Site-1 protease-activated formation of lysosomal targeting motifs is independent of the lipogenic transcription control

Sarah Klünder†, Jörg Heeren§, Sandra Markmann†, René Santer†, Thomas Braulke†, and Sandra Pohl† 1
†Section Biochemistry, Children’s Hospital, §Department of Biochemistry and Molecular Cell Biology, University Medical Center Hamburg-Eppendorf, Martinistrasse 52, 20246 Hamburg, Germany

1To whom correspondence should be addressed: Dr. Sandra Pohl, Section Biochemistry, Children’s Hospital, University Medical Center Hamburg-Eppendorf, Martinistrasse 52, 20246 Hamburg, Germany, Phone: 0049 40 7410 58780, Fax: 0049 40 7410 58504, Email: s.pohl@uke.de

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Abbreviations:
Actb, β-actin; ASB, arylsulfatase B; ATV, atorvastatin; Baf A1, bafilomycin A1; ER, endoplasmic reticulum; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; GlcNAc, N-acetylglucosamine; GM130, Golgi marker protein 130; GNPTAB, α/β-subunit precursor of the GlcNAc-1-phosphotransferase gene; HRP, horseradish peroxidase; LPDS, Lipoprotein-deficient serum; LDLR, LDL receptor; M6P, mannose 6-phosphate; MEF, mouse embryonic fibroblasts; MLII, mucolipidosis type II; MPR, M6P receptor; NBCS, newborn calf serum; NPC1, Niemann-Pick disease type C1; PDI, protein-disulfide isomerase; PCSK9, proprotein convertase subtilisin/kexin type 9; S1P, site-1 protease; S2P, site-2 protease; SCAP, SREBP cleavage-activating protein; SREBP, sterol regulatory element-binding protein

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ABSTRACT

Site-1 protease (S1P) cleaves membrane-bound lipogenic sterol regulatory element-binding proteins (SREBPs) and the α/β-subunit precursor protein of the GlcNAc-1-phosphotransferase forming mannose 6-phosphate (M6P) targeting markers on lysosomal enzymes. The translocation of SREBPs from the endoplasmic reticulum (ER) to the Golgi-resident S1P depends on the intracellular sterol content, but it is unknown whether the ER exit of the α/β-subunit precursor is regulated. Here, we have investigated the effect of cholesterol depletion (atorvastatin treatment) and elevation (LDL overload) on ER-Golgi transport, S1P-mediated cleavage of the α/β-subunit precursor, and the subsequent targeting of lysosomal enzymes along the biosynthetic and endocytic pathway to lysosomes. The data showed that the proteolytic cleavage of the α/β-subunit precursor into mature and enzymatically active subunits does not depend on the cholesterol content. In either treatment, lysosomal enzymes are normally decorated with M6P residues, allowing the proper sorting to lysosomes. In addition we found that in fibroblasts of mucolipidosis type II mice and Niemann-Pick type C patients characterized by aberrant cholesterol accumulation, the proteolytic cleavage of the α/β-subunit precursor was not impaired. We conclude that S1P substrate-dependent regulatory mechanisms for lipid synthesis and biogenesis of lysosomes are different.

Supplementary key words: mannose 6-phosphate, LDL, LDL receptor, cholesterol, statins, Niemann-Pick disease, Golgi apparatus, endocytosis, mucolipidosis type II, lysosome
Lysosomes function in degradation of macromolecules, such as proteins, lipids, glycosaminoglycans or nucleic acids, intracellular organelles and pathogens obtained by endocytosis, autophagy and phagocytosis through the concerted action of more than 50 acid hydrolases (1). To maintain their function, lysosomes require a continuous replenishment of newly synthesized components transported from the endoplasmic reticulum (ER) via the Golgi apparatus to the endosomal/lysosomal compartment. The efficient transport of soluble lysosomal enzymes to lysosomes requires mannose 6-phosphate (M6P) residues on their N-linked glycans which serve as recognition markers for M6P-specific receptors (MPR) (2). The formation of M6P residues is catalyzed by two enzymes, the N-acetylglucosamine-1-phosphotransferase (termed GlcNAc-1-phosphotransferase) and the GlcNAc-1-phosphodiester α-N-acetylglucosaminidase (termed uncovering enzyme) localized in the cis-Golgi apparatus and trans-Golgi network (TGN), respectively (3).

