Mannose 6-Phosphate Receptors Regulate the Formation of Clathrin-coated Vesicles in the TGN
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Abstract. The transport of the two mannose 6-phosphate receptors (MPRs) from the secretory pathway to the endocytic pathway is mediated by carrier vesicles coated with the AP-1 Golgi-specific assembly protein and clathrin. Using an in vitro assay that reconstitutes the ARF-1–dependent translocation of cytosolic AP-1 onto membranes of the TGN, we have previously reported that the MPRs are key components for the efficient recruitment of AP-1 (Le Borgne, R., G. Griffiths, and B. Hoflack. 1996. J. Biol. Chem. 271:2162–2170). Using a polyclonal antibody against the mouse γ-adap-
thetic pathway (Herman et al., 1992). Conversely, the treatment of mammalian cells with wortmannin, a specific inhibitor of PI 3-kinases, results in a drastic missorting of lysosomal enzymes (Brown et al., 1995; Davidson, 1995), suggesting that the AP-1-dependent sorting of their mannose 6-phosphate receptors is impaired.

The fundamental mechanisms by which membranes receptors are specifically segregated into clathrin-coated vesicles has been a difficult question to address. Two distinct models have been proposed. First, receptors can simply be trapped into preexisting coated structures or, second, the receptors actively initiate the formation of coated vesicles.

Morphometric analyses performed at the EM level first suggested that the massive overexpression of the human transferrin receptor in mouse cells correlated with a higher number of clathrin-coated pits at the plasma membrane (Iacopetta et al., 1988). Such overexpression in chicken embryonic fibroblasts did not change the number of coated pits but rather increased the number of flat clathrin lattices (Miller et al., 1991). More recent immunofluorescence studies in which FcεRI receptors were relocalized to restricted areas of the plasma membrane could not detect any change in the redistribution of clathrin and assembly protein AP-2 (Santini and Keen, 1996). Thus, whether membrane receptor sorting and clathrin-coated vesicle formation are coupled processes or not has remained controversial so far.

Our previous in vitro studies on fibroblasts genetically devoid of the two MPRs have suggested that these transmembrane proteins are key components for the efficient translocation of cytosolic AP-1 onto membranes of the TGN and possibly on membranes of early endosomes (Le Borgne et al., 1993, 1996). In addition, specific protein motifs in the MPR cytoplasmic domains are important for the high affinity interaction of AP-1 with TGN membranes in vitro (Mauxion et al., 1996). In the present study, we have raised an antibody against the mouse γ-adaptin to directly determine the steady state distribution of AP-1 in mouse fibroblasts lacking MPR expression or reexpressing physiological levels of the MPRs. After cellular fractionation, the results indicate that the expression level of the MPRs not only determines the amount of AP-1 bound to membranes but also determines the number of clathrin-coated vesicles formed in the TGN. We conclude that, in vivo, membrane protein sorting and clathrin-coated vesicle formation in the TGN are coupled processes.

Materials and Methods

Materials

All reagents were of analytical grade. Mouse Apo-Transferrin, Ficol, and 2H2O were from Sigma Chemical Co. (St. Louis, MO). Ribonuclease A was from Worthington Enzymes Ltd. (Freehold, NJ). 30% (wt/vol) acrylamide/0.8% (wt/vol) bis-acrylamide solution was from National Diagnostics (Atlanta, GA). Protran nitrocellulose membranes (0.45 μm) were from Schleicher and Schuell (Dassel, Germany). Pronase was from Boehringer Mannheim GmbH (Munich, Germany). Iodogen was from Pierce Chemical Co. (Rockford, IL). Na125I and ECL detection system were from Amersham Life Science (Amersham, UK).

Cell Culture

Mouse MPR-negative fibroblasts and MPR-negative fibroblasts stably expressing either MPR were generated as previously described (Ludwig et al., 1994; Mauxion et al., 1996; Munier-Lehmann et al., 1996) and grown in DME medium complemented with 10% FCS; 2 mM glutamine, 100 U/ml penicillin, and 100 U/ml streptomycin.

Antibodies

γ-Adaptin and the ubiquitously expressed αC-adaptin (Robinson, 1989) were detected using polyclonal sera prepared by immunizing rabbits with the K-259-274 peptide (K-RNDDSSEAMNDILAO) and the K-623-637 peptide (K-STVTLDEETKRERSI), respectively, coupled to keyhole limpet hemocyanin. The crude rabbit sera were further affinity purified by passing them on columns made of the corresponding immunogenic peptides coupled to Affigel-10. The specific igs were eluted at 4 M MgCl2 and extensively dialyzed against PBS before use.

