



COMMUNICATION

Molecular Structure and Flexibility of the Yeast Coatmer as Revealed by Electron Microscopy

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Received 7 January 2011;
received in revised form
14 March 2011;
accepted 15 March 2011
Available online
22 March 2011

Edited by W. Baumeister

Keywords:

coatmer;
single-particle electron
microscopy;
COPI;
membrane trafficking;
yeast

Coat protein complex I (COPI)-coated vesicles, one of three major types of vesicular carriers in the cell, mediate the early secretory pathway and retrograde transport from the Golgi to the endoplasmic reticulum. COPI vesicles are generated through activation of the regulatory GTPase Arf1 at the donor membrane and the subsequent recruitment of coatmer, a coat protein complex consisting of seven stably associated components. Coatmer functions in binding and sequestering cargo molecules and assembles into a polymeric protein shell that encompasses the surface of COPI vesicles. Little is known about the structural properties of this heptameric complex. We have isolated native yeast coatmer and examined its structure and subunit organization by single-particle electron microscopy. Our analyses provide the first three-dimensional picture of the complete coatmer and reveal substantial conformational flexibility likely to be critical for its scaffolding function.

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Transport vesicles shuttle proteins and lipids between different subcellular compartments. These cargo carriers are generated at the donor compartment by a highly regulated, multistep process that involves several essential factors including coat proteins. Coat proteins are a set of cytoplasmic proteins or protein complexes that are recruited to the membrane of the donor compartment by a regulatory GTPase of the Arf family upon its activation by guanosine 5'-

diphosphate-to-guanosine 5'-triphosphate (GTP) exchange.¹ Coat proteins play critical roles in sorting and concentrating cargo molecules and assemble into a polymeric "coat" structure that deforms the donor membrane, a process that ultimately leads to vesicle budding.² It has been suggested that coat proteins, which cover the surface of transport vesicles, might also contribute to vesicle targeting specificity by serving as a molecular identification code for recognition by the vesicle tethering machinery located at the acceptor membrane.³

Clathrin, coat protein complex I (COPI), and coat protein complex II (COPII) are the three major types of coats found on transport vesicles.^{2,4} Clathrin-coated vesicles mediate endocytic uptake and post-endoplasmic reticulum (ER) sorting at the Golgi, while COPI-coated vesicles participate in both intra-Golgi trafficking and retrograde transport from the Golgi to the ER. COPII-coated vesicles are responsible for exporting cargo from the ER to the Golgi. These three coat proteins are divergent in composition but are highly analogous in overall design. In

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Abbreviations used: COPI, coat protein complex I; COPII, coat protein complex II; EM, electron microscopy; GTP, guanosine 5'-triphosphate; ER, endoplasmic reticulum; TAP, tandem affinity purification; 3D, three-dimensional; GFP, green fluorescent protein.

particular, they all consist of an “adaptor” module that interacts directly with the cargo and the donor membrane and a “cage” module that mediates intersubunit interaction and formation of a polymeric coat structure.^{2,4}

For clathrin and COPII, the adaptor and cage modules exist as separate protein complexes, which integrate into a two-layer coat structure during biogenesis of the transport vesicle.² The molecular architecture and organization of the clathrin and COPII coats have been extensively studied. In particular, pseudo-atomic models of the clathrin and COPII coats have been generated based on cryo-

electron microscopy (EM) reconstructions of *in vitro* assembled coats and crystal structures of individual components or subcomplexes.^{5–7} These structures revealed how clathrin and COPII coat proteins, specifically their cage modules, use extended α solenoid and β propeller domains to mediate intersubunit interactions.

