Inhibition of Intracellular Cholesterol Transport Alters Presenilin Localization and Amyloid Precursor Protein Processing in Neuronal Cells

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Generation of amyloid-β (Aβ) from the amyloid precursor protein (APP) requires proteolytic cleavage by two proteases, β- and γ-secretase. Several lines of evidence suggest a role for cholesterol on secretase activities, although the responsible cellular mechanisms remain unclear. Here we show that alterations in cholesterol transport from late endocytic organelles to the endoplasmic reticulum have important consequences for both APP processing and the localization of γ-secretase-associated presenilins (PS). Exposure of neuronal cells to cholesterol transport-inhibiting agents resulted in a marked decrease in β-cleavage of full-length APP. In contrast, γ-secretase activity on APP C-terminal fragments was enhanced, increasing the production of both Aβ40 and Aβ42. Remarkably, retention of cholesterol in endosomal/lysosomal compartments induced PS1 and PS2 to accumulate in Rab7-positive vesicular organelles implicated in cholesterol sorting. Accumulation of PS in vesicular compartments was prominent in both Chinese hamster ovary cells deficient in Niemann–Pick C1 protein as well as in neuronal cells exposed to the cholesterol transport-inhibiting agent U18666A. Because Aβ42 also localized to PS1-containing vesicular compartments, organelles involved in cholesterol transport might represent an important site for γ-secretase activity. Our results suggest that the subcellular distribution of cholesterol may be an important factor in how cholesterol alters Aβ production and the risk of Alzheimer’s disease.

Key words: presenilin; cholesterol; NPC1; APP processing; Rab7; ACAT; amyloid-β

Although its primary function in vivo remains unclear, it generally is accepted that amyloid precursor protein (APP) is processed sequentially by several proteolytic events. Cleavage of the full-length precursor by α-secretase creates APPαsec, which is released into the medium, leaving behind a short membrane-bound C-terminal fragment (αCTF). Less frequently, the holoprotein is cleaved at a slightly more N-terminal site, generating βCTF and APPβsec. APP CTFs can be processed further by γ-secretase. This second cleavage event occurs within the membrane region of the C-terminal stump and requires the multi-transmembrane-spanning proteins presenilin 1 (PS1) or PS2. Cleavage at the γ-site of βCTF generates the 4 kDa fragment amyloid-β (Aβ), which represents a major component of amyloid plaques in Alzheimer’s disease (AD). Additional cleavage of αCTF yields the nonamyloidogenic peptide p3 (for review, see De Strooper and Annaert, 2000).

Increasing evidence indicates a role for cholesterol in the metabolism of APP. Carriers of the ApoE4 allele of apolipoprotein E are predisposed to an earlier onset for developing AD (Corder et al., 1993), and a high cholesterol diet accelerates Aβ deposition in transgenic mice (Refolo et al., 2000). Moreover, suppression of cholesterol neosynthesis strongly reduces the formation of Aβ species in vivo and in vitro (Simons et al., 1998; Frears et al., 1999; Fassbender et al., 2001; Refolo et al., 2001). Low-density lipoprotein (LDL) receptor family members such as LRP contribute to the endocytosis of APP, suggesting a role for internalization processes in APP metabolism (Kounnas et al., 1995). A fraction of APP molecules, as well as its secretases, localizes to cholesterol-rich membrane subdomains (Bouillot et al., 1996; Riddell et al., 2001). Recent data showing that Aβ accumulation is associated with NPC1 function (Yamazaki et al., 2001) as well as a strong correlation of cholesterol ester (CE) levels with Aβ generation (Puglielli et al., 2001) indicate that APP processing may be linked to intracellular cholesterol distribution.

In this study we investigate the role of a pathway that mediates the transfer of internalized cholesterol from late endocytic organelles to a sterol-regulating pool in the endoplasmic reticulum (ER) on APP cleavage and PS localization. Mutations in the genes encoding for NPC1 protein or the lysosomal protein HE1 cause cholesterol and other lipids to accumulate in late endosomes and lysosomes, resulting in Niemann–Pick C disease (NPC; Carstea et al., 1997; Naureckiene et al., 2000). Cells mutant in NPC1 show a block in cholesterol esterification by ER resident acyl-coenzyme A:cholesterol acyltransferase (ACAT), suggesting that NPC1 is involved actively in cholesterol transport between late endocytic compartments and the ER (Liscum et al., 1989; Cruz et al., 2000). Class-2 amphiphiles such as U18666A or imipramine mimic NPC disease by inhibiting either NPC1 pro-
tein itself or an unknown factor along its pathway (Lange et al., 2000).

Here we provide evidence for a link among intracellular cholesterol transport, Aβ generation, and PS localization. We show that the retention of cholesterol in late endosomal/lysosomal compartments is associated with major alterations in APP processing, inversely affecting β- and γ-secretase cleavage. Surprisingly, this is accompanied by an accumulation of PS1 and PS2 in Rab7-positive vesicular compartments that are directly adjacent to cholesterol-laden late endosomes under conditions in which intracellular cholesterol transport is blocked. Our results indicate that the subcellular distribution of cholesterol is an important factor in regulating the proteolytic cleavage of APP and suggest the possibility of a cholesterol-dependent trafficking of PS.

