The Appendage Domain of the AP-2 Subunit Is Not Required for Assembly or Invagination of Clathrin-coated Pits

John S. Peeler, William C. Donzell, and Richard G. W. Anderson

Department of Cell Biology and Neuroscience, University of Texas Southwestern Medical Center at Dallas, Dallas, Texas 75235

Abstract. Coated pits contain a resident membrane molecule(s) that binds clathrin AP-2 with high affinity. AP-2 binding to this site is likely to be the first step in coated pit assembly because this subunit functions as a template for the polymerization of clathrin into flat polygonal lattices. Integral membrane proteins involved in receptor mediated endocytosis cluster in the newly assembled pits as they invaginate and bud from the membrane. The AP-2 subunit is a multi-domain, molecular complex that can be separated by proteolysis into a brick-shaped core and ear-like appendage domains. We have used this property to identify the domain involved in the various stages of coated pit assembly and budding. We found that the core of AP-2 is the domain that binds both to membranes and to triskelions during assembly. Triskelions are perfectly capable of forming lattices on the membrane bound cores. Clathrin lattices bound only to core domains were also able to invaginate normally. Limited proteolysis was also useful for further characterizing the AP-2 binding site. Elastase treatment of the inside membrane surface released a peptide fraction that is able to bind AP-2 in solution and prevent it from interacting with membranes. Affinity purification of binding activity yielded a collection of peptides that was dominated by a 45-kD species. This is the candidate peptide for containing the AP-2-binding site. Therefore, the appendage domain does not directly participate in any of the assembly or invagination events required for coated pit function.

Clathrin-coated pits at the surface of cells are composed of two subunits (18, 19): the triskelion and the clathrin AP complex (Assembly Protein; (12), Adapter Protein; (23, 25). The overlapping legs of the triskelions form the walls of each polygon in the clathrin lattice (12, 25). The location of the AP subunits between the lattice and the membrane (35) suggests that they attach the clathrin lattice to the membrane. The AP-2 subunit is a multi-protein complex that consists of two 100-kD proteins (termed α and β adaptin) (25), one 16-kD protein and one 50-kD protein (2, 24). Each type of subunit can be selectively removed from clathrin coated membranes: the triskelions by high pH and both the triskelions and the APs by high salt (12). These two treatments allow one to study how the two sets of subunits interact with each other and with resident components of the membrane (18, 19, 36).

The clathrin lattice most likely is the molecular device that shapes the coated pit into a coated vesicle because selective removal of triskelions prevents coated pit invagination (17). The conversion of hexagons into pentagons within the lattice occurs during the change in curvature of coated pits (9), but there is no evidence that polygon rearrangement is the force-generating mechanism for invagination.

One function of the AP-2 subunit may be to control the clustering of membrane receptors that enter cells by coated pits (5, 23, 25). It may also anchor the lattice to the membrane (18). The subunit has the capability of performing more than one task because it structurally contains multiple domains. EM has shown that the subunit is composed of a brick-shaped core with two ear-like appendages extending from one end (11). Elastase will remove the appendages without degrading the proteins in the core (38). The appendages released by elastase correspond to the 30-kD COOH terminus of both the α and β adaptin (11, 38).

Several years ago we introduced a method for preparing plasma membranes that is ideal for studying the assembly and function of coated pits in vitro (20). We have shown that the plasma membrane-associated clathrin AP, AP-2 (29, 31), binds to a high affinity site in coated pits (19). The Golgi-associated clathrin AP, AP-1, seems to bind much less tightly to the plasma membrane (19), which suggests that AP-2 binding is specific. The binding site shares some of the properties of an integral membrane protein; for example, it is resistant to high salt but sensitive to proteases (19).

This preparation of coated pits offers a unique opportunity to build on the knowledge gained from studying coated vesicle assembly in vitro and to identify the components required for coated pit assembly and function (12). We have used limited proteolysis and column chromatography to prepare purified cores and appendages from AP-2. We found that AP-2 binding to membranes, clathrin lattice formation, and invagination are all under the control of the core domain within the subunit. We also found that elastase releases AP-2 binding activity from the membrane and that this activity is

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in a peptide fraction with a dominant peptide species of 45 kD. This result suggests that plasma membranes contain an integral membrane protein that functions as an AP-2 receptor during coated pit assembly.