By contrast, the adaptor and cage modules of the COPI coat form a single complex known as coatomer. Coatomer consists of seven components (α , β , β' , γ , δ , ϵ , and ζ) and is recruited *en bloc* to the donor membrane by activated Arf1.^{8,9} Biochemical analyses suggested that the α , β' , and ϵ components of

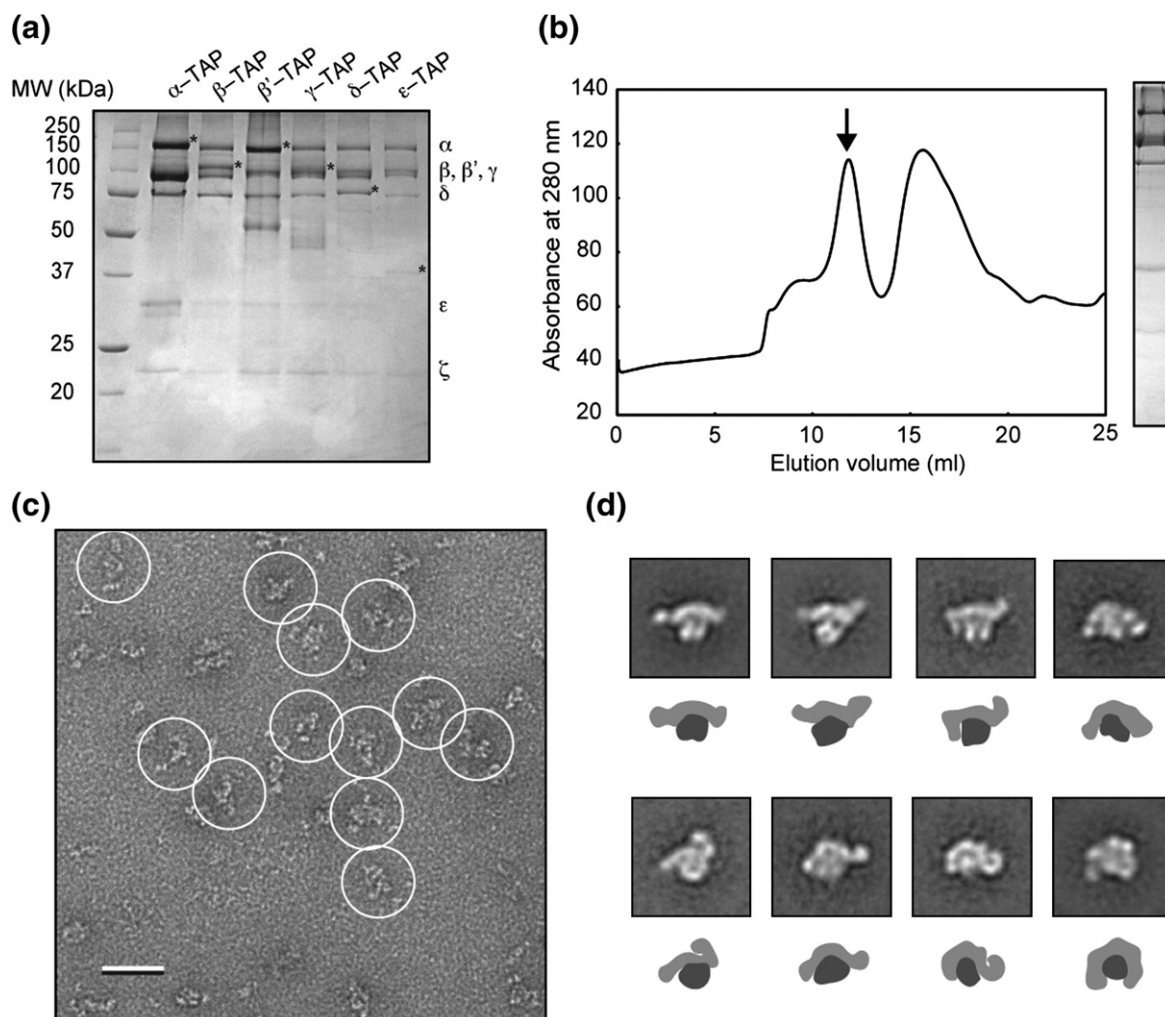


Fig. 1. Purification and negative-stain EM analysis of yeast coatomer. (a) Coomassie-stained 12.5% SDS-PAGE gel of coatomer purified by the conventional two-step TAP procedure.¹⁷ The TAP-tagged strains are shown on the top, with the identities of the individual subunits denoted on the right. Asterisks in each lane indicate the TAP-tagged subunit, which contains only the calmodulin-binding peptide (CBP) part of the tag, as the protein A module is proteolytically removed during TAP purification. (b) Gel-filtration chromatography. The arrow indicates the coatomer peak, with the corresponding silver-stained gel shown on the right. The second peak corresponds to contaminants and dissociated or degraded complexes. (c) A raw image of negatively stained coatomer with individual particles circled. The scale bar represents 50 nm. (d) Eight representative class averages of cryo-negatively stained coatomer with schematic representations underneath each panel showing the two major domains of the coatomer (dark gray for globular domain and light gray for extended domain). The side length of each panel is 38 nm.

coatomer form the cage module or the B subcomplex, while β' , γ , δ , and ζ , four components that share limited sequence similarities with subunits of the canonical clathrin adaptors AP-1 and AP-2, constitute the corresponding adaptor module or the F subcomplex.¹⁰ Although high-resolution structures of two small subunits (ϵ and ζ) and fragments of three larger subunits (α , β' , and γ) have recently been determined,^{11–14} crystallographic analysis of the complete heptameric coatomer has not yet been successful, and little is known about the overall structure of coatomer and its