The antisera against the αC-adaptin detected a single ~110-kD band by Western blotting comigrating with a ~110-kD protein detected with the anti-α-adaptin AP-6 mA. In indirect immunofluorescence, our polyclonal serum and the AP-6 mA decorated the same structures (not shown).

The antisera against the mouse γ-adaptin (the trunk region) recognized two proteins on Western blots. The ~100-kD protein was identified as the γ-adaptin. This protein is also recognized by a polyclonal antibody directed against the hinge region of the mouse γ-adaptin (a kind gift from M. Robinson, Cambridge, U.K., Seaman et al., 1996), and is immunoprecipitated from denatured Hela cells lysates by the monoclonal 100/3 anti-γ-adaptin antibody. The higher molecular mass ~120-kD protein cross-reacting with our anti-peptide antibody does not cross-react with a polyclonal antibody directed against the hinge region of the mouse γ-adaptin, and therefore is most likely a contaminant. In addition, the in vitro recruitment of this 120-kD protein on membranes is insensitive to the addition of brefeldin A or GTPγS (data not shown). In immunofluorescence, the polyclonal anti-peptide antibody against the trunk region of the mouse γ-adaptin decorates the perinuclear region and peripheral structures of mouse fibroblasts and Hela cells as previously observed with the monoclonal 100/3 anti-γ-adaptin antibody. Furthermore, this staining becomes completely cytosolic after a 1–2-min treatment with 5 μg/ml brefeldin A as expected for AP-1.

The rabbit serum against clathrin was as described (Mérresse and Hoflack, 1993). The cation-independent (C1–MPR) was detected with a polyclonal antibody directed against a peptide corresponding to the carboxy-terminal domain of the receptor (Mérresse and Hoflack, 1993). The mouse transferrin receptor was detected using the Hb6-4 mA (Zymed Laboratories, S. San Francisco, CA) directed against the conserved cytosolic domain of human transferrin receptor. β-COP was detected using the E5A5 mA. Goat anti–mouse and goat anti–rabbit secondary antibodies coupled to HRP were from Dianova-Immunotech GmbH (Hamburg, Germany).

Immunoprecipitations

Confluent Hela cells were washed twice with ice-cold PBS, scraped with a rubber policeman, and spun for 5 min at 2,000 g. A postnuclear supernatant was then prepared in lysis buffer (LB; 50 mM Tris, pH 7.4, 100 mM NaCl, 2 mM EDTA, 1 mM PMSF and benzamidine, 5 μg/ml aprotonin, and 1 μg/ml leupeptin). The sample was then supplemented with 1% Triton X-100 (final) for 30 min on ice (native sample) or boiled in 1% SDS for 5 min (denatured sample). After diluting the SDS 10 times, the samples were spun for 15 min in an Eppendorf centrifuge. The supernatant was then preclarified with protein A–Sepharose for 1 h at 4°C and spun for 15 min. The supernatant was incubated overnight at 4°C on a rotating wheel with either 1 μg of our polyclonal anti-γ-adaptin (1573) or 1 μg of the 100/3 monoclonal anti-γ-adaptin antibody. After spinning down for 15 min, 50 μl of a 50% slurry of protein A–Sepharose (Pharmacia Fine Chemicals Inc., Piscataway, NJ) was added to the supernatant for an additional hour. An aliquot of the supernatant of the immunoprecipitation (10%) was kept for Western blotting analysis. The beads were then washed three times in LB, three times in LB + 500 mM NaCl, and once in LB. After boiling in sample buffer twice for 5 min each, the samples were analyzed by Western blotting.

Immunofluorescence

Cocultures of MPR-positive and MPR-negative fibroblasts were grown on glass coverslips, washed in PBS, and fixed with 3% paraformaldehyde. After an extraction with 0.1% Triton X-100, the cells were incubated with the monoclonal 1D4B anti–mouse Lamp-1 antibody (Developmental Stud-
ies Hybridoma Bank, Iowa City, IA) and an affinity-purified rabbit antibody raised against the hinge region of the mouse γ-adaptin (a kind gift from S. Robinson; Seaman et al., 1996). The bound antibodies were detected with FITC- or Texas red–labeled secondary antibodies (Dianova-Immunotech GmbH).