MATERIALS AND METHODS

Cell culture. SH-SY5Y cells and mouse fibroblasts were cultivated in DMEM/5% fetal calf serum (FCS). Mutant Chinese hamster ovary (CHO) cells were grown in Ham-F12/5% FCS. Rab7-GFP-transfected A431 cells were grown in DMEM/5% FCS. Rab7 protein was fused at its N terminus to the C terminus of GFP with the pEGFP-C3 vector provided by Clontech Laboratories (Cambridge, UK). Stable cell lines were selected and maintained in medium containing 0.5 mg/ml G418 (Sigma-Aldrich, Munich, Germany). For the time of exposure to 50 μg/ml LDL (Sigma-Aldrich) and 3 μg/ml U18666A [3-β-2-(diethylamino)ethoxy]androst-5-en-17-one (Biomol, Hamburg, Germany); from a stock solution in ethanol stored at −20°C for <3 months] or 80 μM imipramine (Sigma-Aldrich), the FCS levels were increased to 10% to stimulate cholesterol internalization (Tabas et al., 1998).

Preparation of primary hippocampal neurons from 17-d-old fetal rats was performed as described previously (Gaslin and Banker, 1991). Neurons were plated on poly-L-lysine-coated plastic dishes (Nunc, Dänstadt, Germany) or on poly-L-lysine-coated glass coverslips and cultivated under serum-free conditions in MEM containing N2 and B27 supplements (Invitrogen, San Diego, CA). E14 mixed cortical neurons were prepared and cultivated by following the same protocol, with cerebral cortices used for 12–15 d; cortical neurons were used after 7–10 d in culture.

Antibodies and fluorescent dyes. The following primary antibodies have been described previously: polyclonal anti-PS1 N-terminal fragment (N-18) antibody (Papini et al., 1996), monoclonal 6C4-antibody to lysosomal acid lipase (Papini et al., 1996), polyclonal anti-PS1 antibody 95.23 (diluted 1:10,000 in PBS) or monoclonal anti-PS1 antibody G2-11 (4:10,000) specific for the C-terminal residues of PS1 and PS2, respectively (Ida et al., 1996). Polyclonal antibody G2-10 (2:10,000) was provided by Centers for Proteolysis of Diseases (Center for Proteolysis of Diseases, University of Heidelberg, Germany). Oil red O (Sigma-Aldrich) was a gift from J. McLaughlin (MRc Viology Unit, University of Glasgow, UK). Monoclonal anti-calnexin and monoclonal anti-βI/βII/GRP78 antibodies were from Transduction Laboratories (BD Biosciences, Heidelberg, Germany), monoclonal anti-PS1-loop antibody was from Chemicon (Hofheim, Germany), and polyclonal PS2 antibody was from New England Biolabs (Frankfurt, Germany). For the simultaneous detection of cholesterol, primary antibodies were dissolved in 125 μl/mg filipin (Sigma-Aldrich). Mouse anti-rabbit and rabbit anti-mouse IgG (Alexa-488- or Alexa-568-conjugated) secondary antibodies were from Molecular Probes (Leiden, Netherlands). Horseradish peroxidase-conjugated rabbit anti-mouse or goat anti-rabbit secondary antibodies were from Dako A/S (Glostrup, Denmark).

Immunofluorescence, confocal microscopy, and image processing. Fully polarized hippocampal neurons, SH-SY5Y cells, mutant CHO cells, or A431 cells were incubated with 50 μg/ml LDL in the presence or absence of 0.75 or 3 μg/ml U18666A for 24 hr. Immunofluorescence labeling was performed according to the standard procedures; cells grown on glass coverslips were fixed in 3% paraformaldehyde for 10 min at 4°C and permeabilized with either 0.05% saponin (4°C for 1 min) or a brief dip into ice-cold methanol (−20°C). Alternatively, fixation and permeabilization were performed in ice-cold acetone (−20°C for 30 sec). Primary antibodies were centrifuged to exclude nonspecific aggregation, and the cells were incubated for 30 min at 4°C. After incubation with Alexa-conjugated secondary antibodies the cells were mounted in Mowiol (Calbiochem-Novabiochem, Bad Soden, Germany).

Samples were viewed with a Zeiss LSM 510 confocal microscope (Carl Zeiss, Oberkochen, Germany). Images were acquired with a 63× or 100× Plan-Apochromat III differential interference contrast objective (Carl Zeiss). Pinhole settings were −1 airy unit for all images except for the detection of filipin (2-3 airy units). Fluorescent dyes were imaged sequentially in frame interlace mode to eliminate cross-talk between the channels. Images were sampled accordingly to satisfy the Nyquist criteria for all dimensions and were processed further with Adobe Photoshop (Mountainview, CA).

