Caveolin Transfection Results in Caveolae Formation but Not Apical Sorting of Glycosylphosphatidylinositol (GPI)-anchored Proteins in Epithelial Cells

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Abstract. Most epithelial cells sort glycosylphosphatidylinositol (GPI)-anchored proteins to the apical surface. The “raft” hypothesis, based on data mainly obtained in the prototype cell line MDCK, postulates that apical sorting depends on the incorporation of apical proteins into cholesterol/glycosphingolipid (GSL) rafts, rich in the cholesterol binding protein caveolin/VIP21, in the Golgi apparatus. Fischer rat thyroid (FRT) cells constitute an ideal model to test this hypothesis, since they missort both endogenous and transfected GPI-anchored proteins to the basolateral plasma membrane and fail to incorporate them into cholesterol/glycosphingolipid clusters. Because FRT cells lack caveolin, a major component of the caveolar coat that has been proposed to have a role in apical sorting of GPI-anchored proteins (Zurzolo, C., W. Van’t Hoff, G. van Meer, and E. Rodriguez-Boulan. 1994. EMBO [Eur. Mol. Biol. Organ.] J. 13:42–53.), we carried out experiments to determine whether the lack of caveolin accounted for the sorting/clustering defect of GPI-anchored proteins. We report here that FRT cells lack morphological caveolae, but, upon stable transfection of the caveolin1 gene (cav1), form typical flask-shaped caveolae. However, cav1 expression did not redistribute GPI-anchored proteins to the apical surface, nor promote their inclusion into cholesterol/GSL rafts. Our results demonstrate that the absence of caveolin1 and morphologically identifiable caveolae cannot explain the inability of FRT cells to sort GPI-anchored proteins to the apical domain. Thus, FRT cells may lack additional factors required for apical sorting or for the clustering with GSLs of GPI-anchored proteins, or express factors that inhibit these events. Alternatively, cav1 and caveolae may not be directly involved in these processes.

Epithelial cells are characterized by the presence of polarized plasma membrane domains with different compositions of proteins and lipids (Rodriguez-Boulan and Powell, 1992; Eaton and Simons, 1995; Drubin and Nelson, 1996). In recent years, several sorting signals have been identified that mediate localization of proteins to apical or basolateral plasma membrane domains (Mostov et al., 1992; Matter and Mellman, 1994; Le Gall et al., 1995). Whereas basolateral signals are short, discrete sequences localized in the cytoplasmic domain of the protein, the best characterized apical signal is a glycosphingolipid, glycosylphosphatidylinositol (GPI)1 (Lisanti and Rodriguez-Boulan, 1990), which is used by some proteins as an anchor to the membrane bilayer (Cross, 1987; Ferguson and Williams, 1988; Low and Saltiel, 1988; Low, 1989; Doering et al., 1990; Lisanti et al., 1990). GPI-anchored proteins are selectively localized to the apical membrane of most epithelial cells studied to date (Lisanti et al., 1988; Ali and Evans, 1990; Lisanti et al., 1990; Wilson et al., 1990). Furthermore, a GPI anchor is sufficient to target recombinant GPI-anchored proteins to the apical membrane of MDCK cells (Brown et al., 1989; Lisanti et al., 1989).

Attachment of the GPI moiety occurs in the luminal face of the endoplasmic reticulum by enzymatic replacement of COOH-terminal sequences that act as signals for GPI anchoring (for review see Englund, 1993; McConville and Ferguson, 1993; Vidugiriene and Menon, 1995). The newly synthesized GPI-anchored proteins are then transported to the cell surface, where they are exposed on the topologically equivalent extracytoplasmic face of the plasma membrane (Vidugiriene and Menon, 1994). Sorting of GPI-anchored proteins occurs after their carbohydrates are processed in the Golgi complex (Brown et al., 1989;...
Lisanti et al., 1989), presumably by incorporation into post-Golgi vesicles assembled in the TGN (Lisanti and Rodriguez-Boulan, 1990; Wandinger-Ness et al., 1990).

