Quantitative proteomics and differential protein abundance analysis after depletion of PEX3 from human cells identifies additional aspects of protein targeting to the ER

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Abstract: Protein import into the endoplasmic reticulum (ER) is the first step in the biogenesis of about 10,000 different soluble and membrane proteins in humans. It involves co- or post-translational targeting of precursor polypeptides to the ER and their subsequent membrane insertion or translocation. So far, three pathways for ER targeting of precursor polypeptides plus four pathways for ER targeting of mRNAs were described. Typically, these pathways deliver their substrates to the Sec61 polypeptide-conducting channel in the ER membrane. Next, the precursor polypeptides are inserted into the ER membrane or translocated into the ER lumen, which may involve auxiliary translocation components, such as the TRAP and Sec62/Sec63 complexes, or auxiliary membrane protein insertases, such as EMC and the TMCO1 complex. Recently, the PEX19/PEX3-dependent pathway, which has a well-known function in targeting and inserting various peroxisomal membrane proteins into pre-existent peroxisomal membranes, was also found to act in targeting and, putatively, inserting monotopic hairpin proteins into the ER. These either remain in the ER as resident ER membrane proteins or are pinched off from the ER as components of new lipid droplets. Therefore, the question arose if this pathway may play a more general role in ER protein targeting, i.e. represents a fourth pathway for ER targeting of precursor polypeptides. Thus, we addressed the client spectrum of the PEX19/PEX3-dependent pathway in both PEX3-depleted HeLa cells and PEX3-deficient Zellweger patient fibroblasts by an established approach, which involves label-free quantitative mass spectrometry of the total proteome of depleted or deficient cells and differential protein abundance analysis. The negatively affected proteins included twelve peroxisomal proteins and two hairpin proteins of the ER, thus confirming two previously identified classes of putative PEX19/PEX3-clients in human cells. Interestingly, fourteen collagen-related proteins with signal peptides or N-terminal transmembrane helices and belonging to the secretory pathway were also negatively affected by PEX3-deficiency, which may suggest compromised collagen biogenesis as a hitherto unknown contributor to organ failures in the respective Zellweger patients.

Keywords: endoplasmic reticulum; lipid droplets; peroxisomes; PEX3; protein targeting; membrane protein insertion; protein translocation; label-free quantitative mass spectrometry; differential protein abundance analysis, Zellweger syndrome.
1. Introduction

In analogy to the division of the human body into several organs, the nucleated human cell is divided into various different compartments, the cell organelles, which are surrounded and, thereby, separated from the cytosol by biological membranes. However, the vast majority of the approximately 30,000 types of polypeptides plus their isoforms of a human cell is synthesized in the cytosol. Therefore, proteins of the different organelles have to be sorted to the correct organelles and, subsequently, inserted into or translocated across the organelar membrane(s). The protein import into the endoplasmic reticulum (ER) is the first step in the biogenesis of about 10,000 different soluble and membrane proteins of human cells [1-4]. These were found to fulfill their functions in the membrane or lumen of the ER plus the nuclear envelope, in one of the organelles belonging to the pathways of endo- and exocytosis (i.e. ERGIC, Golgi apparatus, endosome, lysosome, transport vesicles) or at the cell surface as plasma membrane- or secretory-proteins. Excluding resident proteins of the ER, most of the correctly folded and assembled proteins are delivered from the ER to their functional location by vesicular transport, which involves vesicles budding off from subdomains of the tubular ER that are termed exit sites (ERES) [5,6]. In recent years, however, an increasing number of proteins destined to lipid droplets (LDs), peroxisomes or mitochondria was observed to be targeted to the ER as well prior to their integration into budding LDs or peroxisomes or prior to their delivery to mitochondria via the recently-identified ER-SURF pathway [7-11]. LDs and peroxisomes are ER-derived organelles and their biogenesis occurs in specialized subdomains of the tubular ER [8,10].

Protein import into the ER involves ER membrane targeting as the initial step and insertion of nascent or fully-synthesized membrane proteins into or translocation of soluble precursor polypeptides across the ER membrane as the second step [1-4]. Typically, both processes depend on N-terminal signal peptides (SPs) or transmembrane helices (TMHs) in the precursor polypeptides that serve as signals [4,12-14]. Generally, the Sec61 complex of the ER membrane represents the entry point for most of these precursor polypeptides into the organelle (Table 1) [1-4]. A variety of proteins rely on the Sec61 complex for their integration into the ER lumen or their translocation across the ER membrane (Figure 1a). These include SP-containing soluble or GPI-anchored proteins, bitopic type I (C-in N-out) and type II (N-in C-out) proteins as well as multispanning membrane proteins. Sec61 is also implicated in the membrane insertion of proteins without N-terminal SPs such as type III (C-in N-out) [4] and C-tail-anchored (TA) (N-in C-out) membrane proteins [15-17] and even monotopic hairpin proteins (C-in N-in) [18,19]. In these cases, however, the role of Sec61 is less clear and auxiliary factors such as the TRAP complex, the Sec62/Sec63 complex, or insertases such as the ER membrane protein complex (EMC), or the TMCO1 complex are required in addition [4]. Especially the membrane insertion of TA-proteins predominantly relies on the insertase WRB/CAML [15-17]. In conclusion, several co- as well as post-translational protein targeting pathways merge at the Sec61 complex in the ER membrane, including the co-translational SRP/SR-pathway and the posttranslational SRP-independent or SND-pathway [2,4,20-26].

In addition, there is targeting of mRNAs to the ER membrane, which involves mRNA receptors (such as KTN1), or receptors for ribosome nascent chain complexes with nascent chains, which are not yet long enough to be able to interact with SRP (such as RRBP1) [27-33]. However, one general lesson from the analysis of all these pathways is that they are not strictly separated from each other and that there are at least some precursor polypeptides, which can be targeted to the ER by more than one pathway (such as some small presecretory proteins and some tail-anchored membrane proteins) [23,25,26]. Thus, the targeting pathways have overlapping substrate specificities and can substitute for each other at least to a certain extent. Characterization of all these pathways and mechanisms is also of medical importance since several of the components are linked to human hereditary or tumor diseases or being hijacked by viral or bacterial agents [34].
Recent work identified the PEX19/PEX3-dependent pathway as a fourth pathway for ER targeting of precursor polypeptides [35,36]. PEX3 (also termed peroxisomal biogenesis factor 3 or Peroxin-3) was first identified in yeast and is a membrane protein with a N-terminal transmembrane domain plus a large C-terminal domain, which is facing the cytosol both in yeast and in humans [37-41]. Originally, it was characterized as peroxisomal membrane protein, which cooperates with the cytosolic protein PEX19 in targeting of peroxisomal membrane proteins to pre-existent peroxisomes and in facilitating their membrane insertion [38,40]. However, PEX3 is also present in discrete subdomains of ER membranes and is involved in targeting of an unknown number of precursor proteins to ER membranes and, possibly, in their membrane insertion [35,36]. So far, these precursor proteins include membrane proteins, which either remain in the ER (the two-hairpin or reticulon-domain containing proteins ARL6IP1, RTN3A, RTN4C) [35] or are pinched off in LDs (such as the hairpin protein UBXD8) [36]. At the ER, PEX3 cooperates with the farnesylated variant of PEX19 [36,39]. Thus, farnesylation of PEX19 is most likely decisive in delivering precursor polypeptides to either pre-existent peroxisomes or the ER [7,8].

These observations raised the question if this pathway may play a more general role in ER protein targeting. Defects in the human PEX3 gene are linked to a particularly devastating form of Zellweger syndrome, which belongs to the peroxisome biogenesis disorders and is also termed cerebro-hepato-renal syndrome to indicate the most important affected organs [41-44]. Infants with the disease typically die within their first year of life because of the complete absence of peroxisomes in all cells of the body.

Here, we address the client spectrum of PEX3 in ER protein targeting in human cells and, simultaneously, the question if the PEX19/PEX3-dependent pathway to the ER can also target precursor polypeptides to the Sec61 complex. The approach involves transiently PEX3-depleted HeLa cells or chronically PEX3-deficient Zellweger patient fibroblasts in combination with differential proteomic analysis by label-free quantitative mass spectrometry (MS) and differential protein abundance analysis (Figure 1b). Thus, we report on negatively and positively affected proteins after partial depletion of PEX3 in HeLa cells and in PEX3-deficient Zellweger patient fibroblasts.