subunit organization. Here, we report the structural characterization of native yeast coatomer by single-particle EM. Our analyses provide insights into the overall architecture and subunit organization of the full coatomer complex and reveal substantial conformational flexibility likely to be critical for its scaffolding function.

Purification of native coatomer from yeast

To obtain sufficient quantities of coatomer for structural studies, we first developed a reproducible purification method for this complex. The fact that coatomer contains seven subunits, four of which are larger than 90 kDa, has precluded the use of recombinant expression systems for this purpose. The traditional method of isolating native coatomer involves repeated ion-exchange chromatography and high-salt elution steps,^{15,16} and these more stringent procedures could compromise the structural integrity of coatomer. Tandem affinity purification (TAP) is a proven method for isolating native complexes from yeast extracts.¹⁷ The TAP tag consists of a calmodulin-binding peptide separated from a protein A tag by a tobacco etch virus (TEV) protease cleavage site.¹⁷ The two-step TAP procedure, involving IgG Sepharose and calmodulin Sepharose, is simple and uses mild buffers and elution conditions. To evaluate the feasibility of using the TAP method to isolate coatomer, we carried out conventional two-step TAP purification from the extracts of six different yeast strains. Each of these strains encodes a TAP tag fused to the C terminus of the essential α , β , β' , γ , and δ subunits or the nonessential ϵ subunit. The addition of the tag does not appear to interfere with the function of coatomer, as all six strains exhibited wild-type growth rates in yeast/peptone/dextrose (YPD) medium. We were able to recover a seven-component complex from the six independent purifications (Fig. 1a), although the subunit stoichiometry varied to some degree. Mass spectrometry confirmed that the bands on the SDS-PAGE gels corresponded to the seven coatomer subunits (Table S1).