As they migrate through the proximal Golgi complex, GPI-anchored proteins undergo a dramatic change in their biophysical properties, reflected by their becoming insoluble in certain nonionic detergents, such as Triton X-100 (TX-100) (Brown and Rose, 1992; Garcia et al., 1992; Zurzolo et al., 1994). This appears to reflect the association of GPI-anchored proteins with glycosphingolipid–cholesterol clusters in the Golgi complex, which are also detergent insoluble (Thompson and Tillack, 1985). When purified by flotation in sucrose density gradients, these aggregates, denoted TIIFF (Triton-insoluble floating fraction; Kurzchalia et al., 1995) or detergent-insoluble glycosphingolipid-enriched domains (DIG; Parton, 1996) can be shown to be rich in GPI-anchored proteins, sphingomyelin, glycosphingolipids (GSLs) and cholesterol (Brown and Rose, 1992; Garcia et al., 1993; Sargiacomo et al., 1993; Zurzolo et al., 1994). Fluorescence energy transfer experiments indicate that GPI-anchored proteins are still clustered when they arrive at the cell surface, but slowly disperse in the next few hours (Hannan et al., 1993).

The “raft hypothesis” postulates that clustering with GSLs is required for the sorting of apical proteins (Simons and Ikonen, 1997; van Meer and Simons, 1988). GSLs are sorted apically in MDCK (kidney) and Caco-2 (intestinal) epithelial cells, at least as indicated by experiments with short acyl chain fluorescent glycolipids (van Meer et al., 1987; van’t Hof and van Meer, 1990). In its extended version, the raft includes transmembrane apical proteins, such as influenza hemagglutinin, which also becomes detergent insoluble as it traverses the Golgi complex (Skibbens et al., 1989). The mechanisms involved in the formation of the raft are still obscure. The length and saturation of the acyl chains of GSLs appear to be important in this process (Schroeder et al., 1994). A similar GSL–cholesterol enrichment as in the Golgi raft was reported for plasma membrane caveoleae (Fra et al., 1994; Gorodinski et al., 1995; Kurzchalia et al., 1995; Mayor et al., 1995; Parton, 1996; Schnitzer et al., 1995b). Initial reports suggested that GPI-anchored proteins are also enriched in caveoleae (Rotheberg et al., 1990; Sargiacomo et al., 1993); however, recent evidence suggests that GPI-anchored proteins are distributed evenly throughout the plasma membrane, but redistribute to caveolar regions upon cross-linking, e.g., with antibodies (Mayor et al., 1994; Parton et al., 1994; Schnitzer et al., 1995a). Caveoleae are coated on the cytoplasmic side by a coat that includes a phosphorylated cholesterol-binding protein of 21 kD, caveolin (Rotheberg et al., 1992; Murata et al., 1995). Recent transient transfection experiments in lymphocytes indicate that caveolin is required for the assembly of morphological caveoleae (Fra et al., 1995). The observations that caveolin is identical to VIP21, a protein present in post-Golgi vesicles and the TGN (Wandinger-Ness et al., 1990; Kurzchalia et al., 1992), and that caveolin localizes into TIIFF (Sargiacomo et al., 1993; Zurzolo et al., 1994; Kurzchalia et al., 1995), suggest a possible involvement of this protein in the formation of post-Golgi transport vesicles (Dupree et al., 1993). However, caveoleae have not been identified in the TGN to date.

The polarized epithelial cell line Fischer rat thyroid (FRT) provides an excellent model to study the role of caveolin in both GPI-anchored protein sorting and caveolar assembly (Zurzolo et al., 1992; Nitsch et al., 1985). FRT cells do not express caveolin (Sargiacomo et al., 1993; Zurzolo et al., 1994) and do not form caveoleae (this paper). Although FRT cells can sort transmembrane apical proteins to their correct plasma membrane domain (Zurzolo et al., 1992), they deliver endogenous and transfected GPI-anchored proteins and fluorescent GSLs to the basolateral membrane and fail to incorporate GPI proteins into TIIFF (Zurzolo et al., 1993, 1994). In this paper, we report the results of experiments in which we transfected caveolin1 (cav1) into FRT cells to examine its effect on GPI-anchored protein sorting, clustering, and caveolar assembly. Although caveolin was sufficient to promote the assembly of morphologically normal caveoleae in FRT cells, it failed to redistribute GPI proteins to the apical surface or to affect their clustering into TIIFF.