Table 1. Protein transport components and associated proteins in HeLa cells and linked diseases.

| Component/subunit          | Abundance¹ | Localization² | Linked Diseases                                  |
|----------------------------|------------|---------------|-------------------------------------------------|
| for ER targeting           |            |               |                                                 |
| #p34 (LRC59, LRRC59)³      | 2480       | ERM           |                                                 |
| #p180 (RRBP1)              | 135        | ERM           | Hepatocellular Carcinoma, Colorectal Cancer     |
| Kinectin 1 (KTN1)          | 263        | ERM           |                                                 |
| AEG-1 (LYRIC, MTDH)        | 575        | ERM           |                                                 |
| #SRP⁴                      |            | C             |                                                 |
| - SRP72                    | 355        |               | Aplasia, Myelodysplasia                         |
| - SRP68                    | 197        |               |                                                 |
| - SRP54                    | 228        |               | Neutropenia, Pancreas Insufficiency             |
| - SRP19                    | 33         |               |                                                 |
| - SRP14                    | 4295       |               |                                                 |
| - SRP9                     | 3436       |               |                                                 |
| - 7SL RNA                  |            |               |                                                 |
| SRP receptor               |            |               |                                                 |
| - SRα#(docking protein)    | 249        | ERM           |                                                 |
| - SRβ                      | 173        |               |                                                 |
| Calmodulin                 | 9428       | C             |                                                 |

for ER targeting and, possibly, for membrane integration
| **hSnd1** | unknown |
|---|---|
| **Snd receptor** | ERM |
| - **hSnd2 (TMEM208)** | 81 |
| - **hSnd3##** | 49 |
| **PEX9** | 80 |
| **PEX3** | 103 |
| **for ER targeting** | **Zellweger Syndrome** |
| **plus membrane integration** |  |
| **#Bag6 complex** | **C** |
| - **TRC35 (Get4)** | 171 |
| - **Ub14A** | 177 |
| - **Bag6 (Bat3)** | 133 |
| **SGTA** | 549 |
| **TRC40 (Asna1, Get3)** | 381 |
| **TA receptor** | **ERM** |
| - **CAML (CAMLG, Get2)** | 5 |
| - **WRB (CHD5, Get1)** | 4 |
| **for ER membrane integration** |  |
| **ERM protein complex** | **ERM** |
| - **EMC1** | 124 |
| - **EMC2** | 300 |
| - **EMC3** | 270 |
| - **EMC4** | 70 |
| - **EMC5 (MMGT1)** | 35 |
| - **EMC6 (TMEM93)** | 5 |
| - **EMC7** | 247 |
| - **EMC8** | 209 |
| - **EMC9** | 1 |
| - **EMC10** | 3 |
| **#TMCO1 complex##** | **ERM** |
| - **TMCO1** | 2013 |
| - **Nicalin** | 99 |
| - **TMEM147** | 21 |
| - **CCDC47 (Calumin)** | 193 |
| - **NOMO** | 267 |
| **PAT complex** | **ERM** |
| - **PAT10 (Asterix)** |  |
| - **CCDC47 (Calumin)** | 193 |
| **for ER membrane integration** |  |
| **plus translocation** |  |
| **#Sec61 complex##** | **ERM** |
| - **Sec61α1** | 139 |
| - **Sec61β** | 456 |
| - **Sec61γ** | 400 |
| **#Sec62 (TLOC1)** | 26 |
| **ER Chaperones** | **ERM** |
| - **Sec63 (ERj2)** | 168 |
| - **αErj1 (DNAJC1)** | 8 |
| - **BiP (Grp78, HSPA5)** | 8253 |
| - **Grp170 (HYOU1)** | 923 |
| - **Sil1 (BAP)** | 149 |
| **#Calnexin palmitoylated** | 7278 |
| **#TRAP complex** | **ERM** |
| - **TRAPαα(SSR1)** | 568 |
| - **TRAPββ(SSR2)** |  |
| - **TRAPγγ(SSR3)** | 1701 |

**Diabetes**, **CVID**

**CDG, Hepatocellular Carcinoma**
| Protein Name | Abundance (nM) | Localization |
|--------------|----------------|--------------|
| TRAPα (SSR4) | 3212           | CDG          |
| RAMP4 (SERP1) |                | ERM          |

### for covalent modification

#### Oligosaccharyltransferase (OST-A)
- Ribophorin I (Rpn1) 1956 ERM
- Ribophorin II (Rpn2) 527 ERM
- OST48 273 CDG
- Dad1 464 ERM
- OST4 1956 ERM
- TMEM258 527 ERM
- Stt3A* 430 CDG
- DC2 1956 ERM
- Kcp2

#### Oligosaccharyltransferase (OST-B)
- Ribophorin I (Rpn1) 1956 ERM
- Ribophorin II (Rpn2) 527 ERM
- OST48 273 CDG
- Dad1 464 ERM
- OST4 1956 ERM
- TMEM258 527 ERM
- Stt3B* 150 CDG
- TUSC3 1956 ERM
- MagT1 33 ERM

#### Signal peptidase (SPC-A)
- SPC12 2733 ERM
- SPC18* (SEC11A) 334 ERM
- SPC22/23 94 ERM
- SPC25 94 ERM

#### Signal peptidase (SPC-C)
- SPC12 2733 ERM
- SPC21* (SEC11C) 334 ERM
- SPC22/23 94 ERM
- SPC25 94 ERM

#### GPI transamidase (GPI-T)
- GPAA1 9 ERM
- PIG-K 38 ERM
- PIG-S 86 ERM
- PIG-T 20 ERM
- PIG-U 42 ERM

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1. Abundance refers to the concentration (nM) of the respective protein in HeLa cells, as reported by Hein et al. [45].
2. Localization refers to the functional intracellular localization(s) of the respective protein [1-4,33,36,41], i.e. C, Cytosol, ERL, ER lumen, ERM, ER membrane, PexM, Peroxisome membrane.
3. Alternative protein names are given in parentheses.
4. Complexes are indicated by italics. Abbreviations for protein names, EMC, ER membrane (protein) complex, GET, Guided entry of tail-anchored proteins, SEC, (Protein involved in) secretion, SND, SRP-independent, SR, SRP receptor, SRP, Signal recognition particle, SSR, Signal sequence receptor, TMEM, Transmembrane (protein), TRAM, translocating chain-associating membrane (protein), TRAP, Translocon-associated protein, TRC, Transmembrane recognition complex.
5. Diabetes was linked to the particular protein in mouse.
6. Abbreviation for diseases, i.e. CDG, Congenital disorder of glycosylation, CVID, Common variable immunodeficiency, GBM, Glioblastoma multiforme, HUS, Hemolytic-uremic syndrome, MSS, Marinesco-Sjögren syndrome, PLD, Polycystic liver disease, TKD, Tubulointerstitial kidney disease, as reported by Sicking et al. [34].
Figure 1. Types of ER membrane proteins and our experimental strategy to address their biogenesis. (a) The cartoon depicts a signal peptide (SP) (in yellow) and six types of ER membrane proteins (MP) (in black), together with their membrane protein type and mechanism of membrane insertion (both indicated below the cartoon). Cleavable SPs (in yellow) can facilitate ER import of secretory proteins (in green), glycosylphosphatidylinositol (GPI)-anchored membrane proteins (in green) and several types of membrane proteins, including single-spanning type I membrane proteins. Positively charged amino acid residues (+) play an important role in membrane protein and SP orientation, i.e. typically follow the positive inside rule [14]. Amino-terminal transmembrane helices (TMHs) can serve as signal-anchor sequences to facilitate membrane insertion of type II, type III- and many multi-spanning membrane proteins. In the case of membrane proteins with amino-terminal TMHs, membrane insertion typically involves the same components and mechanisms, which deliver secretory proteins (in green) and GPI-anchored membrane proteins (in green) to the ER lumen. The central component here is the Sec61 complex. In some cases, however, auxiliary membrane protein insertases, such as EMC or TMCO1 complex play a role. These can also operate as stand-alone membrane protein insertases, an activity they have in common with the WRB/CAML complex [4]. Hairpin (HP) proteins have a monotopic topology with N- and C-termini facing the cytosol and some of them require PEX3 for membrane targeting. C, carboxy-terminus; N, amino-terminus. (b) The experimental strategy was as follows: siRNA-mediated gene silencing using two different siRNAs for each target and one non-targeting (control) siRNA, respectively, with three replicates for each siRNA for 96 h; label-free quantitative analysis of the total cellular proteome; differential protein abundance analysis to identify negatively affected proteins (i.e. putative clients of the target) and positively affected proteins (i.e. putative compensatory mechanisms); validation by western blot. In addition, PEX3-deficient Zellweger patient cells were analyzed in triplicates.
2. Results

2.1. Quantitative proteomic analysis of HeLa cells after transient and partial depletion of PEX3 by siRNA

Our approach to characterize the client spectrum of PEX3 in ER protein targeting involves gene silencing of the putative receptor PEX3 in HeLa Kyoto cells with two different targeting siRNAs in parallel to a non-targeting or control siRNA and differential proteomic analysis by label-free quantitative MS analysis and differential protein abundance analysis (Figure 1b). This protocol was developed and previously used to characterize the client spectrum and client SP features of ER protein translocation components including Sec61 complex (as a proof of concept), TRAP complex, Sec62/Sec63 complex, TRAM1- protein, ERj1, plus BiP, and the mRNA targeting components KTN1 and RRBP1 [33,46-48]. The approach is based on the assumption that polypeptide precursors, which have to be imported into the ER, are degraded by the proteasome in the cytosol upon interference with their ER targeting or translocation because their SPs or TMHs are not easily compatible with the aqueous character of the cytosol. Therefore, their cellular levels are decreased as compared to control cells, and this change is detected by quantitative MS and subsequent differential protein abundance analysis [46]. Typically, the decrease was observed to be accompanied by an increase of ubiquitin-conjugating enzymes [33,46-48]. Furthermore, a simultaneous increase in other ER import components was detected, which is consistent with the overlap of pathways and, additionally, may indicate a genetic interaction between different pathways and cellular compensation.