Materials and Methods

Materials

Immuron 1Remowwell 96-well plates (01-010-6301) were purchased from Dynatech Laboratories, Inc. (Alexandria, VA). Human plasma fibronectin was obtained from the New York Blood Center (New York). Elastase (LSO6363) was from Worthington Biomedical Corp. (Freehold, NJ). Diisopropyl phospate (DFP) was from Sigma Chemical Company (DO879). Biotinylated horse anti-mouse IgG (BA-2000) was from Vector Laboratories, Inc. (Burlingame, CA). [35S]methionine (Trans 35S-label) was from ICN Biomedical, Inc. (Irvine, CA). All other reagents were analytical grade and obtained as previously described (18-20).

Methods

Buffers and Cell Culture Medium

Cell Attachment Medium. 20 mM Hepes-buffered (pH 7.4) MEM with Earles salts and without NaHCO3 (330-1435; Gibco Laboratories, Grand Island, NY).

Cell Culture Medium. DME (320-1885; Gibco Laboratories) buffered with 20 mM Hepes (pH 7.4), and supplemented with 2 mM l-glutamine, 100 mM MgCl2, 100 mM KCl, 20 mM Hepes (pH 7.2), 68 mM KCl, 4 mM Mg acetate, 100 mM KCI, 1 mM DTT, protease inhibitor cocktail.

Protease Inhibitor Cocktail. 10 μM leupeptin, 1 mM 1,10-orthophenanthroline, 0.5 mM benzamidine, 2 μg/ml soybean trypsin inhibitor, 0.5 mM PMSF.

Sonication Buffer. 20 mM Mes (pH 6.2), 2.5 mM EGTA, 2.5 mM MgCl2, 100 mM KC1, 1 mM DTT, protease inhibitor cocktail.

Cytosol Buffer. Sonicated Hepes (pH 7.2), 58 mM KCI, 4 mM Mg acetate, 1 mM DTT, protease inhibitor cocktail.

Tris-Stripping Buffer. Cytosol buffer containing 0.6 M Tris (pH 7.2, 4°C), protease inhibitor cocktail.

Taps-Stripping Buffer. 20 mM Taps (pH 9.0), 1 mM DTT, protease inhibitor cocktail.

Cell Culture

 Cultured fibroblasts were derived from a skin biopsy obtained from a normal human subject. Cells were grown in monolayer and set up for experiments according to a standard format (8). On day 0, 7 × 10^4 cells were seeded into each Petri dish (100 × 15 mm) containing 10 ml DME supplemented with 100 U/ml penicillin, 100 μg/ml streptomycin, and 10% (vol/vol) CS. Fresh medium of the same composition was added on day 3. On day 5 of cell growth, each monolayer received 8 ml DME supplemented with penicillin, streptomycin, 5 μg/ml selenium, 5 μg/ml insulin, 5 μg/ml transferrin, and 10% (vol/vol) human lipoprotein-depleted serum. Cells were used for all binding experiments on day 7 of cell growth. SV-40 transformed human fibroblasts (SV-589 cells) were grown as a monolayer (100-μm-diam dishes) in cell culture medium as previously described (20). Cells were harvested in log phase of growth and used to prepare elastase fragments (see below).

Preparation of Coat Proteins

Clathrin and associated proteins were extracted from isolated bovine-coated vesicles with 0.6 M Tris and either used directly or purified clathrin triskelions and clathrin AP subunits by gel filtration (Superose 6) as previously described (18). The AP-1 and AP-2 subunits were separated by hydroxypatite chromatography as described (24).

To prepare AP-2 cores and appendages, purified AP-2 (2–4 mg/ml) was dialyzed into cytosol buffer without protease inhibitors and 400 μl of this solution was treated with 1 μg/ml elastase at 4°C for 45 min. The reaction was stopped by adding DFP to a final concentration of 10 mM. The solution was loaded onto a 1 × 24 cm Sephacryl S-200 SF column and chromatographed with cytosol buffer containing the protease inhibitors PMSF and leupeptin. Fractions containing the appendages and the cores of the proteolysed AP-2 were identified by SDS-PAGE. Virtually complete removal of both the α and the β appendage domains was achieved by these treatments.