The GlcNAc-1-phosphotransferase is a heterohexameric complex of three subunits (α2β2γ2) which are encoded by two genes (4-6). The α- and β-subunits of the GlcNAc-1-phosphotransferase are synthesized in the ER as a single highly N-glycosylated 190-kDa type III membrane protein (6). For the transport to the Golgi apparatus combinatorial dileucine and dibasic sorting motifs in the N- and C-terminal cytosolic domains, respectively, are required (7). Upon arrival in the cis-Golgi compartment, the α/β-subunit precursor is proteolytically cleaved by the site-1 protease (S1P) into mature α- and β-subunits (8) prerequisite for the enzymatic activity of the GlcNAc-1-phosphotransferase complex (9).

S1P (also known as subtilisin kexin isoenzyme-1, SKI-1) is a type I membrane serine protease and plays a crucial role in the proteolytic activation of the sterol regulatory element (SRE)-binding proteins, SREBP-1 and -2, necessary for synthesis of fatty acids, triglycerides, and cholesterol, respectively (10, 11). SREBPs are type III membrane-bound transcription factors with both cytosolic N-terminal transcription factor domains and C-terminal regulatory domains (12). At conditions of cholesterol abundance SREBPs form ER-anchored complexes with the polytopic sterol-sensing SREBP cleavage-activating protein (SCAP) and INSIG, an insulin-induced gene product. Upon cholesterol depletion, the complex dissociates and SCAP escorts SREBPs from the ER to the Golgi apparatus followed by S1P-initiated release of the active transcription factor domain entering the nucleus (13, 14).
In cells deficient for S1P, the α/β-subunit precursor of GlcNAc-1-phosphotransferase cannot be cleaved and activated, which is subsequently associated with the lack of M6P residues, missorting of newly synthesized lysosomal enzymes, and accumulation of non-degraded storage material in lysosomes (8), hallmarks of mucolipidosis type II (MLII). MLII (also called I-cell disease) is a severe multi-systemic inherited disorder of childhood caused by mutations in the GNPTAB gene encoding the α/β-subunit precursor protein of the GlcNAc-1-phosphotransferase (6, 15). Although the nature of storage material in MLII cells and tissues has not been fully characterized, analysis of fibroblasts of MLII patients and MLII mice revealed in addition to increased level of phospholipids, fucosylated oligosaccharides, and sialic acid-containing glycosphingolipids, an elevation and lysosomal accumulation of cholesterol (16-18).

In the present study, we investigated whether alterations in the cholesterol content of murine and human fibroblasts affects the ER-Golgi transport, the proteolytic activation of the α/β-subunit precursor of GlcNAc-1-phosphotransferase, and the subsequent lysosomal targeting of lysosomal enzymes. The results were compared with data from fibroblasts of Niemann-Pick type C1 (NPC1) patients and MLII mice both characterized by abnormal cholesterol metabolism (18, 19). These analyses demonstrate that the export and the S1P-mediated cleavage of the α/β-subunit precursor protein of GlcNAc-1-phosphotransferase is independent of the cellular cholesterol content. Furthermore, missorting of lysosomal enzymes and lysosomal dysfunction in MLII fibroblasts lead to up-regulation of endocytic receptors such as LDL receptor (LDLR) and the 300-kDa MPR (MPR300), and accumulation of their degradation intermediates.
MATERIALS AND METHODS

Reagents
Sodium $^{125}$I (74 TBq/mmol) and $[^{35}S]$methionine (37 Bq/mmoll) were purchased from Hartmann Analytik. Atorvastatin, newborn calf serum (NBCS), saponine, paraformaldehyde, mannose 6-phosphate sodium salt, penicillin/streptomycin, cycloheximide, TRI™ Reagent, protease inhibitor cocktail (P2714), and BSA were obtained from Sigma-Aldrich. Maxima™ Probe qPCR Master Mix, Iodination reagent, and prestained protein molecular mass marker Pageruler™ were from Thermo Fisher Scientific. Prestained molecular mass marker Full-Range Rainbow™, FCS and methionine-free DMEM were from GE Healthcare. TaqMan® Gene Expression Assays, High Capacity cDNA Reverse Transcription Kit, DMEM, and Opti-MEM® were purchased from Life Technologies. Transfection reagent JetPEI® was purchased from Peqlab. Roti® Quant Protein Assay was from Roth. Lipoprotein-deficient serum (LPDS) and human LDL were prepared from human plasma as previously described (20). Recombinant human arylsulfatase B (ASB) was kindly provided by Dr. Vellard (BioMarin/Genzyme LLC, Novato, CA).