Here, we applied the established experimental strategy to identify precursor polypeptides that may depend on PEX3-dependent targeting to the ER [46]. They were expected among the negatively affected proteins in the label-free quantitative MS and subsequent differential protein abundance analysis. HeLa cells were treated in triplicates with two different PEX3-targeting siRNAs (PEX3 #1 siRNA, PEX3 #2 siRNA) in parallel to a non-targeting (control) siRNA for 96 h. Each MS experiment provided proteome-wide abundance data as LFQ intensities for three sample groups—one control (non-targeting siRNA treated) and two stimuli (down-regulation by two different targeting siRNAs directed against the same gene)—each having three data points (Figure 1b). To identify which proteins were affected by knock-down in siRNA-treated cells relative to the non-targeting (control) siRNA treated sample, we log2-transformed the ratio between siRNA and control siRNA samples, and performed two separate unpaired t-tests for each siRNA against the control siRNA sample according to [46]. The p values obtained by unpaired t-tests were corrected for multiple testing using a permutation false discovery rate (FDR) test. Proteins with an FDR-adjusted p value of below 5% were considered significantly affected by knock-down of the targeted protein.

After PEX3 depletion, 6,488 different proteins were quantitatively detected by MS in all samples (Figure S1, Table 2, Tables S1 – S2). The MS data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository [49] with the dataset identifier: PXD012005 (http://www.proteomexchange.org; last accessed May 2, 2021). They included the expected representation of proteins with cleaved SP (7%), N-glycosylated proteins (9%), and membrane proteins (13%), which was comparable to the previously published Sec61 or TRAP experiment (Figures S1c, left small pies) [46]. Applying the established statistical analysis, we found that transient PEX3 depletion significantly affected the steady-state levels of 13 proteins: 13 negatively and null positively (permutation false discovery rate-adjusted p value < 0.05). As had to be expected, PEX3 itself was negatively affected (Figure S1a, volcano plots), which was confirmed by western blot (Figure S1b). The identified precursors included one protein with cleavable SP (the ER membrane protein AGPAT1), and two membrane proteins with TMH (not counting PEX3), i.e. the endo- and lysosomal membrane protein TMEM192 with four transmembrane domains and the single-pass type II plasma
membrane protein SGCD (Figure S1b, Table S2). Of these three negatively affected proteins one is N-glycosylated (SGCD). There were no proteins with annotated functional location in peroxisomes or LDs among the negatively affected proteins. There was no positively affected protein observed. These results raise the question why PEX3 depletion from HeLa cells had hardly any effect on the cellular proteome. There are several possibilities. The simplest answer would be that the depletion efficiency of 85% may not have been high enough to cause accumulation of precursor proteins. Another answer could be that PEX3 function in ER protein import in HeLa cells is not essential, i.e., can be substituted by other proteins or pathways. Another possibility is that the accumulating precursors were not degraded but aggregated in the cytosol or ended up in other organelles, where they were protected from degradation. These possibilities will be considered in further detail in the discussion.

Table 2. Statistics for the identification of putative PEX3-clients in comparison to previously identified clients for ER membrane targeting and translocation components.

| Proteins | PEX3 | Z1 | RRBPl2 | KTN12 | SEC612 | TRAP2 | CDG1,2 |
|----------|------|----|--------|-------|--------|-------|--------|
| Quantified proteins | 8178 | 6328 | 4813 | 4947 | 7212 | 7670 | 5920 |
| Statistically analyzed proteins | 6488 | 6328 | 4813 | 4947 | 5129 | 5911 | 5920 |
| representing the secretory pathway (%) | 29 | 26 | 27 | 26 | 27 | 36 |
| Proteins with SP (%) | 7 | 6 | 6 | 6 | 7 | nd |
| N-Glycoproteins (%) | 9 | 8 | 8 | 8 | 8 | nd |
| Membrane proteins (%) | 13 | 12 | 13 | 12 | 13 | nd |
| Positively affected proteins | 0 | 97 | 157 | 25 | 342 | 77 | 39 |
| Negatively affected proteins | 13 | 141 | 141 | 45 | 482 | 180 | 279 |
| representing the secretory pathway (%) | 54 | 39 | 37 | 41 | 61 | 40 | 36 |
| Negatively affected proteins with SP (%) | 8 | 19 | 18 | 7 | 41 | 22 | 12 |
| Negatively affected N-glycoproteins (%) | 8 | 21 | 17 | 18 | 45 | 23 | 17 |
| Negatively affected membrane proteins (%) | 31 | 18 | 18 | 22 | 36 | 26 | 23 |
| Negatively affected proteins with SP | 1 | 27 | 21 | 3 | 197 | 38 | 34 |
| including N-glycoproteins | 0 | 23 | 16 | 3 | 158 | 28 | 30 |
| corresponding to % | | 0 | 85 | 76 | 100 | 80 | 74 | 88 |
| including membrane proteins | 1 | 6 | 16 | 1 | 77 | 19 | 16 |
| corresponding to % | | 100 | 22 | 29 | 33 | 39 | 50 | 53 |
| Negatively affected proteins with TMH | 3 | 16 | 18 | 8 | 98 | 22 | 41 |
| including N-glycoproteins | 1 | 6 | 7 | 4 | 56 | 11 | 17 |
| corresponding to % | | 33 | 38 | 39 | 50 | 57 | 50 | 41 |
| Negatively affected peroxisomal proteins | 1 | 12 | 0 | 1 | 1 | 0 | 1 |
| corresponding to % | 8 | 9 | nd | 2 | 0 | nd | 0 |
| including membrane proteins | 1 | 6 | nd | 0 | 1 | 0 | 1 |
| corresponding to % | | 100 | 50 | nd | nd | 100 | nd | 100 |
| Negatively affected mitochondrial proteins | 0 | 14 | 6 | 1 | 29 | 14 | 21 |
| corresponding to % | nd | 10 | 4 | 2 | 1 | 1 | 1 |
| including membrane proteins | nd | 4 | 3 | 0 | 11 | 3 | 8 |
| corresponding to % | nd | 29 | 50 | nd | 38 | 21 | 38 |

1 Z and CDG refer to immortalized fibroblasts from patients, suffering from Zellweger syndrome or Congenital disorder of glycosylation. 2 refers to siRNA-mediated knock-down HeLa cells and was previously published [33,46]. 3 nd, not determined.
2.2. Quantitative proteomic analysis of PEX3-deficient Zellweger patient fibroblasts

In the course of our previous analysis of Sec62- and Sec63-clients we moved from the respective siRNA-treated and incompletely depleted HeLa cells with low client numbers on to CRISPR/Cas9-treated and deficient HEK293 cells and, indeed, identified many more clients [47]. Therefore, we sought to test whether a complete PEX3 knock out in cells leads to the depletion of putative PEX3-client proteins. Therefore, we subjected control fibroblasts and immortalized Zellweger patient fibroblasts with PEX3-deficiency [41], which had been grown in triplicates, to label-free quantitative proteomic analysis and differential protein abundance analysis and analyzed the data for negatively affected proteins, i.e. potential PEX3-clients (Figure 2, Table 2, Tables S3 – S5). The MS data have been deposited to the ProteomeXchange Consortium via the PRIDE partner repository [49] with the dataset identifier: PXD012005 (http://www.proteomexchange.org; last accessed May 2, 2021).

We quantitatively identified a total of 6,328 different proteins by MS, 141 of which were negatively affected by PEX3-deficiency in the patient fibroblasts versus control fibroblasts. As had to be expected, PEX3 itself was negatively affected (Figure 2a, volcano plots), which was confirmed by western blot (Figure 2b). Applying the established statistical analysis, we found that PEX3 deficiency significantly affected the steady-state levels of 238 proteins: 141 negatively and 97 positively (permutation false discovery rate-adjusted p value < 0.05). Of the negatively affected proteins, GO terms assigned 39.2% to organelles of the endocytic and exocytic pathways (Figure 2c, large pies), which corresponds to a 1.36-fold enrichment (Figure 2c, large pies, 39.2% divided by 28.91% = 1.36) and is below the average values of 1.46 and 1.94 observed after depletion of the mRNA targeting components KTN1 (1.55) plus RRBP1 (1.37) and the translocation components Sec61 (2.37) plus TRAP (1.5), respectively (Table 2) [33,46]. In contrast to the PEX3-knock-down cells (Figure S1c), we also detected enrichment of proteins with SP (2.65-fold), N-glycosylated proteins (2.2-fold), and membrane proteins (1.36–fold) (Figure 2c, small pies), which was lower as compared to the Sec61- (6.51, 2.83, 2.51) or TRAP-experiment (3.3, 2.7, 2.1) but higher as compared to the KTN1- (1, 2.09, 1.76) and RRBP1-experiment (2.44, 2.12, 1.46) [33,46].

The negatively affected proteins of the secretory pathway included 27 proteins with cleavable SP (including six collagens or collagen-like proteins, three collagen modifying proteins, three ER luminal proteins FKBP7, PCSK9, PDIA5, the lysosomal cathepsin CTSB, eight secretory proteins, the ER membrane protein PLOD2, and the plasma membrane proteins ENPP4, HLA-C, ICAM1, ITGB5, LRRC15 (Figure 2a).

