this has more recently been mediated by pull-down via an overexpressed protein component from the expression host (insect, yeast, and HEK293 cells). Here, we transform the strategy of endogenous isolation without involvement of an overexpressed component. Using CRISPR knock-in technology to introduce purification tags in TRiC, we have captured TRiC bound with abundant proteins in the cell, during their acts of folding. Additionally the use of a subunit-specific nanobody as a structural biology tool and improved 3D image classification for cryo-EM data have enhanced the resolution beyond recent TRiC structures.

Our series of substrate-bound TRiC structures have reinforced the current concept that, although TRiC adopts a double-ring architecture forming two largely separated cavities, only one client substrate is bound at any one time, in support of the proposed inter-ring negative cooperativity and asymmetry between the two rings. This concept is further extended by our class of open TRiC particles, which reveals substrate density at the inter-ring septum around the ‘bottom’ of both cavities, formed by the partly disordered N- and C-termini in the equatorial domains of all subunits. Proposed by early studies to form a barrier separating the two cavities, the septum has recently been shown to be the binding site for the substrate MLST8 (ref. 42). Although the identity/identities of our substrate density are not resolved, the inter-ring septum clearly plays a role in holding a substrate protein in the open state, and the less well featured density may imply that proteins held here have only adopted some degree of structures, likely at the early folding stages.

We opted to incubate the TRiC sample with ADP–AlFx, in order to mediate a closed state with double-ring closure, reasoning that this offers the best opportunity for atomic-level data detailing TRiC-substrate interactions. Since this state has not been observed in other ATP analogs, its physiological importance remains to be determined. Nevertheless, our closed-TRiC models reveal that tubulin and actin have now attained near-native fold and are confined in one of the two chamber cavities. For both proteins, the primary contacts with TRiC are through the CCT3/6/8/7 subunits, referred to as the TRiC hemisphere with low ATP binding and hydrolysis.
rates (relative to the CCT1/4/2/5 hemisphere). Our observations and placement of substrate proteins inside one chamber cavity, at the level of CCT3, CCT6, and CCT8 apical domains, are in overall agreement with residual densities identified in the yeast TRiC–actin structure of the closed state and the bovine TRIC–tubulin structure of the open state. Our structural information has unleashed a molecular basis for TRIC-mediated folding: subunit-specific contacts stabilize the already structured regions of substrate proteins (such as actin subdomains 1/3 and tubulin N and C domains), allowing the less folded/disordered regions (such as actin subdomains 2/4 and tubulin TBD) to achieve native structure, presumably by using chamber space and additional subunit contacts if necessary. The disordered TBD in our tubulin model could therefore well explain the smaller-than-expected substrate density inside the bovine TRIC structure.

Furthermore, our TRIC–actin–PhLP2A model reveals for the first time the structure of full-length PhLP2A, belonging to the conserved family of thioredoxin-fold phosducin-like proteins. The yeast ortholog Ppl2p is involved in actin biogenesis and stimulates actin folding by thirtifold in vitro. Our structure suggests how the PhLP2A cochaperone plays two roles in TRIC quality control, via extensive helical segments encoded in its amino acid sequence. By holding the substrate within closed-TRIC (as a strut) until a folded state is reached for chamber release (as a sensor), the cochaperone therefore prevents premature chamber opening, prolongs actin residence time in the chamber, and minimizes abortive folding. This is of particular importance for the folding cycle of actin intermediates, which have an otherwise high dissociation rate from TRIC. In our structure, PhLP2A is fully localized inside the chamber, contrary to the localization at the tip of apical domains by its paralog, revealed in the 18-Å EM reconstruction of insect TRIC overexpressed with human PhLP1 and client Gβ. It remains to be determined whether PhLP1, harboring a 70-aa-longer N terminus and a distinct set of clients compared with PhLP2A, could engage with TRIC differently.

In summary, we speculate a working model for TRIC (Fig. 6e), whereby the client substrate could first engage the open apical domains and then traverse different sections of TRIC depending on its folding status, and in coordination with an ATP-driven conformational cascade that surveys asymmetry and multiple states within the chamber. For example, in the open TRIC, the septum could act as the reception point, loosely holding nascent or partly folded substrates upon their entry into the TRIC chamber. ATP binding and hydrolysis then ensue, possibly causing long-range conformational changes from lid closure at the apical domains all the way to rearrangement of subunit termini at the septum. Through this, a combination of near-native structure and its steric hindrance at the rearranged septum could help translocate the substrate to the ‘top’ of the chamber, readying for the final folding stages. At this juncture, cochaperones such as PhLP2A could ‘step in’ and hold the substrate for as long as it is needed inside the chamber. This work sets the stage for future studies aimed at verifying our model of multiple binding modes for the same/different substrates inside TRiC, in order to piece together the inner workings of this nanomachine.