Subcellular Fractionation

Mouse fibroblasts were grown in complete DME medium supplemented with 20 mM Hepes. The cells were washed twice with ice-cold PBS and once with vesicle buffer (140 mM sucrose, 75 mM potassium acetate, 10 mM MES, pH 6.6, 1 mM EGTA, 0.5 mM magnesium acetate). The cells were then scraped with a rubber policeman, and DTT (1 mM) and protease inhibitors (1 μg/ml chymostatin, 1 μg/ml pepstatin, 1 mM PMSF, 1 mM benzamidin, 1 μg/ml antipain) were added. After centrifugation for 10 min at 2,000 g, the cells were resuspended in vesicle buffer. An aliquot was directly boiled in Laemmli buffer (total cell extract). Cells were then homogenized by boiling in Laemmli buffer (total cell extract). Cells were then homogenized by SDS-PAGE and Western blotting. Different cell types were analyzed by SDS-PAGE and Western blotting. For each clone, six to eight different cell types were analyzed by SDS-PAGE and Western blotting.

Electrophoresis, Western Blotting, and Software Analysis

SDS-PAGE was performed using the mini-PROTEAN II electrophoresis system (Bio Rad Laboratories, Hercules, CA). After SDS-PAGE, the proteins were electroblotted onto nitrocellulose for 2 h at 80 V. The membrane was then incubated in blocking buffer (5% defatted milk and 0.05% Tween-20 in PBS) for 2 h and with the primary antibodies diluted in blocking buffer for 2 h at room temperature. After washing, the membranes were incubated with secondary antibodies for 1 h in blocking buffer. HRP-labeled secondary antibodies were detected using the ECL system (Amersham Life Science). The same blots were sequentially probed with different antibodies. For this process, between each immunodetection, the membranes were stripped in 62.5 mM Tris-HCl, pH 6.7, 2% SDS, and 100 mM 2-mercaptoethanol for 30 min at 50°C. For quantitations of the signals, different undersaturating exposures of the autoradiograms were scanned using a Microtek scanner III scanner at a resolution of 300 dpi (Microtek Electronics Europe, GmbH, Düsseldorf, Germany). The intensities of the signals were quantitated using the NIH Image 1.59 pc software. Mounting of the figures was performed using Adobe Photoshop™ 3.0.5 (Adobe Systems, Inc., Mountain View, CA).

Endocytosis of Transferrin

Purified mouse Apo-transferrin (mTf) was first loaded with iron by incubation in a saturation buffer (250 mM Tris-HCl, 10 μM NaHCO3, 2 mM sodium nitritolbicarbonate, and 6 mM FeCl3) for 3 h at room temperature. After extensive dialysis against PBS, mTf was iodinated with iodogen as described in Podbilewicz and Mellman (1990). For internalization studies, triplicate samples of each cell type were grown in 35-mm-diam dishes, washed with PBS, and incubated for 1 h at 37°C in DME medium containing 0.1% BSA and 20 mM Hepes to deplete the cells of bound, bovine Tf from the culture medium. The cells were then placed on ice and incubated for 1 h with 100 nM mTf (25 nM 125I-mTf + 75 nM cold mTf) in DME medium containing 0.1% BSA and 20 mM Hepes. Cells were then incubated at 37°C for different periods of time (0.5, 1, 2.5, and 5 min). After extensive washing in DME medium, 0.1% BSA, 20 mM Hepes, and DME, and 20 mM Hepes, cell surface–bound mTf was removed by pronase treatment (3 mg/ml for 1 h on ice). The cells and the medium were collected and spun down in an Eppendorf centrifuge for 2 min. Both the supernatant (released transferrin) and the cell pellet were quantitated by γ-radiation counting. Endocytosed 125I-Tf was estimated as the residual radioactivity in the cell pellet after pronase treatment and expressed as a percentage of the total cell surface–bound mTf at 4°C.