Antibodies
A myc-tagged single-chain antibody fragment against M6P residues (scFv M6P-1) and the monoclonal rat antibody against the α-subunit of the GlcNAc-1-phosphotransferase has been described recently (21,22). The monoclonal anti-myc antibody was purchased from Cell Signaling. The polyclonal antibody against cathepsin Z was obtained from R&D Systems. Monoclonal antibodies against glyceraldehydes 3-phosphate dehydrogenase (Gapdh), LDL receptor (LDLR) and the Golgi marker protein (GM130) were obtained from Santa Cruz Biotechnology, Epitomics-Abcam and BD Biosciences, respectively. The monoclonal antibody against β-tubulin was obtained from the Developmental Studies Hybridoma Bank (University of Iowa). Anti rat Mp r300 antibodies were a kind gift from Dr. von Figura (Göttingen, Germany). Secondary antibodies conjugated to horseradish peroxidase (HRP) or Alexa Fluor® were purchased from Dianova and Life Technologies, respectively.
Cell Culture and cDNA Transfection

Mouse embryonic fibroblasts (MEF) of wild-type (WT) and mucolipidosis type II (MLII) ‘knock-in’ mice were prepared as previously described (18). Human fibroblasts were obtained from skin biopsies with informed consent of two controls and three NPC1 patients carrying the following mutations encoded by the NPC1 gene: Patient 1 (P1) was heterozygous for p.G910S and p.G1034R, patient 2 (P2) and patient 3 (P3) were homozygous for p.S940L and p.I1095del, respectively. MEF and fibroblasts from healthy controls and NPC1-defective patients were cultured in DMEM containing 10% FCS and penicillin/streptomycin. To change the cholesterol content, MEF were grown on 6- or 3.5-cm plates or glass coverslips, rinsed with PBS (pH 7.4) and further incubated in DMEM containing 10% NBCS or LPDS. To deplete or overload the cholesterol content in cells, 10 µM atorvastatin or 100 µg/ml human LDL was added to the DMEM/LPDS medium for 48 h, respectively. Twenty four h after addition of atorvastatin (ATV) or LDL, MEF and human fibroblasts were transfected with cDNA of the α*/β-subunit mini construct #3 (8, 22) using JetPEI® according to the manufacturer’s instructions. The α*/β-subunit mini construct misses amino acids 431-819 and results in a shortened α*-subunit exhibiting identical topology, N- and C-terminal ER exit structures, and structural requirements for efficient proteolytic cleavage by S1P (8, 22). The medium was replaced 6 h after transfection, and cells were further incubated with cholesterol depletion medium or LDL loading medium for 18 h.

mRNA Analysis

Total RNA isolation, cDNA synthesis and real-time PCR were performed as previously described (8). TaqMan™ Gene Expression Assays including pre-designed probes and primer sets for mouse Ldlr (Mm00440169_m1), Gnptab (Mm01773334_m1), Actb (Mm00607939_s1) and human LDLR (Hs00181192_m1), ACTB (Hs9999903_m1) were used. The relative expression of human and murine LDLR, and Gnptab mRNA was normalized to the level of ACTB mRNA in the same cDNA using the comparative C_T method (2^-ΔΔCT).
Internalization Assay of $[^{125}\text{I}]\text{ASB}$

Recombinant human arylsulfatase B (ASB) was iodinated with Iodination Reagent and sodium $^{125}\text{I}$ to a specific activity of 6 µCi/µg as previously described (23). Cells grown on 3.5-cm plates were washed with PBS, pre-incubated with DMEM containing 0.1% BSA (DMEM/BSA) for 30 min at 37°C followed by incubation with $[^{125}\text{I}]\text{ASB}$ (625,000 cpm/ml DMEM/BSA) in the presence or absence of 10 mM M6P for 20 min at 37°C. After four washes with PBS to remove non-bound $[^{125}\text{I}]\text{ASB}$ cells were either harvested or chased in DMEM/BSA for 2 to 16 h at 37°C. Cells were lysed in 0.1 ml PBS containing 0.2% Triton X-100 and protease inhibitors for 30 min at 4°C, centrifuged at 15,000 x g at 4°C for 15 min and the supernatants were analyzed by SDS-PAGE and autoradiography.

Western Blotting

Cells were lysed for 30 min at 4°C in PBS containing 1% Triton X-100 and protease inhibitor cocktail. After centrifugation at 10,000 x g supernatants were used for measurement of the protein content by the Roti® Quant Protein Assay. Aliquots of cell extracts (75 µg protein) were solubilized, separated by SDS-PAGE and analyzed by western blotting either with monoclonal antibody against the α-subunit of the GlcNAc-1-phosphotransferase (dilution 1:25) (22), cathepsin Z (1:500), LDLR (1:1,000) or Mpr300 (1:500). β-Tubulin or Gapdh/GAPDH were used as loading controls. After incubation with secondary HRP-conjugated antibodies, the immunoreactive bands were visualized by enhanced chemoluminescence (Molecular Imager ChemiDoc XRS, Bio-Rad). The content of M6P-containing proteins in cell extracts was analyzed by scFv M6P-1 western blotting (21).