Furthermore, 15 membrane proteins of the secretory pathway with TMH (not counting PEX3) were negatively affected, i.e. most notably the two ER-resident hairpin proteins ATL1 and RTN3 as well as the three tail anchored proteins CCDC136, STX6 plus VAMP3, and the ER membrane proteins DHR57B, ERMP1, TMUB2, the Golgi protein MAN1A1, as well as the plasma membrane proteins AIFM2 (which has additional locations, see below), COLEC12, CYBRD1, ENPP1, TMEM237, and the nuclear envelope protein TOR1AIP1 (Figure 2a, Tables 3 and Table S4). Of these 42 negatively affected proteins, 29 are N-glycosylated proteins (23 with SP and 6 with TMH).

Interestingly, there were 14 precursors of mitochondrial proteins negatively affected by PEX3-deficiency (Table S4), a phenomenon previously observed after depletion of RRBP1 from HeLa cells and attributed to their physiological trafficking from the ER to mitochondria via the newly identified ER-SURF pathway [11,33]. Among these negatively affected mitochondrial proteins were three outer membrane proteins (AIFM2, RHOST1, VAT1), one inner membrane protein (NDUFV3), and ten matrix proteins, two of which have a dual localization in mitochondria and peroxisomes (ACAD11, SCP2). Alternatively, our observation of mitochondrial proteins among the negatively affected ones may be due to the findings that at least PEX19 associates with peroxisomes and mitochondria and, for example, facilitates protein biogenesis of tail anchored membrane proteins Fis1 and Gem1 for both organelles [50],
Positively affected proteins in the PEX3-deficient fibroblasts included cell adhesion molecules (such as ITGA1, L1CAM, NCAM1), signal transduction components of the plasma membrane (ANXA3, PKD2), mitochondrial membrane proteins of the inner membrane with functions in protein import into mitochondria (DNAJC15) or protein quality control (OMA1), and the cytosolic chaperone HSPB6.

2.3. Negatively affected precursor proteins in Zellweger patient fibroblasts are specific for PEX3-deficiency and partially affected in PEX3 depleted HeLa cells

To validate the proteomic data on putative PEX3-substrates, we conducted independent western blot experiments with the PEX3-deficient Zellweger- and control-fibroblasts for the SP-containing candidate PDIA5, the hairpin protein RTN3, the dual topology LD/peroxisome protein Far1, and the peroxisomal protein ACBD5 (Figure 3). All of these proteins were depleted in PEX3-deficient fibroblasts, fully confirming the proteomic analysis and verifying them as putative PEX3-clients. Interestingly, for RTN3 we observed that specifically the 100 kDa isoform was depleted in the PEX3-deficient cells, while the 25 kDa isoform remained largely unaltered (Figure 3, e versus f).

For PEX3 depleted HeLa cells, we observed that 17 of the 48 negative hits were not quantified and that 18 out of 31 putative PEX3 negative clients in Zellweger patient fibroblasts (i.e. 58%) were also negatively affected by PEX3 depletion in HeLa cells (including two out of three tail anchored proteins, one out of two hairpin proteins, and four out of six peroxisomal proteins), but did not meet the stringent significance threshold (Table S6 and proteins indicated in red in Figure 2). Notably, the PEX3 depletion in HeLa cells was not as efficient as in fibroblasts (log2-fold change: -3.5 versus -4.6) (Figures 2 and S1). Hence, it is understandable that the levels of the putative PEX3-clients are less perturbed in HeLa cells than in fibroblasts. Still, the majority of them (18 versus 13) are perturbed in the same negative direction as in fibroblasts (Table S1).

We consider these putative clients of PEX3 for organelles of the secretory pathway as specific for additional reasons. First, the negatively affected proteins included, as expected, twelve precursors of proteins with a functional location in peroxisomes (not counting PEX3), including six peroxisomal membrane proteins (Figure 2a) (Table 3). The peroxisomal membrane proteins were ABCD3, ACBD5, AGPS, FAR1, PEX13, and PXMP2. Notably, the tail anchored membrane protein FAR1 exhibits a dual topology and can locate to peroxisomes as well as to LDs [51]. Second, with two hairpin proteins, ATL1 and RTN3, among the negatively affected proteins, we also confirmed a second class of already known PEX3-clients in human cells under physiological conditions (Table 3). Third, only one of the negatively affected peroxisomal proteins, PEX13, had previously been observed for TRAP-deficient fibroblasts from patients who suffer from congenital disorders of glycosylation (CDG) and are either SSR3 (coding for TRAPγ) or SSR4 (TRAPδ) deficient [46]. Furthermore, there was no overlap in the positively affected proteins that accumulate in either CDG- or Zellweger-patient fibroblasts.

As outlined in Table 2 and for comparison with the PEX3-deficient Zellweger patient fibroblasts, 5919 different proteins were previously quantified for CDG patient fibroblasts, 279 of which were negatively affected by TRAP absence, 39 were positively affected. 100 of the negatively affected proteins were assigned to the secretory pathway, including 34 precursor polypeptides with SP and 41 with TMH (including the subunits of the heterotetrameric TRAP complex) [46]. A total of 47 of the negatively affected proteins were N-glycoproteins (30 with SP and 17 with TMH). The peroxisomal membrane protein PEX13 was among the negatively affected proteins, which was also negatively affected in PEX3-deficient Zellweger patient fibroblasts (Table 2).
PEX3-deficient Zellweger patient fibroblasts

**Negatively affected proteins, i.e. putative PEX3 substrates**

with SP:
CCDC80, CDCP1, COL1A1, COL6A1, COL6A2, COL6A3, COL11A1, CTHRC1, CTSB, DHRSX, ENP44, FBLN1, FKBP7, HLA-C, HTRA1, ICAM1, ITGB5, LAMA4, LEPREL1, LOX, LRRC15, MFGE8, PCOLCE, PCSK9, Pdia5, PLOD2, PRSS23,

with TMH:
AIFM2, COLEC12, CYBRD1, DHRS7B, ENPP1, ERMP1, MAN1A1, TMEM237, TMUB2, TOR1AIP1

with tail anchor:
CCDC136, STX6, VAMP3

with hairpin(s):
ATL1, RTN3

of peroxisomal membranes:
ABCD3, ACBD5, AGPS, FAR1, PEX13, PXMP2

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**Notably, the proteins in red were also negatively affected by PEX3 depletion in HeLa cells but did not meet the stringent significance threshold.**

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**C**

**Total quantified proteome**

- nucleus: 28.91%
- mitochondrion: 9.36%
- cytosol: 12.86%
- ribosome: 7.24%
- cytoskeleton: 20.57%
- peroxisome: 39.20%
- plasma membrane: 18.44%
- ER: 19.15%
- golgi: 20.57%
- extracellular: 19.15%
- lysosome: 20.57%
- endosome: 19.15%
- vacuole: 19.15%

**Negatively affected proteome**

- signal peptide: 7.24%
- N-glycosylated protein: 20.57%
- membrane protein: 18.44%
Figure 2 (on the previous page). Volcano plots and Gene Ontology (GO) enrichment for PEX3-deficient Zellweger patient fibroblasts. (a) Differentially affected proteins were characterized by the mean difference of their intensities plotted against the respective permutation false discovery rate-adjusted p-values in volcano plots, PEX3 is highlighted. In addition, the proteins, which were negatively-affected by PEX3-deficiency are given in the right panel. (b) PEX3-deficiency was evaluated by western blot. Molecular mass values are indicated in kilodaltons (KDa). Only the area of interest of the blot is shown, the original images are shown in Supplementary Materials. (c) Classification of putative PEX3-clients was based on GO enrichment factors where the results from the complete set of quantified proteins in the left panel are compared with the negatively affected proteome. Protein annotations of SPs, membrane location, and N-glycosylation in humans were extracted from UniProtKB, and used to determine the enrichment of GO annotations among the negatively affected proteins.

Figure 3. Validation of PEX3-clients by western blot analyses. Three independent cell lysates from control- (WT) and PEX3-deficient fibroblasts (PEX3-/-), respectively, were analyzed by western blotting using antibodies as indicated. Left panels: Relevant sections of representative western blots are shown, tubulin served as a loading control. We note that the full scans of all blots are shown in the supplement. Right panels: Scatter plots indicate relative protein abundances in control- and PEX3-deficient fibroblasts as derived from quantitative western blots as shown in the left panels. Signals were quantified by densitometry and relative abundances were calculated as the ratio of the signal of interest to the corresponding tubulin signal in the same lane and normalized against one control sample. Mean values with SEM from three independent lysates per cell line are indicated as well as the individual data points for each replicate.
Table 3. Negatively affected proteins in PEX3-deficient cells, i.e. putative PEX3 substrates.