EM-negative Staining

Material contained in the dense fractions of the last linear density gradient (fractions 8 to 10) was pooled, diluted in MES buffer (0.1 M MES, pH 6.5,
Results

Our previous in vitro studies on MPR-deficient mouse fibroblasts have strongly suggested that the MPRs are key components for the efficient translocation of cytosolic AP-1 onto membranes. These studies were based on an in vitro assay reconstituting the binding of cytosolic bovine AP-1 on membranes of permeabilized mouse fibroblasts. The newly bound AP-1 was detected with the 100/3 anti-γ-adaptin mAb, unable to recognize the endogenous mouse AP-1 (Le Borgne et al., 1993, 1996). When labeled with an antibody against the hinge region of γ-adaptin (Seaman et al., 1996), the mouse fibroblasts expressing the two MPRs exhibit the typical, perinuclear γ-adaptin staining (Fig. 1). In contrast, the fibroblasts devoid of the MPRs, easily identified by the large number of Lamp-1-positive structures as a result of the massive mis-sorting of lysosom-
mal enzymes (Ludwig et al., 1994), show a reduced γ-adaptin staining (Fig. 1). To quantitate by Western blotting the steady state distribution of γ-adaptin in these cells, we have used a polyclonal antibody against a peptide corresponding to a stretch of amino acids (R259–Q274) contained in the trunk domain of the mouse γ-adaptin, also conserved in the human γ-adaptin. Fig. 2 shows that, in Western blot analysis, this anti–mouse γ-adaptin antibody reacts with two proteins of mouse fibroblasts. The first protein is the ~100-kD γ-adaptin because it is also recognized by a polyclonal antibody directed against the hinge region of the mouse γ-adaptin, while the second, ~120-kD protein, not recognized by the polyclonal antibody against the hinge region of the mouse γ-adaptin and absent from purified clathrin-coated vesicles (see Fig. 5), most likely represents a contaminant. Furthermore, Fig. 2B shows that this anti-peptide antibody immunoprecipitates the γ-adaptin from a total, denatured Hela cell lysate as does the 100/3 mAb. However, this antibody does not immunoprecipitate the native protein.

Expression of Protein Markers in MPR-deficient Fibroblasts

We first determined by quantitative Western blotting the level of expression of different markers in MPR-negative fibroblasts and in MPR-negative fibroblasts stably reexpressing physiological levels of either the cation-dependent (CD) MPR or the insulin-like growth factor II/Cl-MPR (Ludwig et al., 1994; Le Borgne et al., 1996; Mauxion et al., 1996). Fig. 3 shows the relative amounts of γ-adaptin (AP-1), α-adaptin (AP-2), lysosomal marker Lamp-1, or transferrin receptor, a membrane protein recycling between the plasma membrane and the endosomes, as well as that of β-COP, a subunit of the coatomer chosen as a marker of the early secretory pathway in the total cell lysates of the different clones examined. The expression level of each of these markers was nearly identical in all of the different cell types examined. In addition, the ratios of γ-adaptin/α-adaptin (Fig. 3B), γ-adaptin/transferrin receptor (Fig. 3C), or γ-adaptin/β-COP (not shown) were very similar in all of the different cell types examined. This indicates that the expression level of these different markers is, as expected, independent of the expression of the MPRs.

AP-1 Bound to Membranes of MPR-deficient Fibroblasts

We then prepared microsomal fractions from the different clones and determined by quantitative Western blot analysis the amount of α- and γ-adaptins bound to membranes at steady state. The amount of membrane-bound α-adaptin was very similar in the different clones tested and represented ~36.7% ± 4.7% of the total α-adaptin. We also determined the internalization rate of the transferrin receptor to follow the dynamics of its AP-2-dependent endocytosis. For this process, the different mouse fibroblasts were first incubated at 4°C with iodinated mouse transferrin, washed to remove the unbound ligand, and subsequently incubated at 37°C for various periods of time. Table I shows that the internalization rate of the transferrin receptor was similar in all of the different clones tested. These results indicate that the dynamics of endocytosis is not significantly affected by the reexpression of physiological levels of MPRs. Furthermore, our previous studies showing that these different clones secrete similar amounts of proteins (Mauxion et al., 1996; Munier-Lehmann et al., 1996) and the fact that the level of β-COP expression is unchanged suggest that the dynamics of the COPII-dependent vesicular traffic in the early secretory pathway is also not affected.