Other Methods

Confocal immunofluorescence microscopy of transfected cells, activity determination of the enzyme β-hexosaminidase and metabolic $[^{35}\text{S}]\text{methionine labeling of MEF}$ followed by immunoprecipitation of cathepsin Z were described recently (18, 22). Densitometric analyses were performed with the Image J software (http://rsbweb.nih.gov/ij/) to quantify band intensities in western blots and X-ray films.
Statistical analysis

For statistical analysis student’s t-test was performed. $P$-values $< 0.05$ were considered significant.
RESULTS

High LDL receptor expression in MLII mouse embryonic fibroblasts

To investigate, whether the α/β-subunit precursor protein of GlcNAc-1-phosphotransferase is transported from the ER to the Golgi apparatus independently of cholesterol/LDL or not, the effects of the cholesterol-depleting drug atorvastatin (ATV) and the LDL-mediated cholesterol overload were examined in mouse embryonic fibroblasts (MEF). The efficiency of both treatments has been demonstrated by determination of LDL receptor (Ldlr) expression known to be regulated by cellular cholesterol level (24). Treatment of WT MEF with 10 µM ATV for 48 h increased the mouse Ldlr mRNA expression 6-fold (Fig. 1). When cholesterol levels are high after LDL loading, the Ldlr mRNA expression dropped to 20% of controls. For comparison, the transcript concentration of Gnptab was not affected by changes in the cholesterol level in WT cells. These data indicate that both treatments to increase or decrease the cellular cholesterol concentration in MEF were effective and reproducible. In MEF of MLII mice harboring the mutant Gnptab<sup>c.3082insC</sup> (18) the Gnptab transcript level was reduced whereas the Ldlr mRNA concentration was 3-fold higher under basal conditions (DMEM supplemented by 10% NBCS) than in WT MEF which could be further increased by 50% after treatment with ATV (Fig. 1) suggesting low amounts of cholesterol in ER membranes under basal conditions of MLII fibroblasts. Incubation of MLII MEF with LDL reduced the Ldlr mRNA level to 20% of non-treated MLII cells.

The cholesterol-dependent alterations in the Ldlr mRNA expression in WT MEF could be confirmed on the LDLR protein level (Fig. 2). ATV treatment resulted in 12-fold increase whereas LDL overload suppressed the LDLR concentration to 30% of control cells (Fig. 2A). No or marginal effects on the LDLR expression has been observed when the cells were incubated in the presence of 10% NBCS or LPDS alone (supplementary Fig. 1). In MLII MEF the basal LDLR protein level was 3-fold higher than in wild-type MEF (Fig. 2A,B). Incubation of MLII MEF with ATV or LDL led to a 7-fold increase and 6-fold reduction of the LDL receptor expression, respectively.
Cholesterol-independent transport and proteolytic cleavage of the GlcNAc-1-phosphotransferase

The site-1 protease substrates SREBPs are translocated from the ER to the Golgi apparatus in dependency of the membrane cholesterol content (12). To examine whether the α/β-subunit precursor is transported to the Golgi apparatus in a cholesterol-dependent manner, MEF of WT mice (not shown), and human fibroblasts were treated with and without ATV, LDL, and transfected with the human α*/β-subunit precursor construct (8, 22) of the GlcNAc-1-phosphotransferase. Immunofluorescence microscopy revealed complete co-staining of α*-subunit immunopositive material with the cis-Golgi marker protein GM130 independent of high (LDL) or low (ATV) levels of cholesterol (Fig. 3A). When fibroblasts of a Niemann-Pick type C1 (NPC1) patient characterized by high level of non-esterified cholesterol in lysosomes (19) and 2- to 3.5-fold increased Ldlr mRNA level (supplementary Fig. 2), were analyzed, α*-subunit immunoreactive material was found in GM130-positive Golgi membranes (Fig. 3B). These data were complemented by western blot analysis of cell extracts of ATV or LDL treated WT MEF transiently transfected with the α*/β-subunit precursor construct. At steady state, small amounts of immunoreactive 120-kDa α*/β-subunit precursor polypeptides were observed in WT MEF (Fig. 3C) and human control fibroblasts (Fig. 3D). In contrast, strong signals of mature 90-kDa α*-subunits were detected both in MEF (Fig. 3C) and human fibroblasts (Fig. 3D). The proteolytic pattern of α*/β-subunit precursor and mature α*-subunit was neither affected by high nor low cholesterol content (Fig. 3C). Similarly, α*/β-subunit precursor expressed in NPC1 fibroblasts were proteolytically cleaved to mature α*-subunits (Fig. 3D). Identical cleavage patterns have been observed in two other NPC1 fibroblast cell lines (Fig. 3E). Of note, the endogenous α/β-subunit precursor or mature α-subunit cannot be visualized by the anti α-subunit monoclonal antibody due to their low expression. Finally, we tested whether MII MEF exhibiting high lysosomal cholesterol content (18) and characterized by low cholesterol sensing (Fig. 2A,B) were impaired in their capability to cleave the α*/β-subunit precursor protein. As shown in Fig. 3C, the majority of immunoreactive material is presented by the mature α*-subunit. These data demonstrate that neither
increase nor reduction of cellular cholesterol level affected the ER-Golgi transport and the proteolytic activation of the α*/β-subunit precursor protein of GlcNAc-1-phosphotransferase.