AP-2 Binding Assay

Immuron 1removewells were coated with 1 μg/ml poly-L-lysine for 60 min at 37°C, followed by 1 μg/ml fibronectin for 60 min at 37°C. Normal human fibroblasts were harvested during log-phase growth, washed in serum-free cell attachment medium and 250,000 cells were added to each well. The cells were allowed to spread for 90 min at 37°C without serum and then 10% FBS was added to each well for an additional 15 min. The plates were placed on ice and all subsequent treatments were carried out at 4°C. Wells that did not contain cells were used as a control in all experiments. After 60 min, the wells were washed with sonication buffer, attached to the bottom of the buffer reservoir containing 700 μl of sonication buffer, and then sonicated at constant energy (15 joules) using a 3-mm-microtip step probe (Vibra Cell VC60; Sonics and Materials, Danbury, CT) positioned 2 mm above the top of each well. The wells were washed with sonication buffer and used for experiments as described below. In some experiments AP-2 binding was assessed by incubating membranes with unfractionated coat proteins whereas in other trials the purified AP-2 subunit was used (see figure legends). Wells were subsequently fixed with 3% paraformaldehyde and assayed for the presence of either clathrin or specific domains of the AP-2 subunit using an 125I-streptavidin radioimmunoassay as previously described (19).

Preparation of Elastase Fragments

SV-40 transformed human fibroblasts (SV-589) were harvested with Trypsin-EDTA, washed with cell attachment medium and 400,000 cells were allowed to attach to the bottom of each polylysine/fibronectin-coated well of a 24-well culture dish. 10 to 20 24-well culture dishes were prepared for each experiment. The cells were incubated in a normal atmosphere at 37°C for 90 min, 10% FBS was added to each well and the cells were incubated for an additional 15 min at 37°C before being placed on ice for 60 min. All subsequent treatments were carried out at 4°C. Each plate was washed twice in sonication buffer and then seven times in Tris-stripping buffer to remove clathrin and AP-2. The plates were washed seven times in cytosol buffer without protease inhibitors and incubated for 45 min on ice with 400 μl/well cytosol buffer containing 2 μg/ml elastase. Proteolysis was terminated by the addition of DFP to a final concentration of 2 mM. The proteolytic fragments were then harvested from each well and 20% glycerol was added. 10-ml aliquots were frozen in liquid nitrogen and stored at −80°C until use. In some experiments, the cells were incubated for 6 h in methionine-free DME culture medium containing 125 μCi/ml [35S]methionine and 10% dialyzed FBS before plating and sonication. The radioabeled fragments prepared from these membranes had a specific activity of 400,000 to 600,000 cpm/mg protein. Elastase fragments prepared from the outside membrane of fibroblasts were treated identically except that the cells were not sonicated.

Affinity Purification of Elastase Fragments

Purified APs were dialyzed at 4°C overnight against Ca2+, Mg2+-free PBS (pH 6.5) containing 1.0 M NaCI (coupling buffer). 4 ml of the AP solution (0.87 mg/ml) was mixed by gentle rocking with 0.5 grams glutaraldehyde-activated silica overnight at 4°C. The AP-coupled silica was placed in a 5-mL
column and washed with coupling buffer until no protein eluted from the column. The remaining free aldehyde groups were quenched with coupling buffer containing 0.3 M ethanolamine. The column was then washed with cytosol buffer containing 1.0 mM DFP and 0.6 M Tris, pH 7.2, and then equilibrated in cytosol buffer without Tris before use. 95% of the protein was coupled to the column. Either unlabeled or 35S-labeled elastase fragments prepared as described above were thawed and fresh DFP (1 mM) was added before pouring over the AP-2-silica column. The flow-through was recirculated over the column for two hours at 0.5 ml/min and then the column was washed extensively with cytosol buffer. The bound protein was then eluted with 5.0 ml of cytosol buffer containing 0.6 M Tris, pH 7.2. Fractions (0.5 ml) were collected and dialyzed into cytosol buffer containing 1.0 mM DFP before testing in competition binding assays. A portion of each 35S-labeled sample was precipitated with ethanol and prepared for SDS-PAGE. The gels were impregnated with 22% PPO in DMSO and fluorographed using pre-flashed x-ray film.