Online content
Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41594-022-00755-1.

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Methods

No statistical methods were used to predetermine sample size. The experiments were not randomized. The investigators were not blinded to allocation during experiments and outcome assessment.

Tagging of endogenous TRIC/CCT with CRISPR-Cas9. To insert a purification tag to the endogenous TRIC/CCT, HEK293T cells were transfected with plasmid encoding SpCas9(1.1) and sgRNAs targeting CCT5 and ATP1A1 (modified from Addgene no. 66113). These cells were also transfected with linear dsDNAs to act as HDR templates by which to insert 3xFLAG/SpTag at the C terminus of CCT5, as well as the Q18R and N129D ouabain resistance conferring point mutations to ATP1A1.[55] Following ouabain selection, the polyclonal cell pool was assessed for FLAG-tagged CCT5 via immunoblotting. Single cells from confirmed pools were seeded into 96-well plates by FACS, and resulting monoclonal lines were again screened via In-Cell Western Assay (LI-COR) followed by western blots. Three positive monoclonal lines were identified and each was further verified by PCR, sequencing, and western blots.

Llama immunization, nanobody selection, and nanobody purification. We raised CCT5-specific nanobodies by immunization of a llama,[51] using as antigen the recombinant Escherichia coli-expressed human CCT5, which was previously shown to exist as a TRIC-like homo-hexadecamer (CCT5,[52]). The llama received 6 injections of the antigen over a span of 6 weeks, after which the ORF of nanobodies were isolated by extracting blood, separating out peripheral blood lymphocytes, and isolating RNA for use in creating cDNA libraries. In vitro selection of nanobodies was performed using single display, Online Score (a) was evaluated by evaluated by enzyme-linked immunoblot assay (ELISA), detecting presence of nanobody in the sample. Twenty-eight nanobodies from 18 families were identified, among which nanobody Nb18 was selected for cryo-EM experiments on the basis of characterization by pull-down and biolayer interferometry studies (Extended Data Fig. 3). Nb18 (ref. CA14679) is constructed in a pM5Sv vector incorporating C-terminal His, and CaptureSelect C-tag (or EPEA-tag), which allows recombinant expression in E. coli. and purification by Ni-NTA affinity and size-exclusion chromatography.

Isolation and purification of human TRIC. Monoclonal HEK293T cells that were confirmed to contain FLAG-tagged CCT5 were grown in suspension and collected by freeze–thaw of cell pellet and were resuspended in lysis buffer (100 mM HEPES pH 7.5, 100 mM NaCl, protease inhibitors, 1 mM DTT, benzamidine, and 1% Triton X-100) and incubated at 4 °C for one hour. Lysate was centrifuged at 35,000g, and supernatant was passed through a 0.4-μm filter and was incubated at 4 °C for 1 hour with 5 ml of anti-FLAG resin (Anti-DYKDDDK Tag (LS) Affinity Gel Protocol BioLegend (cat no. 651503). Resin/lysate mixture was passed through a gravity column, washed with 100 CV of wash buffer (20 mM HEPES pH 7.5, 100 mM NaCl) and eluted with wash buffer containing 0.15 mg/ml 3x FLAG peptide (Sigma cat. no. F4799) (Extended Data Fig. 1b). The eluates were concentrated and run through a Superose 6 10/30 size-exclusion chromatography column (Extended Data Fig. 1c,d). Fractions containing purified TRIC were concentrated and flash frozen and stored at −80°C.

Cryo-EM structure determination. Purified TRIC/CCT complex (1.5 mg/ml) was premixed with Nb18 at a 1:1 molar ratio, with 1 mM ATP, 5 mM MgCl2, and ALF (5 mM Al(NO3)3 and 30 mM NaF) A 3-μl aliquot was pipetted on a holey carbon cryo-EtcoCar (1.3 nm mesh). Quantitative determination in liquid ethane (Vitrobot Mark IV, Thermo Fisher Scientific). The data were collected using a 300-kV transmission electron microscope (Titan Krios, TES) on a direct electron detector (Gatan K3). Data collection parameters are in Table 1. Cryo-EM data were processed in Scipion software framework.[56] Movie frames were aligned, dose-weighted and averaged using MotionCor2 (ref. 57).