Since the rate of endocytosis (membrane-bound AP-2 and number of plasma membrane–derived clathrin-coated vesicles formed) is unchanged in these different clones, the amount of membrane-bound γ-adaptin could be normalized to that of α-adaptin or the transferrin receptor. In MPR-positive fibroblasts, 30.5 ± 3.2% of the total AP-1 was typically found associated with membranes. Fig. 4 shows that membranes of MPR-negative fibroblasts contain three times less γ-adaptin than membranes of the corresponding MPR-positive fibroblasts (~9.6 ± 1.8% of the total AP-1). In addition, the examination of stable clones of MPR-negative fibroblasts reexpressing various amounts of either CD-MPR or CI-MPR indicated that the amount of membrane-bound γ-adaptin increased upon the reexpression of either MPR in MPR-negative fibroblasts. These results show that, in vivo, the amount of γ-adaptin bound to membranes at steady state correlates, to some extent, with physiological levels of expression of the MPRs. It is unlikely that these differences in amounts of bound-AP-1 reflect an indirect effect as a result of the large number of Lamp-1–positive structures in MPR-negative fibroblasts. MPR-negative fibroblasts reexpressing physiological levels of either MPR still contain numerous Lamp-1–positive structures (Munier-Lehmann et al., 1996). However, the amount of bound AP-1 is significantly increased in these cells.

AP-1 Associated with Clathrin-coated Vesicles of MPR-deficient Fibroblasts

The membrane-bound γ-adaptin represents AP-1 bound to the donor compartment as well as AP-1 present in transport vesicles. Thus, clathrin-coated vesicles were prepared from the different MPR-deficient fibroblasts using the fractionation protocol described previously by Woodward and Warren (1991). Fig. 5 shows a typical example of the distribution of several marker proteins throughout the density gradient of the last purification step after fractionation of

| Cell type          | Internalized m-Tf (percentage of prebound) |
|--------------------|------------------------------------------|
| MOCK               | 65.6 ± 3.8                               |
| CD-4               | 62.1 ± 3.5                               |
| CD-1               | 60.0 ± 7.9                               |
| CD-2               | 70.6 ± 8.7                               |
| CI-3               | 68.9 ± 7.2                               |
| CI-4               | 67.5 ± 7.2                               |

Mock-transfected MPR-negative fibroblasts or MPR-negative fibroblasts reexpressing the CD-MPR (1.4-, 3.5-, and 4.4-fold the endogenous level of CD-MPR for CD-4, -1, and -2, respectively) or the CI-MPR (one- and fivefold the endogenous level for CI-3 and -4, respectively) were incubated at 4°C with iodinated mouse transferrin, washed, and subsequently incubated at 37°C for various periods of time (see Materials and Methods). The indicated values represent the percentage of prebound transferrin internalized during the first minute of uptake. Values correspond to means ± standard error of two independent experiments performed in triplicate.
MPR-deficient fibroblasts. Typically, the dense fractions contained the clathrin light chain, the \(\gamma\)- and \(\alpha\)-adaptins, as well as transmembrane proteins like the transferrin receptor and the CI-MPR, as determined by Western blotting. Very low amounts of \(\beta\)-COP were occasionally detected in the lighter fractions of this gradient (not shown). When analyzed by SDS-PAGE followed by protein staining, the dense fractions exhibited the typical protein profile of purified clathrin-coated vesicles (Fig. 6A). Beside a few contaminants, the clathrin heavy and light chains as well as the different subunits of the APs were easily detected. At the morphological level, these fractions contained only spherical structures of \(\approx 50–100\) nm in diameter with a coat lattice reminiscent of clathrin-coated vesicles (Fig. 6B).

The amount of \(\gamma\)-adaptin present in clathrin-coated vesicles isolated from the different MPR-deficient fibroblasts was analyzed by quantitative Western blotting. Since the dynamics of endocytosis is unchanged in the different cell types, indicating that the number of plasma membrane-derived clathrin-coated vesicles remains unchanged, these values for \(\gamma\)-adaptin could be normalized to the amount of \(\alpha\)-adaptin or transferrin receptor present in these fractions. Fig. 7 shows that the clathrin-coated vesicle fractions obtained from MPR-negative fibroblasts contain three times less \(\gamma\)-adaptin than those isolated from MPR-positive fibroblasts. Moreover, the purification of clathrin-coated vesicles from MPR-negative fibroblasts reexpressing the MPRs indicated that the amount of \(\gamma\)-adaptin recovered in clathrin-coated vesicle fractions increased upon the reexpression of physiological levels of either MPR. Since the amount of \(\gamma\)-adaptin found in these fractions reflects a number of vesicles, we conclude from these data that the number of TGN-derived, clathrin-coated vesicles found at steady state in mouse fibroblasts depends on the number of MPR molecules expressed at physiological levels.