Other Methods

SDS-PAGE was carried out according to the method of Laemmli (16). Protein determinations were made using the micro bicinchoninic acid (BCA) assay (33). Rapid-freeze, deep-etch microscopy was carried out as previously described (18).

Results

The Core of AP-2 Links Triskelions to the Membrane

The domain within the AP-2 subunit that interacts with the high affinity binding site on the membrane should compete for the binding of intact AP-2 to isolated plasma membranes. We used elastase and column chromatography to purify AP-2 appendages and cores. We then mixed different concentrations of each domain with a constant amount of intact AP-2 subunits and incubated the membranes in the presence of the mixture (Fig. 1). The cores abolished the binding of AP-2 to the membrane in a concentration dependent manner (Fig. 1, ●) but the appendages had no effect on binding (Fig. 1, ○). In other trials, we used a core specific mAb (29) to confirm that the isolated core binds to the membrane with about the same affinity as the whole AP-2 (data not shown).

We used the same approach to determine if the core also contains the triskelion binding site. We prepared membranes and treated them with a concentration of elastase that removes both the α and the β appendages from endogenous AP-2 (Fig. 2 A; elastase, ○) and does not affect the ability of purified triskelions to rebind (B). (A) Isolated membranes were prepared and either not treated (Untreated), washed with high pH buffer (Taps), treated with 1 μg/ml of elastase for 30 min at 4°C (Elastase), or treated with elastase followed by washing with high pH buffer (Elastase + Taps). Membranes were fixed and then processed to measure the presence of either the AP-2 appendages with monoclonal AP-6 (●) or the AP-2 cores with monoclonal B1M6 (●) as described. (B) In the same experiment, membranes that had been treated with elastase and stripped of clathrin were incubated with increasing amounts of purified triskelion (●) for 30 min at 4°C. The membranes were then fixed and assayed for the presence of clathrin heavy chain using mAb X-22. The amount of clathrin on the membrane before (●) and after (◆) the high pH treatment is indicated on the ordinate. Half maximal binding was calculated to be 12 nM. Clathrin did not bind to membranes that had been stripped of elastase-treated AP-2 before incubation with 100 μg/ml of purified triskelions for 30 min at 4°C (○, ordinate). All values are the average of duplicate measurements.

Figure 1. The core of AP-2 binds to the membrane binding site. AP-2 was treated with elastase and the appendages were separated from the cores by gel filtration. The indicated amount of either the appendages (○) or the cores (●) was added to each well containing sonicated membranes. While keeping the volume constant, a constant amount of AP-2 was added to each well and the mixture was incubated at 4°C for 30 min. The amount of bound AP-2 was measured using monoclonal AP-6 as described in Materials and Methods. The amount of AP-2 on the membrane before (●) and after (◆) the membranes were treated with Tris is indicated on the ordinate. All values are the average of duplicate measurements.

Figure 2. Elastase removes the appendages from endogenous AP-2 on isolated membranes but does not release either the cores or clathrin (A) and does not affect the ability of purified triskelions to rebind (B). (A) Isolated membranes were prepared and either not treated (Untreated), washed with high pH buffer (Taps), treated with 1 μg/ml of elastase for 30 min at 4°C (Elastase), or treated with elastase followed by washing with high pH buffer (Elastase + Taps). Membranes were fixed and then processed to measure the presence of either the AP-2 appendages with monoclonal AP-6 (●) or the AP-2 cores with monoclonal B1M6 (●) as described. (B) In the same experiment, membranes that had been treated with elastase and stripped of clathrin were incubated with increasing amounts of purified triskelion (●) for 30 min at 4°C. The membranes were then fixed and assayed for the presence of clathrin heavy chain using mAb X-22. The amount of clathrin on the membrane before (●) and after (◆) the high pH treatment is indicated on the ordinate. Half maximal binding was calculated to be 12 nM. Clathrin did not bind to membranes that had been stripped of elastase-treated AP-2 before incubation with 100 μg/ml of purified triskelions for 30 min at 4°C (○, ordinate). All values are the average of duplicate measurements.
Membrane Binding Site Is Released by Elastase