| Gene   | Subcellular location                                      | Membrane protein type                      | SS or TMH |
|--------|-----------------------------------------------------------|---------------------------------------------|-----------|
| ACBD5  | Peroxisome membrane                                      | Single-spanning membrane protein            |           |
| COLEC12| Membrane                                                 | Single-spanning type II membrane protein    | TMH       |
| LRRC15 | Membrane                                                 | Single-spanning type I membrane protein     | SP        |
| PEX3   | Peroxisome membrane                                      | Single-spanning membrane protein            |           |
| TOR1AIP1| Nuclear envelope inner membrane                           | Single-spanning membrane protein            | TMH       |
| COL1A1 | Secreted, Extracellular space, Extracellular matrix       | Single-spanning membrane protein            | SP        |
| AGPS   | Peroxisome membrane                                      |                                             |           |
| ACAD11 | Peroxisome, Mitochondrion                                |                                             |           |
| STX6   | Golgi apparatus membrane                                  | Tail anchored membrane protein              | tail anchor|
| CCDC136| Acrosome membrane, Secretory vesicle, Cytoplasmic vesicle| Tail anchored membrane protein              | tail anchor|
| FAR1   | Peroxisome membrane                                      | Tail anchored membrane protein              | tail anchor|
| PXMP2  | Peroxisome membrane                                      | Multi-spanning membrane protein             |           |
| ATL1   | Cell projection, Golgi apparatus membrane, Axon           | Hairpin membrane protein with one hairpin   | hairpin   |
| COL6A2 | Extracellular matrix, Membrane, Secreted, Extracellular space|                                         | SP        |
| LOX    | Extracellular space, Secreted                            |                                             | SP        |
| ERMP1  | ER membrane                                              | Multi-spanning membrane protein             | TMH       |
| CYBRD1 | Membrane                                                 | Multi-spanning membrane protein             | TMH       |
| TMUB2  | Membrane                                                 | Multi-spanning membrane protein             | TMH       |
| ABCD3  | Peroxisome membrane                                      | Multi-spanning membrane protein             |           |
| SCP2   | Peroxisome, Mitochondrion, Cytoplasm                     |                                             |           |
| CDCP1  | Secreted, Cell membrane                                  |                                             | SP        |
| COL6A3 | Extracellular space, Secreted, Extracellular matrix      |                                             | SP        |
| TMEM237| Cell projection, Membrane, Cilium                        | Multi-spanning membrane protein             | TMH       |
| ENPP4  | Cell membrane                                            | Single-spanning type I membrane protein     | SP        |
| HTRA1  | Cell membrane, Secreted, Cytoplasm, Cytosol              |                                             | SP        |
| VAMP3  | Synapse, Membrane, Cell junction, Synaptosome             | Tail anchored membrane protein              | tail anchor|
| MFGE8  | Membrane, Secreted                                       |                                             | SP        |
| PRSS23 | Secreted                                                 |                                             | SP        |
| DHR54  | Peroxisome, Nucleus                                      |                                             |           |
| ITGB5  | Membrane                                                 | Single-pass type I membrane protein         | SP        |
| FBLN1  | Extracellular space, Secreted, Extracellular matrix      |                                             | SP        |
| COL6A1 | Extracellular space, Secreted, Extracellular matrix      |                                             | SP        |
| PCSK9  | Endosome, Golgi apparatus, Cell surface, Secreted, ER, Lysosome|                                         | SP        |
Proteins are listed according to decreasing negative effects of PEX3 depletion. The colors refer to peroxisomal proteins (yellow), mitochondrial proteins (brown), proteins of the secretory pathway with SP, TMH, tail anchor (green) or hairpin (orange). As compared to Table S4, the GO annotation for TOR1AIP1, the hairpin of RTN3 as well as the definitions of membrane protein type were taken from GeneCards ([https://www.genecards.org](https://www.genecards.org); last accessed September 1, 2021). In addition, the term TMH is used here only for proteins of the secretory pathway. Red letters refer to incomplete annotations (see text for details).
3. Discussion

Here we addressed the question which precursor polypeptides employ the PEX3-dependent pathway for targeting to or insertion and translocation, respectively, into the ER of human cells. First, we employed our previously established approach of siRNA-mediated depletion of PEX3 in HeLa cells, label-free quantitative MS of the total cellular proteome and differential protein abundance analysis. Next, we quantified negatively and positively affected proteins under conditions of PEX3-deficiency in Zellweger patient fibroblasts.

On first sight, the result of siRNA-mediated depletion of PEX3 in HeLa cells was not very informative (Figure S1). In addition to the depletion of PEX3, only three precursor polypeptides with SP or TMH were found among the negatively affected proteins and may be considered as PEX3-clients in ER protein targeting. Notably, the one with a SP, AGPAT1, is an important enzyme in lipid metabolism and required for synthesis of phosphatidic acid and triacylglycerides [52]. Therefore, it may affect LD biogenesis and LD localization would not be unexpected. The proteins with a TMH are the endo- and lysosomal membrane protein TMEM192 with four transmembrane domains and the single-spanning type II plasma membrane protein SGCD. There was no peroxisomal protein negatively affected by PEX3 depletion, which has to be interpreted...
in light of the facts that PEX19 and PEX3 are essential for peroxisome formation [37,38] and that PEX3-deficiency causes the complete absence of peroxisomes [42-44]. Together, these results raise the question why PEX3 depletion from HeLa cells had hardly any effect on the cellular proteome. There are several possibilities. i) The simplest answer would be that the depletion efficiency and its duration may not have been sufficient to cause significant accumulation of precursor proteins. According to the MS data the log2 fold change was -3.4942, i.e. the depletion efficiency was higher than 90%, which is consistent with the western blot analysis (Figure S1b). This residual amount of PEX3, however, may have been sufficient for physiological functions and could explain the absence of an effect on peroxisomal proteins. ii) Another answer may be that PEX3 function in ER protein import in HeLa cells is not essential, i.e. may also be provided by other proteins or pathways. Indeed, it was shown in cell-free ER protein import studies that certain peroxisomal membrane proteins can be targeted to the mammalian ER by SRP or TRC40 (including PEX3) [35,53-55]. Furthermore, some collagens as well as some hairpin membrane proteins were previously observed as RRBP1-clients (ATL2, ATL3, COL1A1, COL1A2, COL4A2) and SRP-clients (ARL6IP1, RTN3), respectively, in cell biological or proteomic experiments [30,33,35]. iii) Another possibility is that some accumulating precursors were not degraded but either stayed soluble in the cytosol (as is known for catalase), aggregated, or ended up in other organelles (as is known for PEX14) [41], where they were protected from degradation. Indeed, we have previously observed mis-targeting of certain precursors of secretory proteins into mitochondria in the absence of Sec61 function in HeLa cells [56]. Furthermore, the known PEX3-client UBXD8 accumulates in mitochondria when PEX3 function is compromised [36]. iv) Last but not least, all three possibilities may have contributed to the result we obtained in HeLa cells upon siRNA-mediated knock-down and we consider this the most likely explanation.

Under conditions of PEX3-deficiency in Zellweger patient fibroblasts, the results were more informative (Figure 2). First, the negatively affected proteins included twelve precursors of proteins with a functional location on or in peroxisomes, including six peroxisomal membrane proteins (Table 3). This was expected and demonstrated once more the feasibility of the approach. However, this does not mean that all these precursors of peroxisomal proteins are targeted to the ER. Rather, at least for the peroxisomal matrix proteins it must be due to the complete absence of peroxisomes from the patient fibroblasts with PEX3-deficiency [42-44]. Likewise, we cannot rule out that the depletion of some precursors could derive from transcriptional effects. However, we consider this unlikely since previous transcriptomics studies revealed that solely the absence of peroxisomes does not generally result in global mRNA expression changes [57].

Second, the identified precursors included the two ER-resident hairpin proteins ATL1 (with one hairpin) and RTN3 (with two hairpins), which is consistent with the previous findings that PEX3 is involved in ER targeting of hairpin proteins [35,36]. It remains to be tested whether PEX3, possibly in cooperation with PEX16 [41], also facilitates membrane insertion of these hairpin proteins and whether additional membrane protein insertases, such as Sec61, EMC, TMCO1, and WRB/CAML, contribute to membrane insertion. Likewise, it is conceivable that also the members from the DHRS- and ACBD-protein families (ACBD5, DHR54, DHR7b, and DHR8X; Figure 2, Table 3) are first inserted into the ER membrane by PEX3 and additional insertion components of the ER. This will also have to be addressed in future research.

Furthermore, 14 α-helical membrane proteins including four tail-anchored membrane proteins (CCDC136, FAR1, STX6 and VAMP3) and ten others (not counting PEX3), i.e. DHR7b (type II), ERMP1 (multi-pass), TMUB2 (multi-pass), MAN1A1 (type II), AIFM2, COLEC12 (type II), CYBRD1 (multi-pass), ENPP1 (type II), TOR1AIP1 (single-pass), and TMEM237 (multi-pass) were negatively affected by PEX3 absence and, therefore, can be considered as potential PEX3-clients in ER protein targeting (Table 3).
Importantly, the annotated membrane topology of these potential clients should be considered with care, as they may be, in certain cases, incomplete. For example, it was recently shown that the tail anchored membrane protein FAR1 exhibits a dual topology, including a monotopic hairpin topology, and can therefore, locate to peroxisomes as well as to LDs [51]. Since ERMP1, STX6, and TOR1AIP1 were previously found to depend on Sec61 for their membrane insertion in HeLa cells, it is tempting to speculate that PEX3 is able to target precursors to the Sec61 complex.