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**Figure 4.** Membrane-bound \(\gamma\)-adaptin and MPR expression. Microsomal membranes of MPR-deficient fibroblasts were prepared as described in Materials and Methods and analyzed by Western blotting for their content in \(\gamma\)-adaptin (arrow), \(\alpha\)-adaptin, transferrin receptor, or \(\beta\)-COP (A). The amount of \(\gamma\)-adaptin bound to microsomal membranes was then quantitated and normalized to that of \(\alpha\)-adaptin (B) or transferrin receptor (C). (MPR \(-/-\), MPR-negative fibroblasts; MPR \(+/+\), control fibroblasts expressing the two MPRs; MOCK, mock-transfected MPR-negative cells; CD-4, -1, and -2, MPR-negative fibroblasts reexpressing 1.5-, 3.5-, and 4.4-fold the endogenous level of CD-MPR, respectively; CI-3 and -4, MPR-negative fibroblasts reexpressing one- and fivefold the endogenous level of CI-MPR, respectively. Values represent means \(\pm\) standard error of four independent experiments. When MPR-positive fibroblasts or MPR-negative fibroblasts reexpressing either MPR were compared with the mock-transfected MPR-negative fibroblasts, the confidence limits of the sample populations were found to be >99% in every case based on the \(t\) test.

**Figure 5.** Distribution of marker proteins on linear Ficoll/\(\text{H}_2\text{O}\) density gradients. Clathrin-coated vesicles were purified from mock-transfected MPR-negative fibroblasts (A) and MPR-negative fibroblasts reexpressing the CI-MPR (clone CI-4) (B) as described in Materials and Methods. Each fraction from the last linear density gradient was analyzed by Western blotting for its content in CI-MPR, \(\gamma\)-adaptin (arrow), \(\alpha\)-adaptin, transferrin receptor (Tf Rec.), and clathrin light chains (Clathrin L.C.).
Affinity of AP-1 for Membranes In Vitro and Clathrin-coated Vesicle Formation In Vivo

Our previous in vitro analysis based on the reexpression of CD-MPR mutants in MPR-negative fibroblasts has indicated that the high affinity binding of AP-1 to membranes ($K_d \sim 40$ nM) relies on the presence of specific determinants in the CD-MPR cytoplasmic domain (Mauxion et al., 1996). In particular, we observed that mutations introduced in the casein kinase II phosphorylation site present in its carboxyl-terminal domain (mutant A565859 according to Mauxion et al., 1996) significantly reduce the affinity of AP-1 for membranes ($K_d \sim 120$ nM) without affecting the number of AP-1 binding sites. This mutant is also drastically, but not completely, impaired in proper transport of lysosomal enzymes in vivo (35% of sorting efficiency). Therefore, these stable clones gave us the opportunity to investigate the relationship between the affinity of AP-1 for its target membrane in vitro and its efficiency in producing TGN-derived vesicles in vivo. Toward this goal, a similar type of analysis, as described above, was performed on these MPR-negative fibroblasts reexpressing a CD-MPR mutated on the casein kinase II phosphorylation site. In these stable clones, the expression level of the different marker proteins ($\alpha$-adaptin, $\gamma$-adaptin, $\beta$-COP, and transferrin receptor) was very similar to that of MPR-positive or MPR-negative fibroblasts (not shown). Fig. 8 shows that the amount of $\gamma$-adaptin recovered at steady state...
bound to membranes or recovered in clathrin-coated vesicles was far less abundant in MPR-negative fibroblasts reexpressing this CD-MPR mutant than in those reexpressing similar levels of wild-type CD-MPR. However, the amount of γ-adaptin bound to membranes or recovered associated with purified clathrin-coated vesicles was slightly higher in these cells than in MPR-negative fibroblasts. These data show that this CD-MPR mutant is unable to efficiently recruit AP-1 on membranes and to produce TGN-derived clathrin-coated vesicles. Thus, this CD-MPR mutant must be largely excluded from the AP-1–dependent pathway toward the endosomes. These results further highlight the importance of the casein kinase II phosphorylation site in the AP-1–dependent sorting of the CD-MPR and suggest that only high affinity interactions of AP-1 with its target membrane determine the production of AP-1 and clathrin-coated vesicles in vivo.