All that we know about the AP-2 binding site is that it is destroyed by elastase and that it is in coated pits (19). Several studies have suggested that clathrin coat proteins can bind to lipid vesicles (6, 28, 34), which raises the possibility that the AP-2 binding site might be a lipid. Therefore, we performed further tests to determine the macromolecular nature of the binding site. We started by determining whether or not elastase released a molecule that retained the ability to bind AP-2. We treated the cytoplasmic surface (inside) and the environmental surface (outside) of human fibroblast membranes with 0.5 M Tris to remove any AP-2 and clathrin be-

Figure 3. Reassembly of clathrin lattices on untreated membranes (A-C) or membranes that had been treated with elastase (D). Membranes were isolated on coverslips as described. One set was either fixed immediately (A), stripped of clathrin with a high pH buffer (B) or stripped and incubated in the presence of 20 μg/ml of triskelions for 1 h at 4°C (C). Another set (D) was first treated with elastase as described in Fig. 2 to remove the AP-2 appendage domain, stripped of clathrin and then incubated in the presence of 20 μg/ml purified triskelions for 1 h at 4°C. All samples were then fixed and processed for rapid-freeze, deep-etch microscopy as described. Arrows (B) indicate the location of coated pits that have been stripped of clathrin. Bar, 0.5 μm.
The effect of elastase fragments prepared from either the inside (○) or the outside (○) membrane surface on AP-2 binding. Elastase fragments were prepared as described. Keeping the volume of the incubation mixture constant (100 µl), varying amounts of either outside (○) or inside (○) fragments were added to purified AP-2 (10 µg/ml) and incubated for 10 min at 4°C. Membranes were then stripped of AP-2 and clathrin and incubated with this mixture for 30 min at 4°C. At the end of the incubation, the samples were fixed and processed for the detection of membrane bound AP-2 using monoclonal AP-6 (1 µg/ml) as described. The amount of AP-2 on the membrane before (●) and after (△) the membranes were treated with Tris is indicated on the ordinate. All values are the average of duplicate measurements.

Before incubating them in the presence of 2 µg/ml of elastase, we added protease inhibitors to each fraction, mixed different amounts of each fraction with a constant amount of coat proteins (containing both AP-2 and clathrin) and incubated the mixture for 10 min at 4°C. Finally, we removed AP-2 and clathrin from isolated membranes and incubated these stripped membranes in the presence of the mixture for 30 min at 4°C. Fig. 5 shows that the addition of increasing amounts of inside fragments to the mixture caused a progressive inhibition of AP-2 binding to the membranes (Fig. 5, ○). By contrast, the same concentrations of outside fragments did not affect binding (Fig. 5, ○).

These results suggest that elastase releases a polypeptide capable of interacting with AP-2 in solution. To see if AP-2 would bind directly to the fragment, we prepared inside and outside elastase fragments and dried different amounts to the bottom of individual wells of a 96-well plate. We used gel electrophoresis to determine that the two preparations contained numerous peptides (data not shown). We then incubated each well in the presence of 10 µg/ml of coat proteins (Fig. 6). AP-2 did not bind to wells containing even the highest concentration of outside fragments (Fig. 6, ○). By contrast, inside fragments did support binding (Fig. 6, ●) and the amount of AP-2 that bound was proportional to the concentration of fragments added to the well. AP-2 bound to wells that received as little as 100 ng of inside elastase fragments.

Figure 4. Effect of elastase on clathrin coated pit invagination. Membranes were prepared as described and either not treated (A), incubated in the presence of 1 µg/ml of elastase for 30 min at 4°C (B), or incubated in the presence of elastase and then warmed to 37°C for 3 min (C). The samples were then fixed and processed for rapid-freeze, deep-etch microscopy as described. Bar, 0.25 µm.
Figure 6. Binding of AP-2 to wells that contain varying amounts of either inside fragments (●) or outside fragments (○). Elastase fragments were prepared as described and the indicated amount was dried onto the surface of individual wells of a 96-well plate. Each well was then incubated with 10 μg/ml of purified AP-2 for 30 min at 4°C, fixed, and processed to detect AP-2 binding using monoclonal AP-6 (1 μg/ml) as described. All values are the average of duplicate measurements.