**Biogenesis of lysosomes is independent of cellular cholesterol content**

The enzymatic activity of the GlcNAc-1-phosphotransferase was determined indirectly by mannose 6-phosphate (M6P) western blotting showing that neither the reduction of cholesterol content by ATV nor the LDL-treatment affected the intensity or pattern of M6P-containing proteins in WT fibroblasts (Fig. 4A). MLII MEF were used as negative controls lacking M6P-containing proteins (18). To analyze the M6P-dependent targeting of lysosomal enzymes, we performed [35S]methionine pulse-chase experiments followed by immunoprecipitation of the lysosomal protease cathepsin Z (CtsZ) from cell extracts and media. In WT MEF the 38-kDa CtsZ precursor (p) was transported to lysosomes, proteolytically processed to the 36-kDa mature (m) form and partially secreted during the 4 h chase period (Fig. 4B). The synthesis rate of CtsZ, the proteolytic processing to the mature form and the amount of secreted precursor forms were not affected by ATV- or LDL-treatment of WT MEF (Fig. 4B). In contrast, in MLII MEF used as control, the newly synthesized CtsZ was completely missorted into the medium during the 4 h chase period and no CtsZ immunoreactive polypeptides were retained intracellularly (Fig. 4B). This was confirmed by the steady-state expression level of CtsZ. The amount of intracellular detectable 36-kDa mature CtsZ in ATV-, LDL- and non-treated WT MEF was similar (Fig. 4C) whereas no CtsZ immunoreactive material was observed in MLII MEF. Furthermore, the activity of another lysosomal enzyme, β-hexosaminidase, in ATV- or LDL-treated MEF and in their respective media was comparable with non-treated control cells (Fig. 4D). In contrast, the β-hexosaminidase activity in MLII MEF was markedly reduced by approximately 80% and missorted into the medium. These data indicate that neither ATV nor LDL treatment affected the GlcNAc-1-phosphotransferase activity in the Golgi apparatus and the transport and processing of lysosomal enzymes along the biosynthetic pathway to lysosomes.

To examine whether the transport of lysosomal enzymes along the endocytic pathway is altered in dependency of the cholesterol membrane content, WT MEF were incubated for 20 min with 125I-labeled
M6P-containing arylsulfatase B (ASB) followed by variable chase periods. ASB is internalized in an M6P-dependent manner as shown by the complete inhibition of ASB uptake in the presence of 10 mM M6P in the medium (Fig. 5). Subsequently the internalized ASB is proteolytically activated via 47-kDa intermediate to mature 15-kDa forms (25; Fig. 5A). The amount of internalized [125I]ASB was up to 2-fold higher in LDL-treated cells compared to non-treated and ATV-treated cells indicating that an elevated cellular cholesterol content resulted in increased [125I]ASB endocytosis whereas the subsequent proteolytic processing of [125I]ASB was not impaired by ATV nor LDL treatment. In MLII MEF, however, the amount of internalized [125I]ASB was 8-fold higher than in WT MEF and the proteolytic maturation was almost completely inhibited even 22 h after internalization of the ASB precursor (Fig. 5B). These data suggest that sufficient amounts of proteases involved in the maturation of the ASB precursor are present whereas proteases mediating the degradation of ASB polypeptides are missing in MLII lysosomes. Of note, in NPC1 cells (P1) both the amount of internalized [125I]ASB as well as the stability of proteolytically generated ASB forms were comparable with that in human control fibroblasts (Fig. 5C). The addition of the vATPase inhibitor bafilomycin A1 (Baf A1) during the chase periods completely inhibited the maturation and degradation of ASB polypeptides by acid proteases. These data indicate that the deficiency of various lysosomal enzymes, in particular lysosomal proteases, in MLII cells rather than the lysosomal accumulation of non-esterified cholesterol affect the proteolytic maturation of the internalized indicator protein ASB.