We next tested whether we could modify the standard TAP procedure to improve the purity of the isolated coatomer. In particular, we substituted the second affinity purification step (calmodulin

Sepharose) with gel-filtration chromatography, which is more effective than calmodulin Sepharose in separating and removing aggregated protein, subcomplexes, and the tobacco etch virus protease used to elute the complexes from the IgG Sepharose resin. We carried out the two-step IgG Sepharose gel-filtration procedure for all six TAP-tagged strains and observed one major peak in the gel-filtration profile of all samples (α -TAP is shown in Fig. 1b). SDS-PAGE analysis confirmed that the peak fractions contained all seven coatomer subunits and showed that the new procedure resulted in improved purity of the complex (α -TAP is shown in Fig. 1b). Coatomer eluted from the gel-filtration column at a volume similar to that of the mass standard thyroglobulin (669 kDa), indicating that the purified complex (calculated mass of 570 kDa) is likely a “monomer,” with a large hydrodynamic radius (Fig. 1b). We also found that addition of glycerol and the detergent NP40 was necessary to prevent protein aggregation. Furthermore, we noticed that the detergent 3-[(3-cholamidopropyl)dimethylammonio]propanesulfonic acid (CHAPS), although successfully used in the purification of the Nup84¹⁸ and TRAPP¹⁹ complexes, compromised the structural integrity of coatomer.

Negative-stain two-dimensional analysis and three-dimensional reconstruction of yeast coatomer

With the ability to isolate relatively pure yeast coatomer, we next characterized its structure by single-particle EM methods. We first used uranyl formate to prepare negatively stained specimens from the peak fraction of the gel-filtration column (α -TAP) and then examined these specimens on a CM10 electron microscope operated at 100 kV. The raw EM images showed monodispersed particles with different overall shapes (Fig. 1c). To better resolve the structural details of coatomer, we collected more images on imaging plates at a nominal magnification of 67,000 \times and a defocus value of $-1.5\ \mu\text{m}$ using a Tecnai T12 electron microscope operated at 120 kV. We selected 12,515 particles from the digitized images using BOXER, the display program associated with the EMAN software package,²⁰ and classified these particle images into 100 classes using the SPIDER software package.²¹ From the averaged images of these classes, we deduced that coatomer consists of two major domains: a more globular domain connected to the midsection of a more extended domain (Fig. S1a). The observation of these two domains is consistent with the notion that coatomer consists of two subcomplexes, which represent the cage and the adaptor modules, respectively.¹⁰ These two domains appear to adopt multiple conformations, resulting in striking differences in the overall

morphology seen in the calculated class averages (Fig. S1a).

The substantial structural variability of the coatomer population revealed by the two-dimensional averages (Fig. S1a) ruled out cryo-EM analysis of the coatomer structure, as it is currently virtually

impossible to use this technique for very heterogeneous specimens. Negative staining not only provides higher image contrast but also often induces the molecules to adsorb in one or a few preferred orientations to the carbon support film, making it possible to classify the imaged particles into