Lipid extraction and analysis. Fresh human LDL was mixed with 4[14C]cholesterol (Amersham Pharmacia Biotech, Braunschweig, Germany) in ethanol (final concentration <0.1%) and dissolved in labeling medium to a final concentration of 50 μg/ml and a total 14C activity of 0.5 μCi/ml. Cells exposed to U18666A or imipramine and the controls were incubated for 2–24 hr at 37°C, 5% CO2. With the removal of the medium the cells were washed twice with PBS, and the lipids were extracted with hexane/isopropanol (3:2) for 30 min at 4°C. Solvent was evaporated under a gentle N2 stream; dried lipids were resolved in a small aliquot of hexane/isopropanol and subjected to unidimensional thin-layer chromatography (TLC) on general purpose silica gel plates (Sigma-Aldrich) in toluene/ethylacetate (2:1). Quantitative analysis of cholesterol ester formation and internalization of 14C-cholesterol were performed after film exposure of TLC plates and densitometric scanning with MacBAS 2 software. For neuronal cells, equal protein levels were checked by the extraction of proteins from cellular debris with 1N HCl for 30 min at room temperature, followed by precipitation with Coomasie blue protein assay (Pierce, Bezons, France).

Virus infection, immunoprecipitation, and quantitative Western blotting. Recombinant Semliki Forest virus (SEV) encoding human APP695 was prepared and applied as described previously (Tienari et al., 1996). After 8 hr of exposure to 50 μg/ml LDL and U18666A or ethanol as a solvent control (final concentration, <0.1%), conditioned media were collected and total cellular extracts were prepared with lysis buffer consisting of 1% NP-40, 1% Triton X-100, 0.2% SDS, and 5 mm EDTA supplemented with Complete protease inhibitor cocktail (Roche, Mannheim, Germany). SH-SY5Y cells overexpressing SP-C99 (Dyrks et al., 1993) were incubated with LDL and 3 μg/ml U18666A for 24 hr and processed for immunoprecipitation by a similar protocol. Samples were immunoprecipitated with monoclonal antibodies W02 (1 μg/ml), G2-10 (2 μg/ml), or G2-11 (4 μg/ml). For the determination of Aβ species, Western blot detection of immunoprecipitates was performed according to Ida et al. (1996). Standard Aβ solutions at different concentrations were used for calibration. Western blotting from neuronal lysates was performed with anti-PS1 antibody 95.23 (diluted 1:10,000 in PBS) or monoclonal anti-calnexin antibody (diluted 1:1000 in PBS).

RESULTS

Inhibition of cholesterol trafficking in neuronal cells expressing full-length APP decreases proteolytic processing by β-secretase.

To elucidate the cellular mechanisms that connect cholesterol and Aβ generation, we investigated how alterations in intracellular cholesterol distribution might affect APP processing. In peripheral cells the transport of LDL-derived cholesterol to the ER can be inhibited by class-2 amphiphiles (Lange et al., 2000). Because cholesterol homeostasis in brain differs from peripheral tissue (Lütjohann et al., 1996), we first analyzed the transport of internalized cholesterol in neuronal cells by measuring its esterification by ER resident ACAT (Pentchev et al., 1987). SH-SY5Y neuroblastoma cells or fully polarized rat cortical neurons were incubated with LDL enriched in 14C-cholesterol in the presence or absence of U18666A or imipramine, and 14C-cholesterol ester formation was quantified. SH-SY5Y cells showed a reduction in cholesterol esterification by >80% with U18666A treatment and by >40% with exposure to imipramine at all time points that were investigated (Fig. 1A). Cholesterol ester levels in primary neurons decreased by ~60% (39.1% of control; n = 3; p < 0.05) (Fig. 1B) with exposure to U18666A. In contrast, cellular 14C-
cholesterol load was not changed significantly, showing that the diminished esterification did not result from a disruption of cholesterol internalization. These results clearly show that cholesterol ester generation from internalized cholesterol takes place in cultured neurons and that these cells, as known from other cell types, show a marked inhibition of cholesterol transfer to metabolizing compartments in response to amphiphiles.

We then investigated the effect of amphiphile-induced inhibition of cholesterol transport on APP processing in neuronal cells. Primary cultures of mixed cortical neurons were infected with recombinant SFV encoding for full-length human APP695. This expression system has been used widely to study sorting and processing of APP in neuronal cells and did not affect neuronal viability within the duration of our experiments (Tienari et al., 1996; De Strooper et al., 1998; Skovronsky et al., 1998). Infected neurons were exposed for 8 hr to LDL at different concentrations of U18666A. Conditioned medium was immunoprecipitated with W02, followed by Western blot detection with W02. Immunoprecipitations of conditioned media from neurons exposed to LDL and either 0.75 \( \mu \)g/ml U18666A (+) or not (−) with antibodies G2-10 or G2-11 were detected with G2-10 to visualize secretory Aβ40 and p3 or with W02 for the detection of secretory Aβ42, respectively. D, Western blot of respective neuronal lysates with W02 (after W02 immunoprecipitation), anti-calnexin antibody, or antibody 95.23 against PS1 NTF. C, D, Quantifications of intracellular (ic) and secreted (sec) APP, βCTF, and overall Aβ levels \( (n = 4–7\) experiments). Depicted are ratios of signal intensities from U18666A- or imipramine-treated versus untreated primary cortical neurons or SH-SY5Y cells. Error bars indicate 1 SD.