The increased M6P-dependent endocytosis of [125I]ASB correlated with 3-fold increase of Mpr300 expression in MLII MEF (Fig. 6A). The Mpr300 expression was neither affected by ATV-induced reduction of cholesterol content nor by LDL overload. In addition to the major 300-kDa Mpr immunoreactive protein, ~240- and 160-kDa Mpr300 immunoreactive polypeptide bands were observed in MLII MEF (Fig. 6A). Furthermore, the cholesterol-dependent variations in the 160-kDa LDLR expression (Fig. 2) correlated with the appearance of high amounts of 17-kDa and minor 40-kDa LDLR fragments and intermediates, respectively, in MLII MEF (Fig. 6B; 26). Small percentages of the 17-kDa LDLR fragment were detectable in ATV-treated WT MEF. These data support previous observations that both
Mpr300 and LDLR undergo ectodomain shedding at the plasma membrane or endosomes (26-28) followed by lysosomal degradation of the C-terminal fragments accumulating in high amounts in MLII MEF.
DISCUSSION

It is well known how the prototypical precursor membrane proteins of lipogenic transcription factors, SREBP-1 and SREBP-2, are released from the ER and proteolytically activated by S1P localized in the Golgi apparatus. In the presence of cholesterol the formation of a complex between SREBPs, the sterol-sensing protein SCAP and INSIG results in retention in the ER. When cells are depleted of sterols, the complex dissociates and exposes a Golgi sorting signal which allows the ER exit of SREBP/SCAP complexes, and subsequent first cleavage of SREBPs by S1P (29, 30). Here we examined whether the transport of the GlcNAc-1-phosphotransferase α/β-subunit precursor from the ER to the Golgi apparatus is also regulated by the intracellular sterol content, since the loss of GlcNAc-1-phosphotransferase activity is associated with accumulation of cholesterol (16-18). We demonstrated, however, that neither the induced reduction of the cholesterol content by the HMG-CoA reductase inhibitor atorvastatin nor the LDL-mediated overload of cholesterol, affected the ER-Golgi transport of the GlcNAc-1-phosphotransferase α/β-subunit precursor protein. Both immunofluorescence microscopy and western blots showed co-localization of immunoreactive α-subunits with the cis Golgi marker protein GM130, and the appearance of mature α-subunits, respectively (Fig. 3). Subsequently, neither the pattern of M6P modifications on lysosomal proteins, the transport kinetics of the newly synthesized lysosomal protease cathepsin Z, and its lysosomal steady state concentration, nor the activity of the lysosomal marker enzyme β-hexosaminidase, were altered in cells of low or high cholesterol content in comparison with control cells (Fig. 4). In a second control approach we used fibroblasts from GlcNAc-1-phosphotransferase ‘knock-in’ mice which mimic the biochemical and clinical symptoms of human MLII disease (18, 31, 32). Fibroblasts of the MLII mouse express truncated, inactive α-subunits that are localized in the ER due to the loss of the combinatorial ER exit motif (7, 18). The missorting and lysosomal depletion of the soluble Npc2 protein in these cells (33) result in lysosomal accumulation of non-esterified cholesterol (18). In consequence, the impaired egress from lysosomal compartments causes a lower cholesterol content in ER membranes which stimulates SREBP processing (13). In MLII cells, activated SREBP signaling leads to a 3-fold up-regulated LDLR expression that is further increased by atorvastatin treatment (Fig. 1; Fig. 2) indicating
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intact cholesterol sensing mechanism in these mutant cells.

The overexpression of the human α/β-subunit precursor protein of GlcNAc-1-phosphotransferase in MLII MEF, however, did not affect the transport to the Golgi apparatus and the S1P-mediated formation of mature α-subunits (Fig. 3C). As a second cell model accumulating massive amounts of cholesterol and other lipids, we used fibroblasts of patients with mutations in the polytopic membrane protein NPC1 that mediates the egress of non-esterified cholesterol from lysosomes to the ER and plasma membranes (19). Similar to MLII MEF, the expression of α/β-subunit precursor protein in NPC1 patient fibroblasts, resulted in the cleavage into mature subunits (Fig. 3D) confirming the independency of the α/β-subunit precursor transport from the ER to the Golgi apparatus of the cholesterol content.