Materials and Methods

Reagents and Antibodies

Cell culture reagents were purchased from Gibco Laboratories (Grand Island, NY). Protein A-Sepharose was from Pharmacia Diagnostics AB (Uppsala, Sweden), sulfo-NHS derivatives and streptavidin–agarose beads were from Pierce Chemical Co. (Rockford, IL). The monoclonal antibody (MCA-404) against herpes simplex virus was from Scriotec Ltd. (Kidlington, Oxford, United Kingdom), anticaeoealin polyclonal antibody was from Transduction Laboratories (Lexington, KY). Affinity-purified antibodies (rabbit anti-mouse IgG) were purchased from Cappel Laboratories (Malvern, PA). All other reagents were obtained from Sigma Chemical Co. (St. Louis, MO).

Cell Culture and Transfection

FRT cells stably expressing gD1–DAF (Zurzolo et al., 1993), and FRT cells expressing transfected cav1 were grown in F12 Coon’s modified medium containing 5% FBS. MDCK cells were maintained in DME supplemented with 5% FBS. FRT cells were cotransfected as previously described using a modification of the calcium phosphate precipitation procedure (Zurzolo et al., 1993). The VIP21/cav1 cDNA from MDCK cells was a gift from P. Dupree (European Molecular Biology Laboratory, Heidelberg, Germany, [EMBL]) and was inserted into pCMV5, carrying the gene for resistance to G418, using the EcoRI and KpnI restriction sites. For some experiments we used myc–caveolin cDNA (from R.G.W. Anderson, University of Texas Health Science Center, Dallas, TX). The construct containing gD1–DAF under the Rous sarcoma virus promoter was a gift of I. Caras (Genentech Inc., South San Francisco, CA) and has been described elsewhere (Lisanti et al., 1989; Zurzolo et al., 1993).

Biotinylation Assays

Confluent monolayers on transwells were labeled overnight using 1 mCi/ml of [125I]sodium iodide (Amersham Corp., Arlington Heights, IL), and were biotinylated and processed for immunoprecipitation as previously described (Le Bivic et al., 1989; Sargiacomo et al., 1989; Zurzolo et al., 1993). To recover the immunoprecipitated biotinylated antigens, the beads were boiled in 10 ml of 10% SDS, diluted with lysis buffer, and then centrifuged for 1 min at 14,000 rpm. Biotinylated antigens present in the supernatants were precipitated with streptavidin–agarose beads. Finally, the beads were boiled in Laemmli buffer (Laemmli, 1970) and analyzed by SDS-PAGE. Dried gels were processed as described (Zurzolo et al., 1992) using preflashed films; densitometry analysis was carried out within the linear range of the film. The steady-state distribution of endogenous GPI-anchored proteins was determined using domain-selective biotinylation and treatment with phospholipase C, as previously described (Lisanti et al., 1988; Zurzolo et al., 1993).

Pulse-chase and TX-100 Extraction

TX-100 extractability during pulse-chase experiments was assayed as pre-
viously described (Brown and Rose, 1992; Zurzolo et al., 1994). Briefly, cells that had just reached confluency in 35-mm dishes were starved of methionine and cysteine for 20 min, pulse labeled for 5 min with 100 ml of pulse medium containing ~500 μCi/ml of [35S]met-cys, and then incubated in chase medium (DME containing 10% FBS and 100X met and cys) for different times. After each time point, cells were washed twice with PBS containing 1 mM CaCl2 and 1 mM MgCl2 (PBS CM) on ice as described previously, and then lysed for 20 min on ice in 1 ml TNE/TX-100 buffer (Brown and Rose, 1992; Zurzolo et al., 1994). Lysates were collected and centrifuged at 13,000 rpm for 2 min at 4°C. Supernatants, representing the soluble material, were removed and then pellets were solubilized in 100 ml of solubilization buffer (50 mM Tris-HCl, pH 8.8, 5 mM EDTA, 1% SDS); DNA was sheared through a 22-g needle. Both soluble and insoluble materials were adjusted to 0.1% SDS before immunoprecipitation with specific antibodies as previously described (Le Bivic et al., 1989).