Figure 1. \( ^2 \) Class 2 amphiphiles inhibit intracellular cholesterol transport and decrease β-cleavage of APP in neuronal cells. SH-SY5Y cells (A) or mature rat cortical neurons in culture (B) were labeled with \(^{14}\)C-cholesterol in LDL for 2–12 hr (A) or 24 hr (B). \(^{14}\)C-cholesterol ester formation in cells incubated in the presence or absence of 3 \( \mu \)g/ml (for neuroblastoma cells) or 0.75 \( \mu \)g/ml (for primary neurons) U18666A or 80 \( \mu \)M imipramine was monitored by the extraction of lipids and subsequent TLC. Depicted are densitometric quantifications of \(^{14}\)C-cholesterol ester signal intensities as a percentage of the total \(^{14}\)C activity per lane (means from three experiments). CE, Cholesterol ester; Chol, cholesterol. C, SFV-infected mature rat cortical neurons were incubated for 8 hr with LDL at different concentrations of U18666A. Conditioned medium was immunoprecipitated with W02, followed by Western blot detection with W02. Immunoprecipitations of conditioned media from neurons exposed to LDL and either 0.75 \( \mu \)g/ml U18666A (+) or not (−) with antibodies G2-10 or G2-11 were detected with G2-10 to visualize secretory Aβ40 and p3 or with W02 for the detection of secretory Aβ42, respectively. D, Western blot of respective neuronal lysates with W02 (after W02 immunoprecipitation), anti-calnexin antibody, or antibody 95.23 against PS1 NTF. C, D, Quantifications of intracellular (ic) and secreted (sec) APP, βCTF, and overall Aβ levels \( (n = 4–7\) experiments). Depicted are ratios of signal intensities from U18666A- or imipramine-treated versus untreated primary cortical neurons or SH-SY5Y cells. Error bars indicate 1 SD.
A diminished secretion was prominent for both Aβ/H9252 (0.661 ± 0.053 for APPsec) as well as exposure of primary neurons to imipramine for Aβ/H11005 (0.524 ± 0.085 for APPsec) as well as exposure of primary neurons to imipramine (0.661 ± 0.124, n = 7, p < 0.001 for Aβsec; 0.98 ± 0.139 for APPsec). A diminished secretion was prominent for both Aβ40 and Aββ42, indicating that none of the major γ-secretase activities was perturbed in favor of the other. Reduced secretion of Aβ could not be explained by an increase in the α-secretory cleavage pathway of APP (Kojro et al., 2001), because we did not observe a compensatory upregulation of p3 levels.

Regardless of the treatment, levels of intracellular APP, calnexin, or PS1 fragments in neuronal lysates were unperturbed (Fig. 1D). However, we found remarkably less βCTF with exposure to U18666A. Quantification revealed a decrease in βCTF levels to an extent almost similar to intracellular Aβ levels in rat primary neurons [0.524 ± 0.203, n = 4, p < 0.05 for βCTF; 0.486 ± 0.195 for intracellular Aβ (Aβic); 0.96 ± 0.244 for intracellular APP (APPsec)] and human SH-SY5Y cells exposed to U18666A (0.723 ± 0.163, n = 3, p < 0.05 for βCTF; 0.64 ± 0.304 for Aβic; 1.052 ± 0.053 for APPsec). This effect was not specific for U18666A but also could be found in primary neurons exposed to imipramine (0.848 ± 0.1, n = 7, p < 0.01 for βCTF; 0.755 ± 0.169 for Aβic; 0.991 ± 0.148 for APPsec). Taken together, these results show that Aβ secretion and generation are reduced in neuronal cells inhibited in intracellular cholesterol transport. The similar decrease in Aβ and βCTF levels indicates that this effect is attributable to a reduction in β- rather than γ-cleavage of full-length APP.

**Inhibition of cholesterol transport increases γ-secretase cleavage in neuronal cells expressing APP C-terminal fragment SP-C99**

We now were interested in the specification of cholesterol transport increases γ-secretase cleavage in neuronal cells expressing APP C-terminal fragment SP-C99.