Covalent crosslinking (CCT) parameter estimation was using using CTFind4 (ref. 58). Particles were picked using cryYLO and subjected to several rounds of classification in RELION.[59–61] The particles were classified to closed and open states and actin (PDB 2Q1N) structures were fit in a similar way to the 3.0-Å resolution real space cross-correlation in UCSF Chimera. The tubulin (PDB 6L2I chain B) and actin (PDB 2Q1N) structures were fit in a similar way to the 3.0-Å resolution closed map. Models were manually adjusted in COOT and subjected to several rounds of real space refinement in PHENIX. To model the open state, the previously solved open state (PDB 6NRA) model was first fitted as a rigid body to the 3.5-Å resolution cryo-EM map of the open state and then manually added to the 2.1-Å resolution map built in COOT. Molecular dynamics flexible fitting was performed using Flex-EM software[62] to create a open model containing apical domains that roughly fit the open state map filtered to 4.0-Å reconstruction of the open state. Model refinement and validation statistics are in Table 1. Structure figures were visualized and processed using UCSF ChimeraX.[63]

Tryptic digest mass spectrometry (MSMS) and side-chain modeling confirmed that the predominant copy of CCT6 in the TRIC complex isolated from HEK293T is the ubiquitously expressed CCT6A isoform, and not the alternatively spliced isoform CCT6B.[64] The predominant peptide coverage in the MSMS data was the ubiquitously expressed CCT6A isoform, and not the alternatively spliced isoform CCT6B.[64] The predominant peptide coverage in the MSMS data was the ubiquitously expressed CCT6A isoform, and not the alternatively spliced isoform CCT6B.[64] The predominant peptide coverage in the MSMS data was the ubiquitously expressed CCT6A isoform, and not the alternatively spliced isoform CCT6B.[64]

Binding and activity assays. Biolayer interferometry (BLI) experiments were performed on a 16-channel ForteBio Octet RED384 instrument at 25 °C, in buffer containing 20 mM HEPES pH 7.5, 150 mM NaCl, and 0.1% BSA. Fifty microliters of 100 ng/µl biotinylated CCT5 was loaded to the streptavidin coated sensors. The concentration used for Nb18 ranged from 40 nM to 39 nM. Measurements were performed using a 300-second association step followed by an enzyme-linked dissociation step on a black 384-well plate with tilted bottom (ForteBio) using a serial dip method from low to high Nb concentration. The baseline was stabilized for 30 seconds prior to association, and signal from the reference sensors was subtracted and steady-state kinetics were fit using Octet Data Analysis software.

Malachite green ATPase activity assays were performed by incubating protein samples in ATPase assay buffer (20 mM HEPES pH 7.5, 100 mM NaCl) with 0.1% BSA. and purifying using a 1:1 molar ratio, with 1 mM ATP, 5 mM MgCl2, and AlFx (5 mM Al(NO3)3 and 30 mM NaF) A 3-µl aliquot was pipetted on a holey carbon cryo-EtcoCar (1.3 nm mesh). Quantitative determination in liquid ethane (Vitrobot Mark IV, Thermo Fisher Scientific). The data were collected using a 300-kV transmission electron microscope (Titan Krios, TES) on a direct electron detector (Gatan K3). Data collection parameters are in Table 1. Cryo-EM data were processed in Scipion software framework.[56] Movie frames were aligned, dose-weighted and averaged using MotionCor2 (ref. 57).

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agitation, and the reaction was quenched with 50 mM NH₄HCO₃, (1:20 dilution from 1 M stock, or 5 μL) for 20 min at 25°C.

XL-MS data generated in this study were analyzed using XlinkX in Proteome Discoverer (Thermo Scientific) at a 5% FDR. XL-MS data were filtered with a 10 ppm tolerance. The results were filtered with an E-value of 0.01 or less. For crosslinks to TRiC substrates, all identified crosslinked peptide–pairs were considered.

Reporting Summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

Data sets generated during the current study are available from the Protein Data Bank (PDB) accession codes 7NVL, 7NVN, 7NVN, and 7NVO, and Electron Microscopy Data Bank (EMDB) accession codes EMD-12605, EMD-12606, EMD-12607, EMD-12608, and EMD-13754. All main data supporting the findings of this study are available in this article. Extended Data, and Supplementary Information. Source data are provided with this paper. Other data are available from the corresponding author upon reasonable request. Source data are provided with this paper.