**Sucrose Gradients**

Sucrose gradient analysis of TX-100–insoluble residue was performed using previously published protocols (Brown and Rose, 1992; Zurzolo et al., 1994). Briefly, cells were grown to confluence in 100-mm dishes, labeled for 30 min with 500 μCi/ml of [35S]met-cys, and then incubated in chase medium for 3 h. Monolayers were then rinsed in PBS CM and lysed for 20 min in TNE/TX-100 buffer on ice. Lysates were scraped from the dish, brought to 40% sucrose, and then placed at the bottom of a centrifuge tube. A linear sucrose gradient (5–35% in TNE) was layered on top of the lysates and then the samples were centrifuged at 39,000 rpm for 18–20 h in a rotor (model SW41; Beckman Instrs., Fullerton, CA). 1-ml fractions were harvested from the top of the gradient. Immunoprecipitation of gD1–DAF and other antigens was performed on the different fractions, after bringing them to ~20% sucrose and 1% TX-100. Samples were solubilized in Laemml buffer and either boiled for 5 min, or left at 25°C for 30 min before running on SDS-PAGE, and were then analyzed by autoradiography.

**Immunofluorescence and Electron Microscopy**

Cells grown on coverslips to confluence were processed for immunofluorescence as described (Zurzolo et al., 1993). For EM experiments, cells were washed three times with PBS pH 7.4, containing 2.7 mM KCl, 1 mM CaCl2, and 1 mM MgCl2, and then fixed for 30 min at room temperature with 2.5% glutaraldehyde, 0.1% tannic acid in 0.1 M cacodylate buffer, pH 7.4. Cells were then rinsed three times in 0.1 M cacodylate buffer and postfixed in 1% osmium tetroxide in 0.1 M cacodylate buffer for 30–60 min at room temperature. Cells were further stained en bloc with 3% uranyl acetate in 0.1 M cacodylate buffer for 90–120 min at room temperature and finally embedded in Epon 812 (Polysciences, Inc., Warington, PA). Thin sections were cut parallel to the plane of the culture on a microtome (model MT 5000; Sorvall, Inc., Newtown, CT) and then stained with 1.5% uranyl acetate and 0.1% lead citrate (Venable et al., 1965). For the immunogold localization of gD1–DAF, cav1–FRT cells grown to confluence on coverslips were incubated on ice with the monoclonal antibody anti-gD1–DAF for 1 h. After three washes in PBS, bound antibody was detected with goat anti-mouse IgG 10-nm gold conjugates. After three washes in PBS, the cells were processed as described above through to dehydration in ethanol. In 70% ethanol, cells were scraped from the coverslips with a razor blade and then the pellets were collected in a microtuge tube before embedding in Epon 812. Specimens were examined and photographed in either a 100 C X II (JEOL USA, Inc., Peabody, MA) or electron microscope (model 400T; Philips Electron Optics, Eindhoven, The Netherlands).

**Results**

**Transfection of gD1–DAF and cav1 in FRT Cells**

We have previously shown that unlike MDCK cells (Lisanti et al., 1989), FRT cells sort gD1–DAF, a fusion protein between the ectodomain of the herpes simplex virus glycoprotein gD1 and the GPI-attachment signal from decay accelerating factor (DAF), to the basolateral membrane (Zurzolo et al., 1993) and fail to incorporate it into TIFF (Zurzolo et al., 1994). Because FRT cells do not express caveolin, a cholesterol binding protein, the possibility was raised that caveolin may be required for both association of GPI-anchored proteins with TIFF and for targeting of GPI-anchored proteins to the apical surface (Dupree et al., 1993; Zurzolo et al., 1994).