We stably transfected SH-SY5Y cells with SP-C99 and exposed these cells to LDL-enriched culture medium in the presence or absence of 3 μg/ml U18666A. Western blot detection showed that levels of endogenously synthesized secretory human APP (APPsec) in conditioned medium from U18666A-treated cells primarily were unaffected compared with controls. Surprisingly, however, secretory and intracellular Aβ levels were elevated markedly (1.56 ± 0.221, n = 5, p < 0.01 for Aβsec; 1.715 ± 0.172, n = 5, p < 0.0005 for Aβic) (Fig. 2A,B). In contrast, no major changes were found for secretory or intracellular endogenous APP (1.155 ± 0.134 for APPsec; 0.969 ± 0.074 for APPic). Similarly, treatment did not induce differences in C99 levels, the substrate for γ-secretase after cotranslational cleavage of the signal peptide from SP-C99 (1.059 ± 0.118 for C99). These results indicate a highly significant increase of Aβ in neuroblastoma cells exposed to U18666A, indicating a strong upregulation of γ-secretase activity in response to an altered subcellular cholesterol distribution.

Because of this strong enlargement of overall intracellular and secreted Aβ pools, we were interested in which of the major Aβ species might contribute to this effect. We found a remarkable increase in both Aβ40 and Aβ42 in cells exposed to U18666A compared with controls (Fig. 2C). This increase was prominent for secretory as well as for intracellular forms of both Aβ species, whereas no significant change could be found in the Aβ40/Aβ42 ratio.
Inhibition of cholesterol trafficking leads to an accumulation of presenilin1 in vesicular organelles associated with late endosomal/lysosomal compartments

Given this strong increase in the γ-secretase product Aβ with the inhibition of intracellular cholesterol transport, we were interested in whether this effect might correlate with changes in proteins involved in γ-secretase cleavage. In view of the role for PS in APP processing, we speculated that a stimulation of γ-cleavage might correlate with an altered subcellular distribution of PS1 (De Strooper and Annaert, 2001). PS1 has been localized to a number of different subcellular compartments, depending on cell type and methods used for its detection. In neuronal cells the majority of PS1 was proposed to reside in the ER and the ER–Golgi intermediate compartment (Annaert et al., 1999). To investigate whether PS1 is affected by alterations in intracellular cholesterol distribution, we exposed SH-SY5Y neuroblastoma cells to U18666A, and we determined PS1 distribution by indirect immunofluorescence and confocal microscopy.

As expected for neuroblastoma cells (Kovacs et al., 1996; Culvenor et al., 1997) and regardless of the presence of LDL, PS1 in untreated and LDL-exposed cells showed a concentration in perinuclear and membranous organelles, although minor signals also were found at the plasma membrane and in vesicular compartments (Fig. 3A). Human vacuolar ATPase as a marker for late endosomes and lysosomes was spread in small vesicular structures throughout the cytosol (Fig. 3B), showing no major colocalization with filipin (Fig. 3C,D), which allows for visualization of cellular cholesterol (Neufeld et al., 1999). However, when cholesterol transport was inhibited by U18666A, v-ATPase-positive vesicles were enlarged and now retained the major pool of cellular cholesterol (Fig. 3F,G). Surprisingly, under these conditions a large fraction of PS1 signal could be seen in vesicular structures that showed a remarkable overlap with enlarged v-ATPase and filipin-positive vesicles (Fig. 3E–H). This vesicular staining pattern for PS1 in U18666A-treated SH-SY5Y cells was independent of the antibody that was used. Results were identical with a polyclonal antibody against PS2 showed that under these conditions PS2 also localized to these compartments (Fig. 4G–I). However, these compartments were not identical with lipid storage droplets (Pol et al., 2001) visualized with the neutral lipid stain Oil red O (Fig. 4J–L). Instead, with a localization near to the nucleus and a ring-like morphology surrounding a central cavity reported previously (Neufeld et al., 1999; Zhang et al., 2001), our findings argue strongly for an accumulation of PS in vesicular compartments involved in intracellular cholesterol distribution.

PS1 and Aβ42 colocalize in Rab7-positive vesicular compartments in primary hippocampal neurons

We next analyzed the effects of U18666A on primary hippocampal neurons in culture. Colabeling of the late endosomal marker LBPA and filipin at higher magnification showed a notable overlap (Fig. 5C,E), confirming LBPA-positive late endosomes as cholesterol-retaining organelles after treatment of the cells with amphiphiles (Zhang et al., 2001). In contrast, Rab7-positive organelles were associated spatially with, but clearly distinct from, late endosomes enriched in cholesterol (Fig. 5F,H). Aggregated Aβ had been described previously in Rab7-positive endosomal fractions in U18666A-treated peripheral cells (Yamazaki et al., 2001). Thus we tested whether PS1 and Aβ colocalized to the same vesicular compartments in hippocampal neurons. As shown in Figure 5G and like in peripheral cells, PS1 in U18666A-treated hippocampal neurons distributed to vesicular compartments most likely corresponding to Rab7-positive organelles. Under these conditions in a subset of neurons the predominant fraction of endogenous Aβ42 localized to PS1-containing vesicular compartments (Fig. 5G–I), indicating a close relationship of cholesterol retention, PS1 distribution, and increased γ-secretase activity.