In addition, the identified precursors included 27 proteins with cleavable SP, including the five collagens and one collagen-like protein, four collagen modifying proteins LEPREL1, three ER lumenal proteins, and the four plasma membrane proteins with a connection to the extracellular matrix or neighboring cells CDCP1, ICAM1, ITGB5, LRRC15. Thus between 14 and 17 of the total of 43 secretory pathway precursor proteins are collagens or related to collagens, which is a significant enrichment of 33 – 40%. This raises the question of how PEX3 could be involved in targeting of these precursors, which are expected to involve a cotranslational targeting pathway to the Sec61 complex. We hypothesize that in these particular cases, PEX3 may act in concert with mRNA targeting pathways, which would be consistent with the observation that RRPB1-mediated targeting pathways were found to be involved in the biogenesis of these proteins [30,33]. However, their degradation in the absence of PEX3 may have an alternative explanation. We speculate that the combination of farnesylated PEX19 plus ER-membrane resident PEX3 targets the membrane-shaping and hairpin(s)-containing membrane proteins (such as Atlastins, Reticulons and Spastin) to the PEX3-rich ER subdomain and that this enrichment of hairpin proteins creates an environment, which attracts collagens as well as some of their modifying enzymes and future interaction partners [6,58-61]. Consistent with this scenario, the key player of the formation of large cargo secretory vesicles at ER exit sites, the membrane protein TANGO1, has two transmembrane domains, one of which is supposed to form a hairpin in the inner leaflet of the ER membrane [59,60]. Either way, a common ER subdomain may be conducive to budding of both peroxisomal precursor vesicles and large cargo secretory vesicles as well as the formation of LDs. Indeed, very recent evidence suggests, that membrane bridges between ER exit sites and LDs allow protein partitioning of hairpin proteins from the ER to LDs (doi: https://doi.org/10.1101/2021.09.14.460330). In addition, this observation raises the question if defects in collagen biogenesis contribute to the devastating effects of PEX3 deficiency in Zellweger patients and clearly warrants further work.

Furthermore, negatively affected proteins in PEX3-depleted or -deficient human cells need to be discussed with respect to LD biogenesis from a functional point of view (Figure 3, Table 3). DHRS- and ACBD-family proteins, which were negatively affected in PEX3-deficient fibroblasts (Table 3) and the dehydrogenase/reductase SDR family members 4, X and 7B (DHRS4, DHRSX, DHRS7B) play important roles in retinol biosynthesis and may also localize to LDs [62]. The latter was suggested for DHRSX and, actually, shown for the family members DHRS3 and DHRS7B [62] (doi: https://doi.org/10.1101/2021.09.14.460330). On the other hand, acyl-CoA-binding domain-containing proteins 5 and 7 (peroxisomal ACBD5, cytosolic ACBD7) are involved in organelle contacts and, therefore, may be important for lipid metabolism and/or organelle budding from the ER. These aspects, too, warrant further studies.

Taken together, our study revealed a putative client spectrum for PEX3-mediated protein targeting to the ER, including several unexpected protein classes such as secreted collagens. Importantly, several PEX3-client candidates are involved in lipid metabolic pathways or membrane shaping mechanisms, which affect ER-derived organelle biogenesis of peroxisomes and/or LDs. As observed for other ER targeting pathways, also PEX3 likely shares some of its putative clients with other ER targeting components including the Sec61 complex.
4. Materials and Methods

4.1. Cell growth and analysis.

HeLa Kyoto cells [36] were cultivated at 37°C in a humidified environment with 5% CO₂, in DMEM with 10% fetal bovine serum (FBS; Sigma-Aldrich). Cell growth and viability were monitored using the Countess® Automated Cell Counter (Invitrogen) following the manufacturer’s instructions.

For gene silencing, 4 × 10⁵ HeLa cells were seeded per 6-cm culture plate, followed by incubation under normal culture conditions. Next, the cells were transfected with either PEX3-targeting Silencer Select pre-designed siRNA (Life Technologies, IDs s16154 and s16156) or with a scrambled siRNA control (Life Technologies ID 4390843) at a final concentration of 3.3 nM using Lipofectamine 2000 (Life Technologies) and following the manufacturer’s instructions. After 48 h the cells were transfected a second time and grown for additional 48 h. Thus, silencing was performed for a total of 96 h using two different siRNAs. We note that cell viability was not affected by PEX3 depletion for 96h, i.e. the viability values were virtually identical for the two PEX3-targeting siRNAs (95.0% for RNA#1 and 96.0% for RNA#2, n = 3) versus scrambled siRNA control (95.7%).

Silencing efficiencies were evaluated by western blot analysis using PEX3-specific antibodies [63] (1:1,000 dilution), which were kindly donated by Prof. Gabriele Dodt (University of Tübingen, Tübingen, Germany) and anti-tubulin antibodies (T6199, Sigma-Aldrich; 1:10,000 dilution). Donkey-derived, Cy3- and Alexa488-conjugated secondary antibodies (715-165-151, 711-545-152, Jackson Immunoresearch) were detected using the Typhoon-Trio imaging system combined with Image Quant TL software 7.0 (GE Healthcare).

Immortalized PEX3-deficient fibroblasts and control fibroblasts have been obtained from Prof. Gabriele Dodt (University of Tübingen, Tübingen, Germany) and were previously characterized [41,64]. They were cultivated at 37°C in a humidified environment with 5% CO₂, in DMEM/GlutaMAX with 10% fetal bovine serum (FBS; Sigma-Aldrich) for 72 h.

4.2. Label-free quantitative proteomic analysis.

After growth for 96 h, 1 × 10⁶ cells (corresponding to roughly 0.2 mg protein) were harvested, washed twice in PBS, and lysed in buffer containing 6 M GnHCl, 20 mM tris(2-carboxyethyl)phosphine (TCEP; Pierce™, Thermo Fisher Scientific), 40 mM 2-chloroacetamide (CAA; Sigma-Aldrich) in 100 mM Tris, pH 8.0 [33,46-48]. The lysate was heated to 95°C for 2 min, and then sonicated in a Bioruptor sonicator (Diagenode, Seraing, Belgium) at the maximum power setting for 10 cycles of 30 s each. For a 10% aliquot of the sample, the entire process of heating and sonication was repeated once, and then the sample was diluted 10-fold with digestion buffer (25 mM Tris, pH 8, 10% acetonitrile). Protein extracts were digested for 4 h with Lysyl endoproteinase Lys-C (Wako Bioproducts, Fujifilm, Neuss, Germany, enzyme to protein ratio: 1:50), followed by the addition of trypsin (Promega) for overnight digestion (enzyme to protein ratio: 1:100). The next day, booster digestion was performed for 4 h using an additional dose of trypsin (enzyme to protein ratio: 1:100). After digestion, a 10% aliquot of peptides (corresponding to about 2 µg of peptides) were purified via SDB-RPS StageTips [65], eluted as one fraction, and loaded for MS analysis. Purified samples were loaded onto a 50-cm column (inner diameter: 75 microns; packed in-house with ReproSil-Pur C18-AQ 1.9-micron beads, Dr. Maisch HPLC GmbH, Ammerbuch, Germany) via the autosampler of the Thermo Easy-nLC 1000 (Thermo Fisher Scientific) at 60°C. Using the nanoelectrospray interface, eluting peptides were directly sprayed onto the benchtop Orbitrap mass spectrometer Q Exactive HF (Thermo Fisher Scientific) [66]. Peptides were loaded in buffer A (0.1% (v/v) formic acid) at 250 nL/min and the percentage of buffer B was ramped to 30% over 180 min, followed by a ramp to 60 % over 20 min, then 95 % over the next 10 min, and maintained at 95% for another 5 min [33,47]. The mass spectrometer was operated in a data-dependent mode with survey scans from 300 to 1,700 m/z (resolu-
tion of 60,000 at m/z = 200). Up to 15 of the top precursors were selected and fragmented using higher energy collisional dissociation (HCD) with a normalized collision energy value of 28 [33,47]. The MS2 spectra were recorded at a resolution of 17,500 (at m/z = 200). AGC target for MS and MS2 scans were set to 3E6 and 1E5, respectively, within a maximum injection time of 100 and 25 ms for MS and MS2 scans, respectively. Dynamic exclusion was enabled to minimize repeated sequencing of the same precursor ions and set to 30 s [33,47].

4.4. Data analysis

Raw data were processed using the MaxQuant computational platform [67]. The peak list was searched against Human Uniprot databases and proteins were quantified across samples using the label-free quantification algorithm in MaxQuant as label-free quantification (LFQ) intensities [68]. We note that LFQ intensities do not reflect true copy numbers because they depend not only on the amounts of the peptides but also on their ionization efficiencies; thus, they only served to compare abundances of the same protein in different samples [66-71]. Each MS experiment provided proteome-wide abundance data as LFQ intensities for three sample groups—one control (non-targeting siRNA treated) and two stimuli (down-regulation by two different targeting siRNAs directed against the same gene)—each having three data points. Missing data points were generated by imputation, whereby we distinguished two cases [46]. To identify which proteins were affected by PEX3 knock-down in siRNA-treated cells relative to the non-targeting (control) siRNA treated sample, we log2-transformed the ratio between siRNA and control siRNA samples, and performed two separate unpaired t-tests for each siRNA against the control siRNA sample [46]. The p values obtained by unpaired t-tests were corrected for multiple testing using a permutation false discovery rate (FDR) test. Proteins with an FDR-adjusted p value of below 5% were considered significantly affected by knock-down of the targeted protein. The results from the two unpaired t-tests were then intersected for further analysis meaning that the abundance of all reported candidates was statistically significantly affected in both siRNA silencing experiments. For completely missing proteins lacking any valid data points, imputed data points were randomly generated in the bottom tail of the whole proteomics distribution, following the strategy in the Perseus software (http://maxquant.net/perseus/; last accessed May 2, 2021) [70]. For proteins having at least one valid MS data point, missing data points were generated from the valid data points based on the local least squares (LLS) imputation method [71]. The validity of this approach was demonstrated [46]. Subsequent to data imputation, gene-based quantile normalization was applied to homogenize the abundance distributions of each protein with respect to statistical properties. All statistical analyses were performed using the R package SAM (https://statweb.stanford.edu/~tibs/SAM/; last accessed May 2, 2021) [72]. Protein annotations of signal peptides, transmembrane regions, and N-glycosylation sites in humans and yeast were extracted from UniProtKB entries using custom scripts [46]. The enrichment of functional Gene Ontology annotations (cellular components and biological processes) among the secondarily affected proteins was computed using the GOrilla package [73].