To determine whether caveolin was involved in either the apical sorting of gD1–DAF or in its clustering with GSLs, we stably transfected FRT cells with a cDNA encoding the cav1 protein. After initial attempts to transfect cav1 into FRT cells already expressing gD1–DAF failed, we simultaneously transfected a plasmid containing the gD1–DAF gene under control of the RSV promoter (Lisanti et al., 1989), and then a plasmid containing the cav1 coding sequence fused to the cytomegalovirus (CMV) promoter and the gene conferring resistance to neomycin (G418). By selection with G418, we isolated different clones expressing both proteins. Among many clones heterogeneous in their expression of caveolin and gD1–DAF proteins, as visualized by double immunofluorescence experiments (data not shown), we selected two clones (FRTcl1 and FRTcl2) that displayed a homogeneous distribution of both markers within all cells (Fig. 1 A and B). Both clones displayed caveolin as a punctate pattern at the plasma membrane and within the Golgi apparatus (Fig. 1 A). gD1–DAF was similarly enriched at the plasma membrane, albeit with a different, more diffuse, pattern (Fig. 1 B), but was also found intracellularly, as previously shown (Zurzolo et al., 1993).

**gD1–DAF Is Sorted to the Basolateral Membrane in cav1-transfected FRT Cells**

The immunofluorescence experiments demonstrated a characteristic ring-like basolateral distribution of surface gD1–DAF in cav1–FRT cells, identical to the localization we had previously observed in nontransfected cells (Zurzolo et al., 1993). To quantify this distribution, we performed a domain selective–biotinylation assay with both clones expressing caveolin, and with nontransfected cells for comparison. In agreement with the distribution observed by immunofluorescence, we found that gD1–DAF was basolaterally localized in both cav1 expressing clones and in the wild-type cells (Fig. 2 A). In all cells we found that at steady state, 90–95% of the total surface protein was localized on the basolateral membrane. Furthermore, immunoprecipitation experiments demonstrated that the levels of cav1 in clone 1 and clone 2 were 90 and 60%, respectively, of the levels found in MDCK cells (Fig. 2 B).

As additional controls, we also examined the plasma membrane distribution of several endogenous apical (DPPIV) and basolateral (35–40-kD Ag, NaK–ATPase) antigens (Zurzolo et al., 1992, Zurzolo and Rodriguez-Boulan, 1993) in the caveolin-transfected cells by immunofluorescence and surface biotinylation. The polarity of these markers was unchanged compared with nontransfected cells (data not shown). These data clearly demonstrated that cav1 expression did not revert the basolateral sorting of gD1–DAF in FRT cells.

**gD1–DAF Is Excluded from TIFF in FRT Cells Expressing Cav1**

In MDCK cells, GPI-anchored proteins are incorporated into TX-100–insoluble aggregates, presumably reflecting their association with GSLs in the Golgi complex (Brown et al., Caveolin and Sorting of GPI-anchored Proteins, 619
and Rose, 1992). In apparent agreement with the idea that this association may be required for their apical sorting (Van Meer and Simons, 1988; Lisanti and Rodriguez-Boulan, 1990), we previously demonstrated that basolaterally sorted gD1–DAF fails to be incorporated into TIFF in FRT cells, whereas it is apically sorted and becomes detergent insoluble in MDCK cells (Zurzolo et al., 1994). To determine whether transfection of \textit{cav1} promotes clustering of gD1–DAF with GSLs, we performed pulse-chase experiments in the two FRT clones stably expressing \textit{cav1}, and then determined whether gD1–DAF was incorporated into the TX-100–insoluble fraction. We found that in both caveolin expressing clones, gD1–DAF was mainly soluble in TX-100 at all times of chase (Fig. 3), exactly as previously shown in nontransfected cells (Zurzolo et al., 1994). These experiments indicated that transfection of \textit{cav1} was not able to reverse the inability of gD1–DAF to partition with TIFF in FRT cells.

**Caveolin Forms Oligomers and Associates with TIFF in Transfected FRT Cells**

To rule out the possibility that the lack of effect of cav1 on the partitioning of gD1–DAF in TIFF was because of the inability of caveolin itself to localize in the insoluble fractions in the transfected FRT clones, we performed a sucrose density–gradient purification of TIFF in clone 2 was reproducible and corresponded to the small quantity of insoluble gD1–DAF found in this clone (Fig. 3). This result was not related to different levels of caveolin expressed by the two clones, but is simply a consequence of clonal variation among FRT cells. In fact, similar amounts of insoluble and floating gD1–DAF were found in the nontransfected FRT population lacking caveolin, as was previously reported (Zurzolo et al., 1994). These experiments indicated that transfection of \textit{cav1} was not able to reverse the inability of gD1–DAF to partition with TIFF in FRT cells.