To determine the specificity of PS1 accumulation with the exposure of neurons to U18666A, we performed coimmunolocalization with markers for different subcellular compartments. β′-COP as a component of COP-I coats and parental Golgi membranes was largely excluded from PS1 or cholesterol-containing organelles (Fig. 6A–D). Similarly, no spatial correlation to filipin-positive compartments could be found for ER resident BiP/GRP78 (Fig. 6E–H), suggesting that an accumulation in vesicular compartments is not a general response of ER proteins after preventing delivery of cholesterol to the ER. Surprisingly, however, when we investigated the localization of ER protein calnexin in response to membran lipid composition by U18666A and suggest a correlation of cholesterol retention in late endosomal/lysosomal compartments and PS1 compartmentalization.

A more detailed analysis of U18666A-induced PS1-containing structures in SH-SY5Y cells revealed that PS1-containing compartments did not overlap completely with v-ATPase and presumably cholesterol-filled compartments but frequently appeared to surround these in a ring-like manner (Fig. 4A–C). Organelles positive for NPC1 and Rab7 with a very similar morphology and involved in cholesterol sorting have been described in fibroblasts enriched in LDL-derived cholesterol (Neufeld et al., 1999; Zhang et al., 2001). We therefore made use of a human A431 cell line stably expressing Rab7-GFP (M. de Bernard, unpublished data) and performed coimmunolabeling with a monoclonal antibody directed to the cytosolic loop region of PS1 CTF. As shown in Figure 4D–F, U18666A-treated cells showed a prominent colocalization of Rab7 and PS1 in vesicular structures that were concentrated in perinuclear areas. Colocalization with polyclonal antibody against PS2 showed that under these conditions PS2 also localized to these compartments (Fig. 4G–I). However, these compartments were not identical with lipid storage droplets (Pol et al., 2001) visualized with the neutral lipid stain Oil red O (Fig. 4J–L). Instead, with a localization near to the nucleus and a ring-like morphology surrounding a central cavity reported previously (Neufeld et al., 1999; Zhang et al., 2001), our findings argue strongly for an accumulation of PS in vesicular compartments involved in intracellular cholesterol distribution.
U18666A (Fig. 6J), a fraction of calnexin became prominent in punctuate structures associated with PS1-positive and cholesterol-retaining compartments (Fig. 4D–F). Higher magnification suggested that calnexin and PS1 partially colocalized to identical vesicular organelles.

Finally, we tested whether the effects of U18666A on cholesterol transport were attributable to potential direct effects on PS. Fibroblasts from PS1 knock-out and PS1/PS2 double knock-out mice (Armogida et al., 2001) were exposed to U18666A, and CE generation in treated and untreated cells was determined. As shown in Figure 7, CE formation from internalized cholesterol can occur regardless of PS1 expression. Furthermore, U18666A decreased CE generation to a similar degree in wild-type and knock-out cells. Correspondingly, retention of cholesterol in late endosomes with the blocking of cholesterol transport by U18666A was unchanged (data not shown). Together, these results suggest that, although PS1 expression is not essential for an efficient transfer of internalized cholesterol to metabolizing sites, PS1...
localization in hippocampal neurons correlates highly with the intracellular distribution of cholesterol.

**DISCUSSION**

This study reports that the intracellular distribution of cholesterol has substantial effects on the subcellular localization of PS and the amyloidogenic processing of APP. Our results suggest that the compartmentalization of cholesterol in neurons may play an important role in how cholesterol alters Aβ generation. We determined the effects of intracellular cholesterol distribution on PS localization and APP cleavage by two means, first by using compounds that inhibit cholesterol transport from endocytic compartments to the ER and second by examining cells expressing mutant NPC1.

We describe a dose-dependent reduction in both secreted and cellular Aβ species in neurons and neuroblastoma cells exposed to cholesterol transport-inhibiting amphiphiles. For several rea-
subcellular cholesterol distribution could alter the cholesterol metabolism in neuronal cells (Simons et al., 1998). Mal compartments might interfere directly with (Lusa et al., 2001), retention of cholesterol in endosomal/lysosomal rafts (Riddell et al., 2001) and NPC1 plays a role in raft dynamics (Subtil et al., 1999). However, intriguing possibilities argue for a more direct role of the intracellular cholesterol transport on Aβ generation. We cannot rule out that the effects on APP processing caused by alterations in endosomal or TGN function (Pentchev et al., 1987) or consecutive regulatory mechanisms after decreased transport of sterols to the ER (Subtil et al., 1999) have an impact on APP. Whereas cholesterol-enriched late endosomes are positive for Rab7 (D. F. green), G, H, Coimmuno- n stainings of 95.23 for PS1 and monoclonal antibody G2-11 to detect cellular Aβ42. A–C and insets in D–I show areas in the cell body of a hippocampal neuron. C, Merged image from A, B, F, Merged image from D, E, I, Merged image from G, H. Arrows indicate selected compartments. Scale bars, 4 μm.