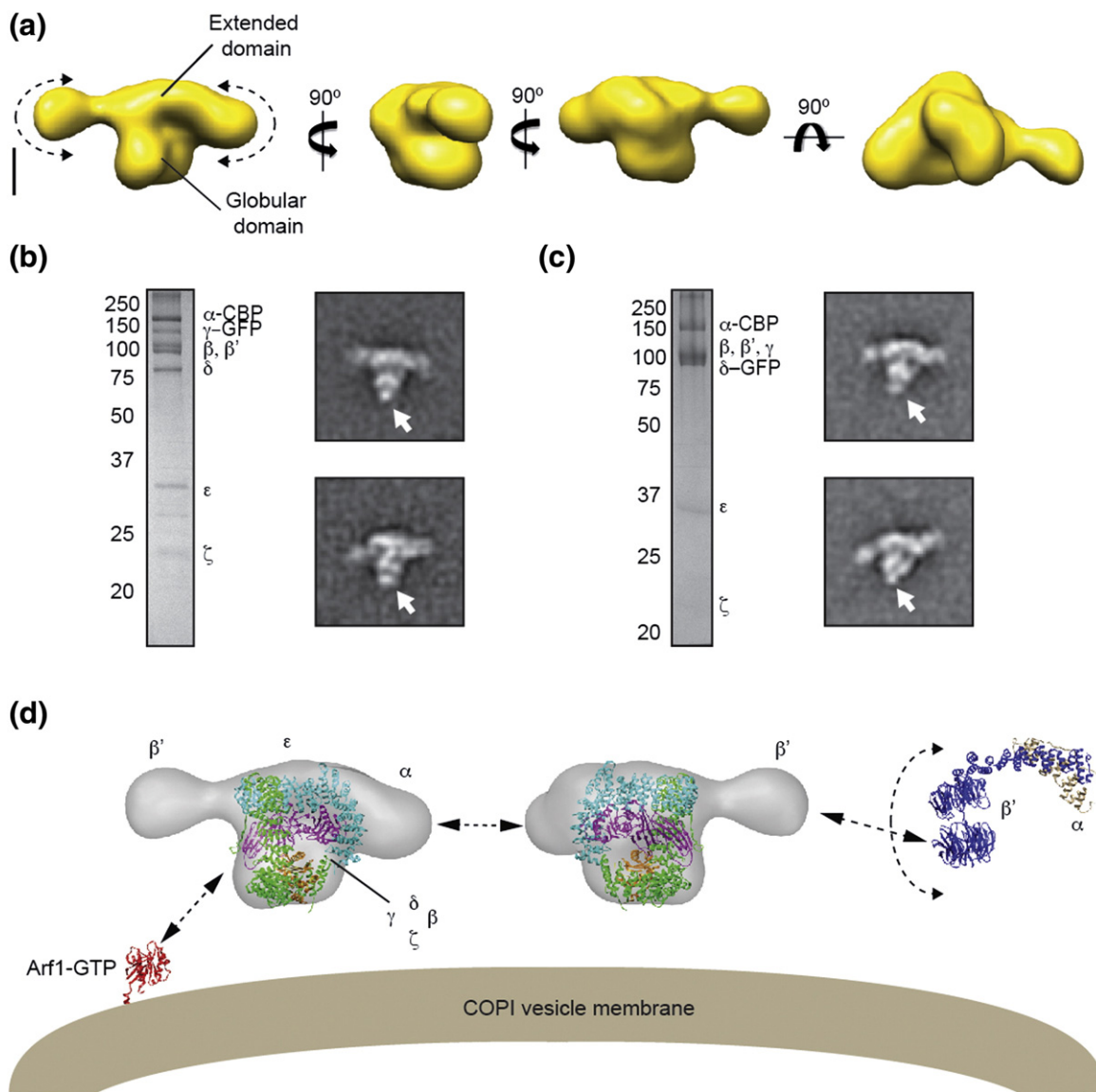


Fig. 2. 3D reconstruction and subunit organization of yeast coatomer. (a) Different views of the 3D reconstruction of coatomer corresponding to the class average shown in the top left panel of Fig. 1a. The arrows indicate the flexibility of the terminal parts of the extended domain. The scale bar represents 5 nm. (b) A silver-stained gel of purified coatomer with GFP fused to the γ subunit (γ -GFP) (left) with two representative averages (right). The arrow indicates the GFP density. The side length of each panel is 43 nm. (c) A silver-stained gel of purified coatomer with GFP fused to the δ subunit (δ -GFP) (left) with two representative averages (right). The arrow indicates the GFP density. The side length of each panel is 43 nm. (d) Placement of the crystal structure of the clathrin AP-2 core complex [Protein Data Bank (PDB) code: 2JKR]²⁴ into the coatomer EM density map. The labels indicate the proposed locations of the seven subunits. A putative model for coatomer recruitment and intersubunit interaction is shown. Upon its activation by GTP exchange, Arf1 (PDB code: 2KSQ)²⁵ recruits coatomer by binding to its β and γ subunits. Coatomer units interact with each other through the terminal regions of the B subcomplex, which contains the β propeller domains of the α and β' subunits. In particular, the tandem β propeller domains of the β' subunits from neighboring complexes interact with each other to form the vertex element of the COPI coat (PDB code of core B subcomplex: 3MKQ).¹²