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These experiments also revealed that similar high molecular weight complexes resistant to boiling were found in the two cell lines. To better resolve these complexes on the gel and to follow their migration on the gradients, we performed a similar experiment in the two cell lines without boiling the samples. These experiments clearly showed that transfected caveolin in FRT cells formed oligomers of molecular weights of $\approx 350$, $300$, and $200$ kD, as well as $45$-kD dimers (Fig. 6, top), similar to that formed by endogenous caveolin in MDCK cells (Fig. 6, bottom) as previously shown (Monier et al., 1995; Sargiacomo et al., 1995). Furthermore, these oligomers appeared to migrate to TIF in both cell lines in similar amounts as the monomer (compare Figs. 5 and 6, top and bottom).

Endogenous GPI-anchored Proteins Distribute Basolaterally in Wild-Type and cav1-transfected FRT Cells

To determine whether caveolin transfection affected the polarity of endogenous GPI-anchored proteins in FRT cells, we studied their steady-state localization using domain-selective biotinylation in combination with partition into Triton X-114 and treatment with phospholipase C (Lisanti et al., 1988). As shown in Fig. 7, the majority of endogenous GPI-anchored proteins were basolaterally localized in the cav1-expressing cells. This result was identical to that obtained in nontransfected cells (Zurzolo et al., 1993). Therefore, caveolin expression appears to have no effect on GPI-anchored protein distribution, which remains predominantly basolateral, exactly as in FRT cells not expressing caveolin.

Cav1 Promotes Formation of Caveolae in Transfected FRT Cells

Previous experiments showed that transient expression of VIP21/cav1 using a Semliki Forest virus vector in lymphocytes lacking caveolin promoted the formation of plasmalemmal caveolae (Fra et al., 1995). Furthermore, it was suggested that the formation of large caveolin oligomers might be required for caveolar assembly (Parton and Simons, 1995; Sargiacomo et al., 1995). To definitively address the role of caveolin in caveolar formation, we assayed for the presence of caveolae in the stably transfected and nontransfected FRT cells. Using 0.1% tannic acid to enhance visualization of caveolar invaginations by EM (Palade and Bruns, 1968), we observed that wild-type FRT cells have no plasmalemmal caveolae (Fig. 8, A and B). In contrast, FRT cells expressing caveolin displayed large numbers of normal flask-shaped caveolae (Palade and Bruns, 1968), frequently organized into characteristic racemose clusters (Fig. 8, C–E). These data showed that cav1 was necessary and sufficient to promote efficient caveolae formation in FRT cells.

To determine whether gD1–DAF was able to localize in these caveolae at the surface of the transfected FRT cells, we performed an immuno-EM localization of gD1–DAF after cross-linking with antibodies. We found that gD1–DAF was distributed upon the entire surface of transfected FRT cells, and that it was able to cluster in caveolae upon antibody cross-linking (Fig. 9, A–D) as was previously shown in other cell lines (Mayor et al., 1994; Parton et al., 1994; Mayor and Maxfield 1995). These experiments indicated that in FRT cells, the newly formed caveolae are...
able to interact with GPI-anchored proteins and, together with our other data, suggest that cav1 and caveolae are not involved in GPI-anchored protein trafficking.