Of interest, with the exception of LDL overload of WT MEF both the induced alterations in the cellular cholesterol content as well as the lysosomal accumulation of cholesterol in NPC1 fibroblasts failed to impair the receptor-mediated internalization and the transport of an M6P-containing lysosomal enzyme, arylsulfatase B (ASB), along the endocytic pathway, or the proteolytic processing into mature enzyme forms (Fig. 5A-C). It is likely that the increased cellular cholesterol content in LDL overloaded WT MEF results in a moderate redistribution of Mpr300 toward the endosomal compartment with a subsequent higher number of receptors recycling via the cell surface. In MLII MEF, however, the internalization rate of ASB is increased due to the elevated MPR300 level (Fig. 6A) caused by higher Mpr300 mRNA level (33), and the intermediate and mature forms of ASB generated by lysosomal protease(s) accumulate. This observation is unexpected, because the majority of lysosomal proteases, such as cathepsin D, L, B, S, C, reach lysosomes in MLII cells in an M6P/sortilin-independent manner, with the exception of cathepsin Z (Fig. 4B and C), carboxypeptidase Q, and dipeptidylpeptidase 7 (33). It remains to be examined whether these missorted peptidases/proteases are involved in the degradation of ASB polypeptides, or of the 17-kDa LDLR fragments (Fig. 6B) formed in the process of proprotein convertase subtilisin/kexin type 9 (PCSK9) initiated and mediated internalization and degradation of LDLR (26, 34).

Similarly to SREBPs which are retained in the ER by SCAP, the ER exit of another S1P substrate, ATF6, is prevented by binding of the ER chaperone BiP that dissociates in response to ER stress and allows the
translocation of ATF6 to the Golgi apparatus (35). Our data showed that the transport of the α/β-subunit precursor of GlcNAc-1-phosphotransferase is independent of cholesterol and ER stress. However, mutational analysis suggests that the transport of the α/β-subunit precursor is not constitutive, but depends on the binding of a yet unknown protein in the lumen of the ER, required for the ER transport of the precursor protein to the Golgi apparatus (36).

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GlcNAc-1-phosphotransferase interaction domain and an alternative site-1 protease cleavage site.

*unpublished results S. Klünder

*unpublished results S. Klünder

FIGURE LEGENDS

Fig. 1. Effect of ATV and LDL on Ldlr mRNA expression in fibroblasts. MEF from WT and MLII mice were cultured in DMEM containing 10% NBCS or 10% LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 48 h. The relative mRNA level of Ldlr and for comparison of Gnptab were determined by real-time PCR and normalized to β-actin mRNA expression. The relative mRNA expression in non-treated WT cells, incubated in DMEM containing 10% NBCS, was set 1. The data are the mean of triplicate PCRs obtained from three independent experiments and expressed as the fold change ± SD. ***P ≤ 0.005, **P ≤ 0.001, *P ≤ 0.05.

Fig. 2. Effect of ATV and LDL on LDLR protein expression in fibroblasts. MEF from (A) WT and (B) MLII were cultured in DMEM containing 10% NBCS or 10% LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 48 h. The protein expression of LDLR was analyzed by western blotting and evaluated by densitometry. The LDLR expression in cells incubated with DMEM containing NBCS was set 1. The positions of the molecular mass marker proteins (in kDa) are indicated. The content of Gapdh in the samples served as loading control. Representative western blots of four experiments (n = 4) are shown. ***P ≤ 0.005, **P ≤ 0.001.

Fig. 3. ATV or LDL overload did neither affect the localization nor proteolytic cleavage of the GlcNAc-1-phosphotransferase α*/β-subunit precursor protein. A-E: human fibroblasts of healthy controls and NPC1 patients, WT and MLII MEF were cultured in DMEM containing 10% NBCS or 10% LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 24 h as indicated. Cells were then transfected with cDNA of α*/β-subunit precursor construct and analyzed after 24 h. A and B: the localization of α*/β-subunit precursor of the GlcNAc-1-phosphotransferase (α*/β, green) was determined in (A) control and (B) NPC1 patient 1 (P1) fibroblasts by co-staining with the cis-Golgi marker protein GM130 (red) using immunofluorescence microscopy. Co-localization in merged images appears yellow. Scale bars, 5 µm. C
and D: the proteolytic cleavage of the α*/β-subunit precursor as indicator for correct transport to the Golgi apparatus was analyzed by α-subunit western blot of transfected (C) WT and MLII MEF, (D) human control and NPC1 (P1) fibroblasts, and (E) three different NPC1 cell lines (P1, P2 and P3). Gapdh/GAPDH western blot analysis, respectively, was used as a loading control. Extracts of non-transfected cells were used as negative control. The positions of molecular mass marker proteins (in kDa), α*/β-subunit precursor and mature α*-subunit are indicated.