structurally homogeneous groups according to their conformation. However, negative staining also introduces preparation artifacts, most notably flattening of the particles upon sample dehydration. Cryo-negative staining has been shown to minimize preparation artifacts and, when combined with the random conical tilt approach, provides a powerful method to determine accurate three-dimensional (3D) reconstructions *de novo*.²² We therefore prepared cryo-negatively stained specimens by embedding purified coatomer (α -TAP) in a layer of glycerol and uranyl formate sandwiched between two carbon films on Quantifoil grids. The specimens were frozen in liquid nitrogen and maintained at liquid nitrogen temperature during data collection. We collected pairs of images (50° tilted/untitled) from these specimens at a magnification of 50,000 \times and a defocus value of $-2\ \mu\text{m}$ using an Oxford CT-3500 cryo-transfer holder and a Tecnai F20 electron microscope operated at 200 kV (Fig. S1b). We selected 20,561 pairs of particles from 56 corresponding pairs of images using WEB, the display program associated with SPIDER,²¹ and classified the particles from the images of the untitled specimen into 100 classes using SPIDER. The resulting class averages showed a lack of stain accumulation around the particles and clear structural features, indicating that the purified complexes were well preserved during specimen preparation and imaging (Fig. 1d and Fig. S1c). The majority of the 100 class averages resemble those seen in the class averages obtained with the negatively stained specimen (Fig. S1b and c). Using the corresponding particles from the images of the tilted specimen, we calculated independent 3D reconstructions using the random conical tilt approach,²³ a procedure implemented in SPIDER.²¹ We only selected classes with at least 200 particles to ensure that the 3D reconstructions we calculated would represent a significant percentage of the coatomer population. We also omitted classes that showed diffuse densities in their averaged images, as these indicate poor alignment and/or substantial remaining heterogeneity in the class.

We determined 3D reconstructions of 20 classes. The density maps show a similar two-domain architecture but display considerable differences in the structural features and orientation of the two domains (Fig. S1d and Fig. S2a–e). In particular, the extended domain, as shown by the two-dimensional class averages (Fig. 1d) and substantiated by the 3D reconstructions, is highly flexible and leads to an overall shape of coatomer that ranges from a “V” to a “horseshoe” (Fig. 1d). The terminal regions of the extended domain project to different directions, resulting in even more structural diversity. The inherent flexibility of the extended domain may be important for coatomer to make the necessary adjustments needed to form coats on vesicles of

varying sizes. This structural property may also allow coatomer to overcome the geometrical constraints imposed by having the cage and the adaptor modules in a single entity. We also found that the size of the globular domain varies between different reconstructions when we displayed the maps at the contour level consistent with the calculated mass of the coatomer. We attribute this problem to the more flexible parts being averaged out in the reconstruction process. Inspection of each density map at different contour levels showed that the 3D reconstruction from class 36 was least affected by this issue (Fig. 2a), indicating that it was the class with the least heterogeneity. We therefore chose this particular reconstruction as the reference structure for our interpretation and further discussion. Based on its strong resemblance to other class averages, this density map also seems to capture most of the key structural elements of coatomer. With an estimated resolution of 36 Å according to the Fourier shell correlation=0.5 criterion (Fig. S1d), the class 36 reconstruction has an overall T shape with two well-defined domains (Fig. 2a). The extended domain is triangular and slightly curved and spans a distance of approximately 250 Å, with an ellipsoidal density at one end (Fig. 2a). In projection, the extended domain resembles the Sec13–Sec31 complex, the cage module of the COPII coat.²⁶ The globular domain, with a dimension of about $100 \times 110 \times 60\ \text{Å}$, connects to the extended domain at an angle and contains a central groove reminiscent of that in AP1 and AP2, two clathrin adaptor complexes that are believed to share structural homology with the adaptor module of the coatomer.^{24,27}