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Author contributions

C.B., J.T.H. and W.W.Y. conceived of the study. J.T.H., K.M.K., J.M.J., J.H. and W.W.Y. supervised the project. D.T., V.O.P., and J.J.K. performed genome editing. E.P., J.J.K., and J.S. carried out nanobody generation and characterization. H.K. and L.H. carried out LC-MS/MS and XL-MS analysis. J.K., G.C., and J.T.H. performed cryo-EM and model building. J.K., J.T.H., and W.W.Y. wrote the manuscript with contributions and critical feedback from all the authors.

Competing interests

K.M.K., J.M.J., and C.B. are employees of Pfizer Inc. J.K., D.T., E.P., G.C., H.K., L.H., J.S., V.O.P., J.T.H., and W.W.Y. declare no competing interests.

Additional information

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Extended Data Fig. 1 | Selection and large-scale purification of monoclonal CCT5 C-term edited cell lines. (a) Anti-FLAG western blot (top) and stain-free imaging (bottom) of wild type HEK293T, mixed pool, and five monoclonal cell lines. (b) Size exclusion profile of large-scale TRiC purification with pooled fractions indicated by pink shading. (c) SDS-PAGE of fractions from size exclusion chromatography showing double-banding patterns of TRiC subunits, and tubulin co-elution with TRiC. (d) Anti-CCT western blots (left) and Coomassie visualisation (right) of monoclonal 3.B5 line showing presence of TRiC subunits. Coomassie gel bands were excised, digested with trypsin and analyzed using LC-MS/MS to confirm presence of TRiC subunits. (e) Negative stain micrograph showing well-defined TRiC particles (data representative of n = 1). The Western Blot experiments in a,d were carried out at least two times. Uncropped images and unprocessed scans in a,c,e are available as source data.
Extended Data Fig. 2 | Volcano plot of peptides identified in solution digest LC-MS/MS. Volcano plots are displayed with the log(2)-fold change on the x-axis and the -log(10)-p-value, as calculated by Perseus software using a two-sided statistical T-test, on the y-axis. Confidently enriched interactors in the endogenous TRiC sample, defined as p < 0.01 and log-fold change > 1.5, are annotated with protein name including actin (red dots), tubulin (blue dots) and PhLP2A (shown as PDCL3, black dot). ACTA1 denotes peptide identification that could not distinguish between α-cardiac muscle 1 ACTC1, aortic smooth muscle ACTA2, γ-enteric smooth muscle ACTG2, and cytoplasmic 2 isoforms ACTB (shown in the figure as ACTA1A). Raw data are found in Supplementary Data 1.
Extended Data Fig. 3 | See next page for caption.
Extended Data Fig. 3 | Nanobody Nb18 characterisation. (a) Nb18 (red box) pulls down overexpressed CCT5 (left, n = 2) and endogenous TRiC (right, n = 1) from affinity chromatography. (b) Nb18 binds recombinant CCT5 in biolayer interferometry (Kd = 86 nM). Inset: Raw sensogram showing that recombinant CCT4 does not bind Nb18. (c) CCT5 and TRiC ATPase activity in the presence of Nb18. Nb18 was added at 5:8 molar ratio to recombinant CCT5, and at 1:1 to endogenous TRiC. Data are presented as mean values +/- SD (n = 6 for CCT5 only, n = 3 for all other samples). (d) Nb18 binds to a hydrophobic interface of CCT5 in proximity to ATP binding pocket. (e) Interacting residues between Nb18 and CCT5 in closed-TRiC (left) and open-TRiC (right). Green dashed lines indicate van der Waals contacts and blue dashed lines indicate hydrogen bonds. (f) Structural representation of CCT5-Nb18 crosslinks. Crosslinked residues are highlighted as spheres in gold. Dotted lines indicate distances between amines. Uncropped gels in a are available as source data.
Extended Data Fig. 4 | EM data processing. (a) A representative micrograph of purified TRiC complexes (n = 1). (b) Cryo-EM data processing workflow. During the processing steps, the data were split in four sets (Set 1 - Set 4) which were processed in parallel for computational efficiency. Uncropped image in a is available as source data.
Extended Data Fig. 5 | EM map resolutions. Angular distribution and local resolution of all maps.