Elastase fragments also bound to immobilized AP-2. We prepared an affinity column to isolate the elastase fragments by attaching bovine brain APs (a mixture of AP-2 and the Golgi specific AP, AP-1) to glutaraldehyde-activated silica gel. We passed the elastase fragments from the cytoplasmic membrane surface over the column and washed extensively until there was no more protein in the flow through (Fig. 7, B, ●). The addition of 0.6 M Tris-HCl immediately released additional protein from the column (Fig. 7, C, ●). We collected each fraction and tested it for AP-2 binding activity using the competition assay (Fig. 7, ○). The starting material had a modest amount of specific inhibitory activity (Fig. 7, △), but there was not any activity in the initial washes from the column. By contrast, the Tris eluate fractions (Fig. 7 C) contained significant inhibitory activity. The fraction with the highest specific activity had 30-fold more binding activity than the starting material.

We wanted to determine the molecular weight range of the peptides that bound to AP-2. Therefore, we radiolabeled elastase fragments from fibroblasts by growing the cells for 6 h in the presence of [35S]methionine. The radiolabeled fragments were applied to an AP affinity column as described in Fig. 7. The column was washed, 0.6 M Tris-HCl was added, and the eluate was collected in six fractions. The starting material (start), the wash fraction (wash), and each eluate fraction (1-6) was tested for its ability to inhibit AP-2 binding to purified membranes (A) and analyzed by SDS-PAGE (B). The amount of AP-2 on the membrane before (●) and after (▲) the membranes were treated with Tris is indicated on the ordinate (A). All values are the average of duplicate measurements.

Figure 7. Affinity purification of an elastase fragment that contains the AP-2 binding site. Elastase fragments from the inside surface of the plasma membrane were prepared as described. 5 ml of this preparation was applied to an AP affinity column by repeated circulation. The flow-through was collected (A) and then the column was washed (B) until the absorbance at 280 nm (●) declined to zero. 0.6 M Tris was added (C) to release any proteins that had bound to the AP-2 and 0.5-ml fractions were collected. Each fraction was then tested for its ability to inhibit AP-2 binding to the inside surface of the plasma membrane (○). Specific inhibitory activity in the starting material is shown on the right ordinate (△). Specific inhibitory activity equals the amount of binding in the absence of the fraction minus the amount of binding in the presence of the fraction per milligram of protein. All values are the average of duplicate measurements.

Figure 8. Identification of the AP-2 binding fragment. Elastase fragments were prepared from cells that had been grown in the presence of [35S]methionine. The radiolabeled fragments were applied to an AP affinity column as described in Fig. 7. The column was washed, 0.6 M Tris-HCl was added, and the eluate was collected in six fractions. The starting material (start), the wash fraction (wash), and each eluate fraction (1-6) was tested for its ability to inhibit AP-2 binding to purified membranes (A) and analyzed by SDS-PAGE (B). The amount of AP-2 on the membrane before (●) and after (▲) the membranes were treated with Tris is indicated on the ordinate (A). All values are the average of duplicate measurements.
Appendage Domains

parable amounts of activity (Fig. 8 A, columns 2-4). As the intensity of the band declined in fraction 5 and 6, there was also a decline in the ability of the fraction to inhibit AP-2 binding (compare Fig. 8 B, lanes 5 and 6 with Fig. 8 A, columns 5 and 6).

Discussion

Different Functions for the Core and the Appendage Domains

Ordinarily investigators study the interaction between AP-2 and clathrin using clathrin in solution, clathrin immobilized on a support or clathrin assembled into cages (12). Keen and co-workers (13, 27, 37, 38) have used these techniques to show that the core of AP-2 will not promote clathrin cage assembly but is able to bind to intact clathrin trimers. There appear to be two binding sites for the core of AP-2 on the triskelion legs (21, 27): one on the terminal domain and one near the hub region. In contrast to these studies, Schroder and Ungewickell (32) found that in a tartrate buffer system only intact APs would bind to clathrin. They concluded from their studies that tartrate had its affect by preventing the APs from aggregating.