Figure 5. Rab7-containing compartments after U18666A exposure of hippocampal neurons are distinct from cholesterol-retaining late endosomes and retain Aβ42 and PS1. Mature hippocampal neurons were incubated with LDL and U18666A for 24 hr, fixed, and stained with filipin (B, E; red) and either monoclonal antibody 6C4 against LBPA (A, C; green) or a polyclonal antibody against Rab7 (D, F; green). G, H, Coimmuno- staining of 95.23 for PS1 and monoclonal antibody G2-11 to detect cellular Aβ42. A–C and insets in D–I show areas in the cell body of a hippocampal neuron. C, Merged image from A, B, F, Merged image from D, E, I, Merged image from G, H. Arrows indicate selected compartments. Scale bars, 4 μm.

There are several possible mechanisms for how changes in subcellular cholesterol distribution could alter the β-secretase cleavage of APP. First, βCTF levels decrease to an almost similar extent as overall secreted and intracellular Aβ, and Aβ40 and Aβ42 both decrease to a similar degree. Second, levels in APPsec and p3 do not rise, indicating that α-secretase cleavage of APP was not increased. Our results confirm previous data showing a strong correlation of β-secretase activity and cholesterol metabolism in neuronal cells (Simons et al., 1998).

There are several possible mechanisms for how changes in subcellular cholesterol distribution could alter the β-secretase cleavage of APP. We cannot exclude indirect effects on APP processing caused by alterations in endosomal or TGN function (Pentchev et al., 1987) or consecutive regulatory mechanisms after decreased transport of sterols to the ER (DeBose-Boyd et al., 1999). However, intriguing possibilities argue for a more direct role of the intracellular cholesterol transport on Aβ production. For example, reinternalization of surface APP during Aβ generation may be affected by a decreased cholesterol content in the plasma membrane in response to cholesterol-depleting agents that also reduce cholesterol transport to the ER (Subtil et al., 1999; H. Runz and T. Hartmann, unpublished results). Because the β-secretase BACE1 has been shown to localize to lipid rafts (Riddell et al., 2001) and NPC1 plays a role in raft dynamics (Lusa et al., 2001), retention of cholesterol in endosomal/lysosomal compartments might interfere directly with β-secretase function. Our findings of a reduced β-cleavage of APP after the inhibition of cholesterol transport to the ER also correspond to a recent study showing a strong correlation of Aβ production and cellular CE levels (Puglielli et al., 2001). Inhibition of ACAT was associated with a prominent reduction in Aβ generation, which resulted from an inhibition of all three major processing events of APP.

Unexpectedly and in contrast to what we observed for β-secretase cleavage, we found a dramatic increase in γ-secretase activity accompanied by higher levels of secreted and intracellular Aβ when the transfer of LDL-derived cholesterol from endocytic compartments to ACAT in the ER was inhibited. This increase was prominent in cells expressing the βCTF analog SP-C99 but was concealed in cells expressing full-length APP, where an increased γ-cleavage could not compensate for a reduction in β-cleavage as the rate-limiting step in Aβ generation. We cannot rule out that the effects on γ-secretase are attributable to downstream effects on ACAT itself or mechanisms important for CE formation. However, a direct influence of NPC1 function or the transport of free cholesterol or other lipids to the ER on APP cleavage cannot be excluded.

This is supported further by the fact that regardless of how cholesterol transport was inhibited, we found a prominent accumulation of PS in vesicular organelles positive for Rab7 and intimately associated, but not identical, with cholesterol-retaining late endosomes. During cholesterol transport from the plasma membrane to the ER vesicular compartments are involved that contain NPC1 protein and are associated spatially but that are distinct from endosomal/lysosomal compartments required for the initial uptake of lipoprotein-derived cholesterol (Neufeld et al., 1999). Whereas cholesterol-enriched late endosomes are pos-
itive for lysosomal hydrolases, LBPA, and endocytic fluorescent-labeled dyes, NPC1-containing structures exclude these markers but are enriched instead in Rab7 and certain glycolipids (Zhang et al., 2001). The NPC1 compartment was described as a membranous organelle often surrounding cholesterol-laden late endosomes in a ring-like manner that may have a role in cholesterol sorting and the rapid transport of material between the plasma membrane and the ER (Cruz et al., 2000; Ko et al., 2001). The colocalization of PS with Rab7 as well as an identical morphology of PS-containing structures in our experiments indicate that, on blocking intracellular cholesterol transport, PS accumulates in these vesicular organelles involved in cholesterol sorting.