Extended Data Fig. 6. | Conserved structural features of TRiC. (a) Conserved structural motifs common to all CCT subunits (here represented by CCT1) including apical lid helices, proximal loop, release loop of substrate (RLS), P-loop, nucleotide-sensing loop (NSL), and stem loop. Insets: Motifs are coloured orange and density map is shown grey. (b,c) Snapshots of ligand density in the nucleotide binding site for each CCT subunit of closed-TRiC (b) and open-TRiC (c). Dashed lines represent hydrogen bonding. ADP molecules are shown in yellow with Mg$^{2+}$ in green and water molecules are represented in red spheres.
Extended Data Fig. 7 | Intra- and Inter-ring contacts. (a) Flattened surface representation showing conformational differences between closed-TRiC and open-TRiC in the equatorial (E), intermediate (I) and apical (A) domains. (b) Views of the network of N-terminal extensions between subunits across the ring interface in closed-TRiC state (top). Note that these interactions were not observed in open-TRiC state (bottom) due to disorder of the N-terminal extensions. (c,d) Flat rendering of the CCT2-CCT2′ inter-ring stacking in closed-TRiC (c) and open-TRiC (d). Compared to closed-TRiC, rearranged helix α15 pushes loop α4-α5 from the trans subunit away from the interface. Inter-ring contacts are now mediated between the entire helical face of α15, and loop α13-α14 from the trans subunit.
Extended Data Fig. 8 | See next page for caption.
Extended Data Fig. 8 | Tubulin-TRiC interactions. For all panels, tubulin is coloured by domain: N-terminal domain (blue), TBD (crimson), and C-terminal domain (dark green). Green dashes indicate van der Waals contacts; blue dashes indicate hydrogen bonds. TRiC structural features labelled are apical domain loop (AL), stem-loop (SL), C-terminus (CT) and helix X (αX). (a) CCT8 interacts with tubulin N-terminus (Arg2, Glu53, Asp74) using its stem-loop (His59, Leu60) and intermediate domain (Arg314, Asn316). (b) CCT6 contacts tubulin N-terminal α4 (Glu125) and α5 (Glu157, Glu158, Tyr159) through a β-sheet of the apical domain. Additionally, CCT6 stem-loop (Ala51) and C-terminus (Met526) interact with tubulin N- and C-termini, along with CCT3 stem-loop (Met54). (c) Tubulin C-terminus (Phe402, Trp405) interacts with CCT3 stem-loop, helix α3 from equatorial domain (Glu89), and intermediate domain. (d) CCT1 apical domain loop (Lys245) and stem-loop (Asp358) form few tubulin contacts (Phe260, Pro261), involving also CCT3 apical domain loop and helix α3.
Extended Data Fig. 9 | Actin-TRiC interactions. *For all panels,* actin is coloured by domain: subdomain 1 (S1, blue), subdomain 2 (S2, crimson), subdomain 3 (S3, dark green), and subdomain 4 (S4, yellow); green dashes indicate van der Waals contacts; blue dashes indicate hydrogen bonds; TRiC structural features labelled are apical domain loop (AL), stem-loop (SL) and C-terminus (CT). (a) CCT8 uses stem-loop and C-terminus to interact with actin subdomains 1 and 3. This network of interactions also involves CCT7 C-terminus. (b) CCT6 uses the β-sheet in apical domain to contact exclusively actin subdomain 1. (c) CCT3 uses its stem-loop and apical domain β-sheet to contact actin subdomain 1. (d,e) CCT1 makes large number of actin contacts, using a groove in the intermediate domain to interact with the partially disordered D-loop of actin subdomain 2 which plays a role in ATP binding. (f) CCT4 I304 and L305 make few contacts with actin subdomain 4 which is not well ordered in our map. (g) CCT7 uses the stem-loop and C-terminus to interact with actin subdomain 3.
Extended Data Fig. 10 | Intra-TRiC and inter-protein crosslinking mass spectrometry corroborates TRiC subunit orientation and protein localization. 

(a) Ten most prevalent mappable crosslinks between TRiC subunits are overlayed onto our model of closed-TRiC. (b) All mappable TRiC-tubulin crosslinks are overlayed onto our model of closed-TRiC complexed with tubulin. (c) All mappable TRiC-actin-PhLP2A crosslinks are overlayed onto our model of closed-TRiC structure complexed with actin and PhLP2A. For all panels, crosslinks indicated as red lines map to the external surface of the chaperonin and are unlikely to represent interactions relevant to the study. Crosslinks indicated in yellow lines are not compatible with the localization or orientations suggested by our model, while those indicated in green lines are.
Reporting Summary

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Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