4.5. Validation of putative PEX3 substrates by quantitative western blotting:

Hits from the proteomics analysis were validated by quantitative western blot analyses of Triton-X100 cell lysates (1% Triton X-100, 50mM Hepes pH 7.5, 150mM NaCl, 10% glycerol, 1 mM EDTA, 1 mM PMSF, Complete EDTA-free protease inhibitors (Roche)) using the following antibodies: anti-PEX3 (gift from G. Dodt; 1:1000 dilution), anti-ACBD5 (HPA012145 Merck; 1:1000 dilution), anti-RTN3 (12055-2-AP Proteintech; 1:1000 dilution), anti-PDIA5 (15545-1 Proteintech; 1:1000 dilution), anti-Far1 (ATA-HPA017322 Biozol, 1:1250 dilution), and anti-tubulin (T6199, Sigma-Aldrich; 1:10,000 dilution), which served as a loading control for normalization. Secondary antibodies were purchased from Licor Biosciences (926-68020, 926-68021, 926-32211,
Signals were detected using the Odyssey CLx system from Licor Biosciences and quantified by densitometry using the Image Studio software (Licor Biosciences). Relative protein abundance was calculated as the ratio of the signal of interest to the corresponding tubulin signal in the same lane and normalized against one control sample. Visualization of the quantification data was performed with the Graphpad Prism software.

5. Conclusions

Recent studies characterized the PEX19/PEX3 pathway, which is best known for its role in the biogenesis of peroxisomal membrane proteins both at the peroxisomal and the ER membrane, as being involved in the biogenesis of hairpin membrane proteins of the ER as well as LDs. Therefore, the question arose if this pathway may play a more general role in ER protein targeting, i.e. represent a fourth pathway for ER targeting of precursor polypeptides next to SRP, SND, and TRC/GET. We have started to address this question by a novel approach, which involves label-free quantitative mass spectrometry of the total proteome of depleted or deficient cells and differential protein abundance analysis. Thus, we addressed the client spectrum of the PEX19/PEX3-dependent pathway in both PEX3 targeting siRNA-treated HeLa cells and PEX3-deficient Zellweger patient fibroblasts. The negatively affected proteins included six peroxisomal membrane proteins and two hairpin proteins of the ER, thus confirming the two previously identified classes of putative PEX19/PEX3-clients for ER targeting in human cells as well as the validity of the experimental approach. In addition, 14 membrane proteins (including four tail anchored proteins) and 27 proteins with SP (including 14 collagens plus collagen-related proteins) and belonging to the secretory pathway were also negatively affected by PEX3-deficiency. The latter findings are consistent with the idea that PEX3 represents a fourth pathway for targeting of precursor polypeptides to the Sec61 complex. Furthermore, it may suggest a hitherto unknown spatial or at least physical relationship between ER subdomains that are involved in ER shaping and the budding of peroxisomal precursor vesicles, large cargo vesicles, and lipid droplets. In addition, these results may suggest compromised collagen biogenesis as a hitherto unknown contributor to organ failures in the respective Zellweger patients. All these suggestions will have to be addressed in future research.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Table S1_pex_3_all, Table S2_pex_3_full_lo, Table S3_pex_3_Zellweger_all, Table S4_pex_3_Zellweger_full_lo, Table S5_pex_3_Zellweger_full_up, Table S6_proteomeXchange_identifiers, PEX3_Description of Supplementary Tables; MS blots PEX3.

Author Contributions: R.Z. and B.S. planned and supervised the sample generation for MS analysis and validation experiments by M.L. D.N. performed the MS data analysis under supervision by V.H. F.F. and R.Z. designed the study. R.Z. and B.S. wrote the manuscript together with F.F., S.L. and V.H. All authors discussed the results and have read and agreed to the published version of the manuscript.

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Abbreviations

- ATL: Atlastin
- CDG: Congenital disorder of glycosylation
- EMC: ER membrane complex
- ER: Endoplasmic reticulum
- GET: Guided entry of tail-anchored proteins
- GO: Gene ontology
- GPI: Glycosylphosphatidylinositol
- LD: Lipid droplet
- PEX: Peroxin
- RAMP: Ribosome-associated membrane protein
- RNC: Ribosome-nascent chain complex
- RTN: Reticulon
- SEC: (Protein involved in) secretion
- SND: SRP-independent
- SP: Signal peptide
- SR: SRP receptor
- SRP: Signal recognition particle
- SSR: Signal sequence receptor
- TMEM: Transmembrane (protein)
- TMH: Transmembrane helix
- TRAM: Translocating chain-associating membrane (protein)
- TRAP: Translocon-associated protein
- TRC: Transmembrane recognition complex
- Z: Zellweger (patient fibroblasts)

References

1. Dudek, J.; Pfeffer, S.; Lee, P.-H.; Jung, M.; Cavalié, A.; Helms, V.; Förster, F.; Zimmermann, R. Protein transport into the human endoplasmic reticulum. *J. Mol. Biol.* 2015, 427, 1159-1175. doi: 10.1016/j.jmb.2014.06.011
2. Aviram, N.; Schuldiner, M. Targeting and translocation of proteins to the endoplasmic reticulum at a glance. *J. Cell Sci.* 2017, 130, 4079-4085. doi: 10.1242/jcs204396
3. Gemmer, M.; Förster, F. A clearer picture of the ER translocon complex. *J. Cell Sci.* 2020, 133, jcs231340. doi: 10.1242/jcs.231340
4. O’Keefe, S.; Pool, M.R.; High, S. Membrane protein biogenesis at the ER: the highways and byways. *FEBS J.* 2021, doi: 10.1111/febs.15905
5. Budnik, A.; Stephens, D.J. ER exit sites - Localization and control of COPII vesicle formation. *FEBS Lett.* 2009, 583, 3796–3803. doi: 10.1016/j.febslet.2009.10.038
6. Stephens, D.J. Collagen secretion explained. *Nature* 2012, 482, 474-475. doi: 10.1038/482474a
7. Schrul, B.; Schliebs, W. Intracellular communication between lipid droplets and peroxisomes: the Janus face of PEX19. *Biol. Chem.* 2018, doi: 10.1515/bch-2018-0125
8. Jansen, R.L.M.; van der Klei, I.J. The peroxisome biogenesis factors Pex3 and Pex19: multitasking proteins with disputed functions. *FEBS Lett.* 2019, 593, 457-474. doi: 10.1002/1873-3468.13340
9. Dhimann, R.; Caesar, S.; Thiam, A.R.; Schrul, B. Mechanisms of protein targeting to lipid droplets: A unified cell biological and biophysical perspective. *Sem. Cell Dev. Biol.* **2020**, *108*, 4-13. doi: 10.1016/j.semcdb.2020.03.004

10. Goodman, J.M. Building the lipid droplet assembly complex. *J. Cell Biol.* **2020**, *219*, e202006025. doi: 10.1083/jcb.202006025

11. Hansen, K.G.; Aviram, N.; Laborenz, J.; Bibi, C.; Meyer, M.; Spang, A.; Schuldiner, M.; Herrmann, J.M. An ER surface retrieval pathway safeguards the import of mitochondrial membrane proteins in yeast. *Science* **2018**, *361*, 1118-122. doi: 10.1126/science.aar8174

12. Hegde, R.S.; Bernstein, H. The surprising complexity of signal peptides. *Trends Biochem. Sci.* **2006**, *31*, 563-571. doi: 10.1016/j.tibs.2006.08.004

13. Goder, V.; Spiess, M. Molecular mechanism of signal sequence orientation in the endoplasmic reticulum. *EMBO J.* **2003**, *22*, 1470-1478. doi: 10.1093/emboj/cdg361

14. Goder, V.; Junne, T.; Spiess, M. Sec61p contributes to signal sequence orientation according to the positive-inside rule. *Mol. Biol. Cell* **2004**, *15*, 1470-1478. doi:10.1091/mbc.e03-08-0599

15. Borgese, N.; Fasana, E. Targeting pathways of C-tail-anchored proteins. *Biochim. Biophys. Acta* **2011**, *1808*, 937-946.