**Discussion**

Formation of clathrin-coated vesicles on TGN membranes requires the translocation of cytosolic AP-1 and clathrin. This process leads to the segregation of the MPRs and their bound lysosomal enzymes into a vesicular intermediate. Our biochemical analysis of MPR-negative fibroblasts reexpressing physiological levels of MPRs shows that the amount of AP-1 bound to membranes or recovered into clathrin-coated vesicles depends on the expression of these membrane receptors. While these results further support the notion that the MPRs are key components for the first step of clathrin coat assembly in the TGN, they demonstrate that the number of TGN-derived vesicles is determined, at least in part, by the number of membrane proteins to be sorted.

This biochemical study confirms our former proposals based on in vitro assays (Le Borgne et al., 1996) and demonstrates that, in vivo, membrane protein sorting in the TGN is tightly coupled to the formation of clathrin-coated vesicles on this organelle. If a rather low, physiological level of MPR expression can induce AP-1 recruitment and vesicle formation in cells with an MPR-negative background, it does not apparently affect the number of AP-2–coated vesicles formed at the plasma membrane. The most simple explanation is that, in these cells, the MPRs are the major membrane proteins sorted along the AP-1–dependent pathway as a result of their continuous recycling between this compartment and endosomes, whereas they probably represent a minor population (~5,000–10,000 molecules of each MPR) among the family of receptors internalized at the plasma membrane via the AP-2–dependent pathway. Thus, the mobilization of AP-2 and clathrin at the plasma membrane would probably require a massive overexpression of the MPRs as previously described for the transferrin receptor (Iacopetta et al., 1988; Miller et al., 1991). It is extremely likely that other membrane proteins sorted along the same AP-1–dependent pathway as the MPRs and sharing similar sorting signals would also contribute to recruit AP-1 on membranes and to generate AP-1–coated vesicles. MPR-negative cells still produce detectable amounts of AP-1 and clathrin-coated vesicles (~30%). In this respect, it is noteworthy that the overexpression of the major histocompatibility complex class II molecules in Hela cells also increases AP-1 binding in vitro (Salamero et al., 1996).

While this study further stresses the importance of membrane proteins, it remains possible that, in their absence, AP-1 can interact albeit weakly with membranes without producing clathrin-coated vesicles. Indeed, our former in vitro binding studies (Le Borgne et al., 1996) have shown that the membrane insertion of massive amounts of ARF-1 in its GTPγS-bound form would essentially result in the creation of low affinity (K_d ~150 nM) binding sites for AP-1, which, according to the present study using a CD-MPR mutant (see below), would not result in the efficient
production of vesicles. Thus, it is also possible that the arrival of the MPRs in the TGN would create a favorable context to stabilize AP-1 on the membrane (transition between low and high affinity interactions) to generate a transport intermediate. This model implies that AP-1 interacting with its binding sites containing the cargo membrane proteins are probably scattered in the plane of the membrane and then clustered to form a nascent transport intermediate. This clustering could be mediated either by the APs themselves, which can self-aggregate in vitro (Chang et al., 1993), or alternatively by the polymerization of clathrin triskelions bound to the APs.

Our results argue that the MPRs are rate-limiting components for TGN-derived vesicle formation at low levels of expression. However, it has to be noted that relatively high levels of MPR expression (three to fivefold the physiological level of one MPR) do not necessarily produce a corresponding increase in the amount of membrane-bound AP-1 and in the number of vesicles formed. In other words, there is not a linear relationship between high MPR expression and the number of AP-1–coated vesicles formed. This may first indicate that some steps in the recycling of the MPRs back to the TGN are rate limiting. On the other hand, clathrin-coated vesicle formation is a multistep process involving several regulatory factors. Thus, this observation could alternatively reflect a saturation of other components (or their products) required in the early steps of clathrin coat assembly. Among potential candidates are the GTPase ARF-1, potentially a phospholipase D or the mammalian homologue of the yeast Vps34p, a PI-3 kinase. A GTPase could also regulate the later steps of vesicle formation, like dynamin in the context of the plasma membrane (van der Bliek et al., 1993). This latter possibility could easily explain the differences seen after massive overexpression of the transferrin receptor in mammalian cells, producing a higher number of clathrin-coated pits at the plasma membrane in one cell type (Iacopetta et al., 1988), while producing a higher number of flat clathrin lattices without modifying the number of clathrin-coated vesicles in another cell type (Miller et al., 1991).