The primary interaction between these two subunits ordinarily occurs within the cell during coated pit formation. We designed the in vitro assay used in the current study to mimic the conditions that favor these interactions (18, 20). When we treated membranes with elastase concentrations that are known to remove both the α and the β appendages from purified coated vesicle APs (32), clathrin coated pit assembly appeared to be unimpaired. We can not rule out the possibility that these conditions did not completely remove both appendages from the membrane bound APs. Nevertheless the affinity of interaction between clathrin and the bound, elastase treated AP-2 was unaltered (compare Fig. 2 B with Fig. 3 in reference 19). Furthermore, the bound triskelions formed into lattices that were normal in appearance.

Rapid-freeze, deep-etch images have previously shown that triskelions make contact with solid surfaces through the terminal portion of each leg (14) and that this is the region that interacts with APs in solution (11). Murphy et al. (21) recently showed that this region of the leg binds to the AP core domain. We propose that coated pit lattice assembly requires the high affinity binding of the triskelion terminal region to the core of AP-2. Possibly β-adaptin contains the binding site for clathrin (1). The core is also the domain that mediates AP-2 binding to the inner membrane surface. Elastase treatment of intact coated pits removed the appendages without releasing the AP-2 core from the membrane, which indicates that the appendage does not have a high affinity for the binding site (Fig. 2 A). What is more important is that only the purified cores inhibited AP-2 binding to membranes (Fig. 1). The AP-2 domain that binds to the membrane could reside in the non-proteolysed, amino terminal portion of either α or β adaptin.

We conclude that all of the major interactions that occur between the membrane and clathrin during assembly of coated pits are mediated by the AP-2 core domain. This leaves the appendage domain available to carry out other key reactions necessary for coated pit function. One activity may be to cluster receptors that enter cells by receptor mediated endocytosis (3) while another may be to control the shape of the lattice.

Molecular evidence has suggested a role for the appendage domain in the adaptive ability of the AP-2 complex to recognize the cytoplasmic tails of multiple receptors that enter coated pits (15, 26, 30). On the other hand, Beltzer and Spies (5) have found that the amino terminal domain of β-adaptin, which is present in the AP-2 core, binds a fusion protein that contains the cytoplasmic tail of the asialoglycoprotein receptor. Therefore, the exact role of the appendages in receptor clustering is not known.

Elastase treatment did not prevent coated pits from rounding up at 37°C. In fact, there seemed to be a tendency for elastase by itself to cause coated pits to change shape. We observed this in the shape change experiments (Fig. 4 B) and in the reassembly trials (Fig. 3 D). This raises the possibility that the appendage domain plays an active role in keeping the lattice in a planar configuration during assembly. This may be important for the proper clustering of receptors before budding and for the ordered addition of clathrin trimers into the forming polygonal lattice.

The AP-2 Receptor

The protein fragments released by elastase from the inside surface of the plasma membrane contain the high affinity, AP-2 binding site. The predominant band in the affinity-purified fraction has an apparent molecular weight of 45,000. This peptide is a candidate for containing the AP-2 binding site. Most likely the peptide or peptides that have the binding activity are derived from integral membrane proteins that are essential for coated pit assembly. Therefore, AP-2 seems not to be binding with high affinity to a specific class of plasma membrane lipids.

The isolation of peptide fragments, released specifically from the inside surface of membranes, that have AP-2 binding activity raises the possibility that plasma membranes contain an AP-2 receptor. The major function of this receptor would be to bind the core of the AP-2 and initiate coated pit assembly. The appendage domain, on the other hand, would be available to interact with membrane proteins that contain a reverse-turn, three-dimensional conformation in their cytoplasmic tail (4) and cause them to cluster before coated pit budding (3). This model agrees with rapid-freeze, deep-etch microscopy studies that show coated pit assembly precedes membrane receptor clustering (10). A hierarchical arrangement of domains within the AP-2 subunit gives each cell type the flexibility of controlling the uptake of a specific ligand without affecting the whole endocytic process (22).

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