- **n/a** Confirmed
- [ ] The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
- [ ] A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- [ ] The statistical test(s) used AND whether they are one- or two-sided
  - *Only common tests should be described solely by name; describe more complex techniques in the Methods section.*
- [ ] A description of all covariates tested
- [ ] A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
- [ ] A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
- [ ] For null hypothesis testing, the test statistic (e.g. F, t, r) with confidence intervals, effect sizes, degrees of freedom and P value noted
  - *Give P values as exact values whenever suitable.*
- [ ] For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
- [ ] For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
- [ ] Estimates of effect sizes (e.g. Cohen’s d, Pearson’s r), indicating how they were calculated

*Our web collection on [statistics for biologists](https://www.nature.com/article/statistics-for-biologists) contains articles on many of the points above.*

Software and code

**Policy information about availability of computer code**

**Data collection** DLS beamline Krios IV, CLARIOstar microplate reader (BMG Labtech), ForteBio Octet RED384, LTQ Orbitrap Velos

**Data analysis** Scipion v3.07, RELION v3.1, GraphPad Prism v8.4.3, COOT v0.9.7, SWISS-MODEL, MotionCor2 v1.4.0, CTFFind4 v4.1.9, cryolo v1.7.6, UCSF Chimera v1.16, UCSF Chimera X v1.3, PHENIX v1.17.1-3660, Octet Data Analysis v11.0, MaxQuant v1.6.2.3, Proteome Discoverer v2.4, FlexEM v2.6.10986.0, Perseus v1.6.2.2

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Data

**Policy information about availability of data**

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

Datasets generated during the current study are available from the Protein Data Bank (PDB) accession codes 7NVL, 7NVN, 7NVO, and Electron Microscopy Data Bank (EMDB) accession codes 12605, 12606, 12607, 12608 and 13754. All main data supporting the findings of this study are available within the article, Extended Data, and Supplementary Information. Source data are provided with this paper. Other data are available from the corresponding author upon reasonable request.
Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

- Life sciences  
- Behavioural & social sciences  
- Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

| Sample size       | No sample size calculation performed. For cryo-EM experiment, the sample size is determined by sufficient signal in the experiment to ensure confidence in conclusions drawn from data. That is, a redundant set of micrographs and huge number of particles is used during processing to ensure the best signal-to-noise. |
|-------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Data exclusions   | No data excluded.                                                                                                                                                                                   |
| Replication       | Activity measurements were carried out in technical/biological replicates of n=2 or 3 as stated in figure legends                                                                                 |
| Randomization     | Not applicable - no experimental groups were involved.                                                                                                                                             |
| Blinding          | Not applicable - no group allocation was involved.                                                                                                                                                   |

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

| Materials & experimental systems | Methods                                                                 |
|----------------------------------|------------------------------------------------------------------------|
| n/a                             | n/a                                                                    |
| □ √ Antibodies                   | □ √ ChIP-seq                                                           |
| □ √ Eukaryotic cell lines        | □ Flow cytometry                                                       |
| √ Palaeontology and archaeology  | □ MRI-based neuroimaging                                               |
| □ Animals and other organisms    |                                                                        |
| □ Human research participants    |                                                                        |
| □ Clinical data                  |                                                                        |
| □ Dual use research of concern   |                                                                        |

Antibodies

Primary antibodies:
1. monoclonal ANTI-FLAG M2 antibody, Sigma-Aldrich, F1804
2. monoclonal Anti-TCP-1 α Antibody (R-3), Santa Cruz Biotechnology, sc-374088
3. monoclonal Anti-TCP-1 β Antibody (D-8), Santa Cruz Biotechnology, sc-374152
4. monoclonal Anti-TCP-1 γ Antibody (F-3), Santa Cruz Biotechnology, sc-271336
5. monoclonal Anti-TCP-1 ε Antibody (D-6), Santa Cruz Biotechnology, sc-374554
6. monoclonal Anti-TCP-1 ζ Antibody (F-12), Santa Cruz Biotechnology, sc-271734

Secondary antibody:
IRDye® 800CW Goat anti-Mouse IgG Secondary Antibody, LI-COR, 926-32210

Validation
Validation information is found on the manufacturers’ websites.

Eukaryotic cell lines

Policy information about cell lines

| Cell line source(s)    | HEK293T                                                              |
|------------------------|----------------------------------------------------------------------|
| Authentication         | No cells lines used were authenticated                               |
| Mycoplasma contamination| Cell lines were not tested for mycoplasma contamination                |
Commonly misidentified lines
(See ILAC register)

No commonly misidentified lines were used