**Fig. 4.** ATV or LDL overload did not affect GlcNAc-1-phosphotransferase activity and transport of lysosomal enzymes. A-D: MEF (WT and MLII) were cultured in DMEM containing 10% NBCS or 10% LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 48 h. A, the GlcNAc-1-phosphotransferase activity in cell extracts was indirectly determined by M6P western blotting. The positions of the molecular mass markers (in kDa) are indicated. B: the biosynthesis and sorting of the lysosomal protease cathepsin Z (CtsZ) was analyzed by labeling of MEFs with [35S]methionine for 1 h followed by 4 h further incubation in non-radioactive medium and immunoprecipitation of CtsZ from cell extracts and media, SDS-PAGE and fluorography. The amounts of secreted CtsZ precursors were estimated by densitometry and expressed as percentage of totally synthesized CtsZ. The positions of the molecular mass marker proteins (in kDa), precursor (p) and mature (m) form of CtsZ are indicated. The identity of the 45-kDa 35S-labeled polypeptide immunoprecipitated from cell extracts (●) is unknown. C: the total intracellular CtsZ expression at steady state was analyzed by western blotting. Anti β-tubulin western blot was used as a loading control. The positions of the molecular mass markers (in kDa) are indicated. D: the relative enzyme activity of the lysosomal enzyme β-hexosaminidase was measured in cell extracts and media conditioned for 24 h. The activities in control WT MEF incubated with DMEM containing 10% NBCS were set 1. The specific activity of β-hexosaminidase in these cells and media were 57.7 mU/h/mg protein ± 14.0 and 4.1 mU/24h/ mg protein ± 1.3, respectively. ***P ≤ 0.005
Cholesterol-independent M6P formation on lysosomal enzymes

**Fig. 5.** Endocytosis and proteolytic processing of arylsulfatase B. A: WT MEF were cultured in DMEM containing 10% NBCS or 10% LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 48 h. A-B: WT and MLII MEF were incubated with $[^{125}\text{I}]$ASB (625,000 cpm/ml) in the presence or absence of 10 mM M6P for 20 min, washed and either harvested (−) or chased for the indicated time points. n = 3 C: human healthy control and NPC1 (P1) fibroblasts were incubated with $[^{125}\text{I}]$ASB (625,000 cpm/ml), washed and either harvested (−) or chased in the presence or absence of 100 nM bafilomycin A1 (Baf A1). Cell extracts containing equal amounts of β-hexosaminidase activity (A: 21.0 mU/h/mg protein ± 3.1, B: 22 mU/mg protein ± 1.9 for WT and 4.5 mU/mg protein ± 0.6 for MLII, C: 19.4 mU/mg protein ± 3.3) were separated by SDS-PAGE and the internalized ASB was visualized by autoradiography. The positions of the molecular mass markers (in kDa) and the migration of precursor (p) and mature (arrow) forms of ASB are indicated.

**Fig. 6.** Accumulation of LDLR in MLII embryonic fibroblasts. MEF (WT and MLII) were cultured in DMEM containing NBCS or LPDS supplemented with 10 µM ATV or 100 µg/ml human LDL for 48 h. A: cell extracts were separated by SDS-PAGE (5% acrylamide) under non-reducing conditions and analyzed by Mpr300 western blotting. B: cell extracts were separated by SDS-PAGE (15% acrylamide) followed by LDLR western blotting. Afterwards the membrane was stripped and reused for Gapdh western blotting as loading control. Western blot analyses were repeated twice with similar results. The positions of the molecular mass markers (in kDa) and the Ldlr fragments are indicated. * unspecific polypeptide band.
Figure 1

Cholesterol-independent M6P formation on lysosomal enzymes
Figure 2

Cholesterol-independent M6P formation on lysosomal enzymes
Figure 3

Cholesterol-independent M6P formation on lysosomal enzymes
Figure 4

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28
**Figure 5**

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Cholesterol-independent M6P formation on lysosomal enzymes
Cholesterol-independent M6P formation on lysosomal enzymes

Figure 6