The localization of PS in vesicular organelles was surprising, because convincing evidence from neuronal cells shows a steady-state localization of PS1 predominantly in the ER (Kovacs et al., 1996; Culvenor et al., 1997; Annaert et al., 1999). Internalization of lipoprotein-derived cholesterol by neurons was reported to be low (Lütjohann et al., 1996). Nevertheless, neurons express LDL receptor family members (Herz et al., 1988), NPC1 (Falk et al., 1999) and ACAT (Sakashita et al., 2000) and thus possess molecular equipment involved in the uptake and utilization of lipoprotein-derived cholesterol. Accordingly, in our experiments amphiphiles diminished esterification of LDL-derived cholesterol in SH-SY5Y cells and primary cortical neurons. The block in cholesterol transfer to ACAT-containing compartments was sufficient to induce a massive accumulation of cholesterol in late endosomes positive for LBPA but distinct from Rab7-containing organelles. These results indicate that the intracellular transport

![Figure 6.](image)

**Figure 6.** PS1-bearing vesicular compartments in hippocampal neurons impaired in cholesterol transport exclude β’COP and BiP but retain calnexin. Fully polarized rat hippocampal neurons in culture were incubated with 50 µg/ml LDL and 0.75 µg/ml U18666A for 24 hr. Neurons were fixed and triple labeled with antibody 95.23 (A, E, I), filipin (C, G, K), and monoclonal antibodies against β’COP (B), BiP/GRP78 (F), or calnexin (J). D, H, L, Merged images. Bottom insets show selected regions of single sections in the cell body of the depicted neuron (top insets). Arrows indicate selected compartments. Scale bars, 3 µm.

![Figure 7.](image)

**Figure 7.** U18666A decreases cholesterol ester formation independently of PS1 expression. Cultured mouse fibroblasts from wild-type (PS1+/+PS2+/+), PS1 single knock-out (PS1−/−PS2+/+), and PS1/PS2 double knock-out (PS1−/−PS2−/−) mice were labeled with 14C-cholesterol in LDL in the presence (+) or absence (−) of 3 µg/ml U18666A for 24 hr. Lipids were extracted and separated by TLC. Effects of U18666A on CE formation in treated cells at equal CE levels in untreated cells are shown.
of internalized cholesterol in primary neurons essentially follows identical routes as in peripheral cells (Zhang et al., 2001). Correspondingly, PS1 localized to vesicular compartments in the cell bodies of hippocampal neurons. Most interestingly, we also could detect a large fraction of Aβ42 in these compartments. These microscopic data from neuronal cells inhibited in intracellular cholesterol transport are in accordance with a study showing a significant accumulation of Aβ in both NPC1 mutant CHO cells and NPC mouse brain (Yamazaki et al., 2001). In addition to their results proposing a pool of aggregated Aβ42 in cholesterol- and Rab7-containing gradient fractions, our data showing a colocalization of PS1 and Aβ42 raise the possibility that under these conditions Aβ42 is produced in vesicular compartments involved in neuronal cholesterol transport.

How might an inhibition of intracellular cholesterol transport induce a shift in PS distribution and an increased γ-secretase activity? One possibility is that alterations in subcellular cholesterol distribution might induce changes in ER membrane composition that could alter ER stability (Orci et al., 1984) and favor a relocation of ER proteins to compartments involved in cholesterol sorting. This might be a reason for the unexpected localization of a fraction of ER membrane protein calnexin to these organelles in hippocampal neurons. The accumulation of PS and presumably other factors in Rab7- and NPC1-bearing organelles also might favor the assembly of the active high-molecular-weight γ-secretase complex, which might stimulate formation of Aβ in these compartments. Because the site for γ-cleavage of BCTF is within the membrane domain of APP, a unique lipid composition of PS- and NPC1-containing compartments might favor Aβ generation further.

Alternatively, the redistribution of PS to organelles involved in cholesterol trafficking might support a proposed ER export model for PS-dependent γ-cleavage of APP (De Strooper and Naeraa, 2001). Analogies between APP processing and sterol-dependent trafficking and proteolysis in the regulation of sterol homeostasis are intriguing. When ER membranes are depleted from cholesterol, the sterol response element binding protein (SREBP) cleavage-activating protein (SCAP) escorts SREBP to the Golgi, where Site-1 protease activates SREBP, which subsequently is cleaved by an intramembranous proteolytic event that is very similar to the γ-cleavage of APP (DeBose-Boyd et al., 1999). Recent data from living cells demonstrate a close interaction of NPC1-bearing organelles and the ER (Ko et al., 2001), and a threshold concentration of cholesterol in the endosomal/lysosomal system was proposed to induce a shift in NPC1 localization from a diffuse membranous organelle to these compartments (Zhang et al., 2001).

In conclusion, our results show that the intracellular compartmentalization of cholesterol is an important factor in determining PS localization and the regulation of APP cleavage. Further experiments in additional cellular and in vivo models systems will be required to elucidate its potential physiological and pathological implications on APP processing. In view of recent clinical studies reporting a potential benefit of regulating cholesterol metabolism as a strategy for preventing the onset of AD symptoms (Jick et al., 2000; Wolozin et al., 2000), our findings may help to design therapeutic approaches to cholesterol-dependent cellular events associated with AD. In providing a plausible link between factors such as ApoE or LRP in the endocytic limb and subsequent cholesterol-regulating events in the ER where, under balanced conditions, the major pool of PS is localized and γ-cleavage may occur, we propose the intracellular cholesterol transport as a potential crossroad in Aβ generation.

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