Subunit organization of the coatomer

To validate our assignment of the two modules, we carried out EM-based green fluorescent protein (GFP) localization studies to investigate the subunit organization of coatomer. In this approach, we take advantage of the globular fold instead of the fluorescence property of GFP, which is sufficiently large to be observed by EM.²⁸ We generated three yeast strains, in which GFP was fused to the C terminus of the components of the B subcomplex, α , β' , and ϵ , and two additional strains, in which GFP was fused to the C terminus of the components of the F subcomplex, γ and δ . We purified the GFP-tagged complexes from these strains based on TAP-tagged α or, in the case of α -GFP, TAP-tagged δ . Although the overall yields varied across the different strains, all five GFP-tagged complexes could be isolated at a purity comparable to that of the non-GFP-tagged complex (Fig. 2b and c and Fig. S2f–h). Furthermore, the GFP tag did not seem to interfere with complex assembly, as all five complexes eluted at volumes from the gel-filtration

column as the one without GFP (data not shown). We prepared negatively stained specimens from the purified GFP-labeled complexes and collected images using a Tecnai T12 electron microscope. For each of the five samples, we selected at least 3000 particles from the digitized images, which we classified into 200 classes using SPIDER. For γ -GFP and δ -GFP, we could unambiguously identify an additional density in the class averages positioned directly underneath the globular domain (Fig. 2b and c). By contrast, we could not detect any additional density in the class averages obtained for α -GFP, β' -GFP, or ϵ -GFP (Fig. S2f–h). A possible explanation is that the GFP tags are positioned within the core density and are thus obscured. Alternatively, the positions of the GFPs could be too variable, causing them to be averaged out. Nevertheless, the GFP labeling data provide evidence supporting our interpretation that the globular domain represents the F subcomplex or adaptor module and imply that the extended domain corresponds to the B subcomplex or cage module.

Using the position of the central groove as a guide, we were able to place the crystal structure of the clathrin AP-2 core into the globular domain of the class 36 density map. According to this placement, coatomer is positioned with its globular domain facing the donor membrane (Fig. 2d), which would position the adaptor module subunits β and γ in close proximity to Arf1. In agreement with this proposal, previous studies have shown that Arf1, upon activation by GTP exchange, localizes to the donor membrane and recruits coatomer by binding to these two proteins^{29,30} (Fig. 2d). We were, however, unable to fit the crystal structure of the core cage module (β' in complex with a fragment of α) into the class 36 density map.¹² One possible explanation is that this particular 3D map captured a conformation of coatomer different from the one found in the crystal structure, but the uncertainty concerning the appropriate threshold for contouring prevented us from using our other density maps for docking experiments. Our 3D reconstruction also does not have sufficient resolution and structural detail to allow unambiguous docking of the crystal and NMR structures of the ϵ and ζ components into the density map. We did notice, however, that the N-terminal β propeller domains of the β' subunit, which engage in a three-way junction hypothesized to mediate intersubunit interactions at the vertex of the COPI cage,¹² are flexible and could potentially represent the ellipsoid subdomain in our EM map. Future experiments will focus on developing methods to reconstitute COPI coats *in vitro* to not only validate this proposal but also further delineate the molecular details of the interaction between different subunits within coatomer and between different coatomer complexes within an assembled coat.

Supplementary materials related to this article can be found online at [doi:10.1016/j.jmb.2011.03.029](https://doi.org/10.1016/j.jmb.2011.03.029)

Acknowledgements

We thank Zongli Li for assistance with microscopy and image processing. C.K.Y. acknowledges fellowships from the Jane Coffin Childs Memorial Fund and the Canadian Institutes for Health Research. T.W. is an investigator in the Howard Hughes Medical Institute. The molecular EM facility at Harvard Medical School was established with a generous donation from the Giovanni Armenise Harvard Center for Structural Biology and is maintained with funds from National Institutes of Health grant PO1 GM62580 (to Stephen C. Harrison).

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