**Discussion**

The raft hypothesis postulates that apically localized plasma membrane proteins laterally associate with GSLs into segregated microdomains as a prerequisite for sorting into apical transport vesicles in the TGN (Simons and Ikonen, 1997). The most frequently used method to identify these lipid microdomains and to follow the partitioning of a protein within them is to determine whether the lipids and the proteins are insoluble in certain nonionic detergents (TX-100, TX-114, CHAPS) and whether they co-float into the lighter density region (TIFF or DIG) of sucrose density gradients (for review see Kurzchalia et al., 1995; Parton, 1996). Although different transmembrane apical proteins show variable association with TIFFs according to this criterion, GSLs and GPI-anchored proteins appear to be consistently incorporated into TIFFs in different epithelial cells, where they are apically targeted (Lisanti and Rodriguez-Boulan, 1990; Brown and Rose, 1992; Mirre et al., 1996). From this perspective, GPI-anchored proteins appear to require GSL association for apical targeting. Our previously published data, showing the failure of FRT cells to target GPI-anchored proteins and GSLs apically and to incorporate GPI-anchored proteins into TIFF, provides additional support for this hypothesis (Zurzolo et al., 1993, 1994). Therefore, if GPI-anchored proteins are indeed cosorted, the question arises as to what is the mechanism that promotes their association in microdomains at the level of the TGN and then leads to their apical sorting.

Recent circumstantial evidence suggests that the caveolar coat protein caveolin might play a role in the clustering of GSLs and GPI-anchored proteins (Dupree et al., 1993; Sargiacomo et al., 1993; Zurzolo et al., 1994). Although TIFF exists in the absence of plasmalemmal caveolae, the lipid composition of caveolae is very similar to that of TIFF (Brown and Rose, 1992; Fra et al., 1995). Furthermore, it has been shown that caveolin, which is 90% enriched in plasma membrane caveolae, associates with TIFF (Sargiacomo et al., 1993; Kurzchalia et al., 1995). Although some controversy exists as to whether GPI-anchored proteins are enriched within caveolae (Rothberg et al., 1990; Anderson, 1993; Mayor et al., 1994), nonetheless, these proteins show affinity for caveolae after they have been cross-linked with antibodies. Finally, the presence of caveolin in the TGN and post-TGN vesicles (Kuzchalia et al., 1992), and the absence of caveolin in FRT cells, which exhibit defective apical targeting and clustering of GPI-anchored proteins, further underlines the need for studies that address the role of caveolin in intracellular sorting of proteins in the Golgi complex.

The approach we used in this report was to transfect the cav1 gene into FRT cells to determine its effect on the sorting of GPI-anchored proteins and its association with TIFF. In two FRT clones stably expressing caveolin at levels comparable with MDCK cells, we observed that although caveolin typically associated with TIFF and formed large oligomers similar to those assembled in MDCK cells, gD1–DAF remained unable to cluster with GSLs that independently fractionated into TIFF, and was still sorted to the basolateral membrane, as in wild-type cells. Furthermore, no changes were detected in the distribution of endogenous GPI-anchored proteins, which remained mainly basolateral or nonpolar. Taken together, our data suggest that cav1 does not mediate the association of GPI-anchored proteins with GSL-enriched microdomains in the Golgi complex, nor participates in apical sorting.

Another possible explanation for our results is that cav1 is, for some reason, nonfunctional in FRT cells and, therefore, fails to carry out the functions that it performs normally in other cells. For example, FRT cells might not be able to perform a critical posttranslational processing modification of cav1, or might fail to form the oligomers observed in MDCK cells. To test this hypothesis, we studied the effect of cav1 transfection on the assembly of plasma membrane caveolae. Indeed, we found that wild-
type FRT cells do not form caveolae, but, upon transfection with \textit{cav1}, formed large numbers of normal flask-shaped caveolae. Our results confirm and extend previous experiments in which caveolin transfection was shown to be necessary and sufficient to promote the assembly of caveolae in lymphocytes (Fra et al., 1995). FRT caveolae were indistinguishable from the caveolae endogenously present in other epithelial and endothelial cells; they frequently showed a classical septum at the neck and often associated into racemose clusters of five or six units (Fig. 8) (Palade and Bruns, 1968). We also found that gD1–DAF clustered into these newly formed caveolae upon cross-linking with antibodies. Furthermore, cav1 transfected into FRT cells forms high molecular weight oligomers as in cells that form caveolae, such as MDCK cells. Therefore, our results show that both cav1 and caveolae are functional in FRT cells, as judged by (a) the ability of cav1 to oligomerize and promote caveolar assembly, and (b) by the capacity of these organelles to interact with GPI-anchored proteins. This is the first report of caveolar assembly in cells permanently transfected with caveolin.