Sorting of membrane proteins and their segregation into specialized transport vesicles is mediated by specific determinants in their cytoplasmic domains. The MPRs contain in their cytoplasmic domains multiple determinants that mediate their sorting along the different steps of their recycling pathway. The endocytosis of the CD-MPR involves a tyrosine-based motif (at position 45) and two phenylalanines (at positions 13 and 18). This receptor as well as the CI-MPR also contain a carboxyl-terminal di-leucine–based motif critical for lysosomal targeting of hydrolases (John-son and Kornfeld, 1992; Chen et al., 1993). According to surface plasmon resonance studies performed on immobilized oligopeptides containing either a tyrosine or a di-leucine (Heilker et al., 1996), both motifs could potentially interact with both AP-1 and AP-2. Genetic approaches have also revealed that tyrosine-based motifs interact with the µ chains of APs (Ohno et al., 1995). We have previously shown that a CD-MPR containing an endocytosis motif and a di-leucine–based motif but mutated around a casein kinase II phosphorylation site adjacent to this di-leucine–based motif provides only low affinity binding sites for AP-1 in vitro and exhibits a reduced capability of trans-

We show here that the expression of such a mutant in MPR-negative fibroblasts does not result in an efficient production of AP-1–coated vesicles in vivo, indicating that this CD-MPR mutant is largely excluded from the AP-1–dependent pathway. These results further stress the functional importance of this casein kinase II phosphorylation site in regulating the AP-1–dependent sorting of the CD-MPR. It is possible that the phosphorylation of this site could induce conformational changes in the CD-MPR cytoplasmic domain to expose the cryptic tyrosine- and di-leucine–based motifs. A similar casein kinase II phosphorylation site is found at the carboxyl-terminal domain of the other CI-MPR and has also been shown to be implicated in lysosomal enzyme transport in vivo (Chen et al., 1993). It could be anticipated from our results that this phosphorylation site in the CI-MPR also regulates its segregation into AP-1–coated vesicles.

Our former and present studies as well as those performed by others in the context of the plasma membrane (Iacopetta et al., 1988; Miller et al., 1991) argue that membrane proteins containing specific sorting motifs in their cytoplasmic domains and concentrating soluble ligands via their luminal domains play a critical role in the formation of clathrin-coated vesicles. Could this represent a general rule for the formation of transport intermediates? Some aspects in the assembly of clathrin and non-clathrin coats, namely COPI and COP II functioning in vesicular traffic in the early secretory pathway (Rothman and Wieland, 1996; Schekman and Orci, 1996), argue that they could be related processes. Both COPI (Palmer et al., 1993) and AP-1 (Stammes and Rothman, 1993; Traub et al., 1993) recruitments onto membranes are regulated by the small GTPase ARF-1, whereas COPII assembly is regulated by the GTPase Sar-1p (Barlowe et al., 1993). The question of whether transmembrane proteins are essential partners in the recruitment of non-clathrin coats is still unresolved. It is interesting to note, however, that both COPI- and COPII-coated vesicles are enriched in members of the p24 family of transmembrane proteins. The latter have been proposed to function as putative cargo receptors, selecting proteins in COPI and COPII-coated vesicles (Stamnes et al., 1995; Schimmoller et al., 1995). These 16 homologous transmembrane proteins contain in their cytoplasmic domain a monoo- or a di-phenylalanine motif that was found to interact in vitro with the F subcomplex of COPI containing β, γ, and ζ COP (Fiedler et al., 1996). A subset of these p24 proteins also contains in their cytoplasmic domains carboxyl-terminal di-lysine motifs known to function in the retrieval of transmembrane proteins from the Golgi complex back to the ER (Nilsson et al., 1989; Jackson et al., 1990). These retrieval motifs, first shown to interact both in vitro and in vivo with COPI (Cosson and Letourneur, 1994; Letourneur et al., 1994), seem to interact in vitro with the B subcomplex of COPI containing α, β’, and ε COP (Fiedler et al., 1996). Thus, these different studies suggest a link between a given type of coat and specific sorting determinants in the cytoplasmic domain of these transmembrane proteins. It would be interesting to know whether these p24 transmembrane proteins also participate in the active recruitment of COPs (or subcomplexes) on their target membranes, as we report here for the MPRs and AP-1.
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