Although our data indicate that cav1 is functional and sufficient to promote the assembly of caveolae in FRT cells, they do not exclude the possibility that FRT cells lack additional factor(s) required for GPI-anchored protein clustering with GSLs, enriched microdomains, and/or apical sorting. For example, GPI-anchored proteins might have different intrinsic properties in FRT cells that prevent their clustering with GSLs. Alternatively, FRT cells might lack a clustering molecule that would be involved in partitioning GPI-anchored proteins into GSL clusters. However, some data already present in the literature do not favor a role for caveolin in these processes (Gorodinsky and Harris, 1995; Kurzchalia et al., 1995; Mirre et al., 1996). Most importantly, CaCo2 cells lack caveolin and are, nonetheless, able to form TX-100–insoluble GSLs/GPI-protein–enriched microdomains, and sort them to the apical plasma membrane (Mirre et al., 1996).

Because we did not check whether FRT cells also lack caveolin2 (Parton, 1996; Scherer et al., 1996), a possibility remains that this isoform of caveolin could have some role in the apical sorting of GPI-anchored proteins. However, considering that such a role was proposed originally for cav1, which was found to be enriched in the TGN and
post-TGN sorting vesicles (Dupree et al., 1993; Kurzchalia et al., 1992), and considering that caveolin2 appears to be targeted largely to the basolateral plasma membrane in MDCK cells (Scheiffele et al., 1998), we believe this scenario is unlikely.

The data reported here support the hypothesis that caveolin oligomerization is the basis of caveolar coat assembly (Anderson, 1993; Fra et al., 1995; Monier et al., 1995) and clearly demonstrate that cav1 is necessary and sufficient to promote formation of these characteristic plasma membrane invaginations. Furthermore, because we found similar levels of TX-100–insoluble GSLs in FRT cells expressing or not expressing cav1 and caveolae, our data definitively rule out the suggested identity between caveolae and detergent-insoluble microdomains, and do not support a role for caveolae as carrier vesicles for the transport of GPI-anchored proteins to the apical plasma membrane.

An alternative model is that clustering with GSLs is indeed a prerequisite for apical sorting of GPI-anchored proteins, and that caveolin and caveolae are not involved in these events. Although it is not yet clear whether incorporation of a protein in these rafts would be sufficient for apical sorting, there could be different mechanisms mediating the association of GPI-anchored proteins with GSLs rafts. One could hypothesize that MDCK and other epithelial cells that sort GPI-anchored proteins to the apical surface possess a clustering factor, such as was originally suggested for caveolin, that could also be recognized as an apical targeting signal. An equivalent function could be performed by the membrane-spanning domains of apical transmembrane proteins (Kundu et al., 1996). Additionally, the ectodomain may also play a role in apical sorting since it has been shown that most of the ectodomains of GPI-anchored proteins are secreted selectively into the apical medium (Brown et al., 1989; Lisanti et al., 1989; Powell et al., 1991). Further experiments will be necessary.
to discriminate among these possibilities. Since the sorting machinery seems to be conserved also in nonpolarized cells (Musch et al., 1996; Yoshimori et al., 1996), we believe that the use of in vitro assays, together with comparative analyses between different epithelial cells displaying different sorting profiles of GPI-anchored proteins and GSLs, will help to address these possibilities.

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Figure 9. Immunogold localization of gD1–DAF on the surface of caveolin-expressing FRT cells. Cav1–FRT cells were grown to confluence on coverslips, incubated with an anti-gD1–DAF monoclonal antibody on ice for 1 h and, after washing in PBS, with a secondary antibody conjugated with 10-nm colloidal gold on ice for 1 h. After fixation, cells were dehydrated through ethanol and were scraped from the coverslips in 70% ethanol. The pellets were then processed as described in Materials and Methods. A–D show different sections of apical and basolateral plasma membranes of cav1–FRT cells that contain gD1–DAF clustered in newly formed (A and B) or classical flask-shaped caveolae (C and D; small arrows). A racemose cluster of caveolae containing gD1–DAF is shown on the apical surface (D; large arrow). Arrowheads indicate the gold particles bound to the anti-gD1–DAF antibody. Bar, 100 nm.
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