16. Yamamoto, Y.; Sakisaka, T. Molecular machinery for insertion of tail-anchored membrane proteins into the endoplasmic reticulum membrane in mammalian cells. *Mol. Cell* **2012**, *48*, 387-397. doi: 10.1016/j.molcel.2012.08.028

17. Borgese, N.; Coy-Vergara, J.; Colombo, S.F.; Schwappach, B. The ways of tails: the GET pathway and more. *Proteins* **2019**, *38*, 289-305. doi: 10.1007/s10930-019-09845-4

18. Abell, B.M.; High, S; Moloney, M.M. Membrane protein topology of oleosin is constrained by its long hydrophobic domain. *J. Biol. Chem.* **2002**, *277*, 8602-8610. doi: 10.1074/jbc.M103712200

19. Leznicki, P.; Schneider, H.O.; Harvey, J.V.; Shi, W.Q.; High, S. Co-translational biogenesis of lipid droplet integral membrane proteins. *J. Cell Sci.* **2021**, *132*, jcs.259220. doi: 10.1242/jcs.259220

20. Gamerdinger, M.; Hanebuth, M.A.; Frickey, T.; Deuerling, E. The principle of antagonism ensures protein targeting specificity at the endoplasmic reticulum. *Science* **2015**, *348*, 201-207. doi: 10.1126/science.aaa5335

21. Hsieh, H.-H.; Lee, J.H.; Chandrasekar, S.; Shan, S-ou. A ribosome-associated chaperone enables substrate triage in a cotranslational protein targeting complex. *Nat. Commun.* **2020**, *11*, 5840. doi: 10.1038/s41467-020-19548-5

22. Aviram, N.; Ast, T.; Costa, E.A.; Arakel, E.; Chuartzman, S.G.; Jan, C.H.; Haßdenteufel, S.; Dudek, J.; Jung, M.; Schorr, S.; Zimmermann, R., Schwappach, B.; Weissman, J.S.; Schuldiner, M. The SND proteins constitute an alternative targeting route to the endoplasmic reticulum. *Nature* **2016**, *540*, 134-138. doi: 10.1038/nature20169

23. Casson, J.; McKenna, M.; Haßdenteufel, S.; Aviram, N.; Zimmermann, R.; High, S. Multiple pathways facilitate the biogenesis of mammalian tail-anchored proteins. *J. Cell Sci.* **2017**, *130*, 3851-3861. doi: 10.1242/jcs.207829

24. Haßdenteufel, S.; Sicking, M.; Schorr, S.; Aviram, N.; Fecher-Trost, C.; Schuldiner, M.; Jung, M.; Zimmermann, R.; Lang, S. hSnd2 protein represents an alternative targeting factor to the endoplasmic reticulum in human cells. *FEBS Lett.* **2019**, *593*, 2506-2524. doi: 10.1002/1873-3468.13542

25. Haßdenteufel, S.; Johnson, N.; Paton, A.W.; Paton, J.C.; High, S.; Zimmermann, R. Chaperone-mediated Sec61 channel gating during ER import of small precursor proteins overcomes Sec61 inhibitor-reinforced energy barrier. *Cell Rep.* **2018**, *23*, 1373-1386. doi: 10.1016/j.celrep.2018.03.122

26. Haßdenteufel, S.; Nguyen, D.; Helms, V.; Lang, S.; Zimmermann, R. Components and mechanisms for ER import of small human presecretory proteins. *FEBS Lett.* **2019**, *593*, 2506-2524. doi: 10.1002/1873-3468.13542

27. Cui, X.A.; Zhang, H.; Palazzo, A.F. p180 promotes the ribosome-independent localization of a subset of mRNA to the endoplasmic reticulum. *PLoS Biol.* **2012**, *10*, e1001336. doi: 10.1371/journal.pbio.1001336.

28. Cui, X.A.; Zhang, Y.; Hong, S.J.; Palazzo, A.F. Identification of a region within the placental alkaline phosphatase mRNA that mediates p180-dependent targeting to the endoplasmic reticulum. *J. Biol. Chem.* **2013**, *288*, 29633-29641. doi: 10.1074/jbc.M113.482505
29. Calvin, H.J.; Williams, C.C.; Weissman, J.S. Principles of ER translocational translocation revealed by proximity-specific ribosome profiling. *Science* 2014, 346, 1257521. doi:10.1126/science.1257521

30. Ueno, T.; Tanaka, K.; Kaneko, K.; Taga, Y.; Sata, T.; Irie, S.; Shunji Hattori, S.; Ogawa-Goto, K. Enhancement of procollagen biosynthesis by p180 through augmented ribosome association on the endoplasmic reticulum in response to stimulated secretion. *J. Biol. Chem.* 2010, 285, 29942-29950. doi: 10.1074/jbc.M109.094607

31. Hsu, J.C.-C.; Reid, D.W.; Hoffman, A.M.; Sarkar, D.; Nicchitta, C.V. Oncoprotein AEG-1 is an endoplasmic reticulum RNA-binding protein whose interactome is enriched in organelle resident protein-encoding mRNAs. *RNA* 2018, 24, 688-703. doi: 10.1261/rna.063313.117

32. Hannigan, M.M.; Hoffman, A.M.; Thompson, J.W.; Zheng, T.; Nicchitta, C.V. Quantitative proteomics links the LRRC59 interactome to mRNA translation on the ER membrane. *Mol. Cell. Proteom.* 2020, 19, 1826-1849. doi: 10.1074/mcp.RA120.002228.

33. Bhadra, P.; Schorr, S.; Lerner, M.; Nguyen, D.; Dudek, J.; Förster, F.; Helms, V.; Lang, S.; Zimmermann, R. Quantitative proteomics and differential protein abundance analysis after depletion of putative mRNA receptors in the ER membrane of human cells identifies novel aspects of mRNA targeting to the ER. *Molecules*, 2021, 26, 3591. doi: 10.3390/molecules26123591

34. Sicking, M., Lang, S., Bochen, F., Drenth, J. P. H., Zacharia, M., Zimmermann, R., Roos, A., Linxweiler M. Cells 2021, 10, 1036. doi: 10.3390/cells10051036

35. Yamamoto, Y.; Sakisaka, T. The peroxisome biogenesis factors posttranslationally target reticulon homology-domain containing proteins to the endoplasmic reticulum membrane. *Sci. Rep.* 2018, 8, 2322. doi: 10.1038/s41598-018-20797-0

36. Schrul, B.; Kopito, R.R. Peroxin-dependent targeting of a lipid-droplet-destined membrane protein to ER subdomains. *Nat. Cell Biol.* 2016, 18, 740-751. doi: 10.1038/ncb3373

37. Dahan, N.; Francisco, T.; Falter, C.; Rodrigues, T.; Kael, V.; Kunze, M.; Hansen, T.; Schliebs, W.; Erdmann, R. Current advances in the function and biogenesis of peroxisomes and their roles in health and disease. *Hist. Cell Biol.* 2021, 155, 513-524. doi: 10.1007/s00418-021-01982-1

38. Schmel, R.; Veenhuis, M.; Mertens, D.; Kunau, W.-H. Isolation of peroxisome-deficient mutants of Saccharomyces cerevisiae. *Proc. Natl. Acad. Sci. USA* 1989, 86, 5419-5423. doi: 10.1073/pnas.86.14.5419

39. Götte, K.; Girzalsky, W.; Linkert, M.; Baumgart, E.; Kammerer, S.; Hunau, W.-H.; Erdmann, R. Pex19p, a farnesylated protein essential for peroxisome biogenesis. *Mol. Cell. Biol.* 1998, 18, 661-628. doi: 10.1128/mcb.18.1.616

40. Hettema, E.H.; Girzalsky, W.; van den Berg, M.; Erdmann, R.; Distel, B. Saccharomyces cerevisiae Pex3p and Pex19p are required for proper localization and stability of peroxisomal membrane proteins. *EMBO J.* 2000, 19, 223-233. doi: 10.1093/emboj/19.2.223

41. Schmidt, F.; Dietrich, D.; Eylenstein, R.; Groemping, Y.; Stehle, T., Dodt, G. The role of conserved PEX3 regions in PEX19-binding and peroxisome biogenesis. *Traffic* 2012, 13, 1244-1260. doi: 10.1111/j.1600-0854-2012.01380.x

42. Muntau, A.C.; Maerhofer, P.U.; Paton, B.C.; Kammerer, S.; Roscher, A.A. Defective peroxisome membrane synthesis due to mutations in human PEX3 causes Zellweger syndrome, complementation group G. *Am. J. Hum. Genet.* 2000, 67, 967-975.

43. Shimozawa, N.; Suzuki, Y.; Zhang, Z.; Imamura, A.; Ghaedi, K.; Fujiki, Y.; Kondo, N. Identification of PEX3 as the gene mutated in a Zellweger syndrome patient lacking peroxisomal remnant structures. *Hum. Mol. Genet.* 2000, 9, 1995-1999.

44. Ghaedi, K.; Honsho, M.; Shimozawa, N.; Suzuki, Y.; Kondo, N.; Fujiki, Y. PEX3 is the causal gene responsible for peroxisome membrane assembly-defective Zellweger syndrome of complementation group G. *Am. J. Hum. Genet.* 2000, 67, 976-981.

45. Hein, M.Y.; Hubner, N.C.; Poser, I.; Cox, J.; Nagaraj, N.; Toyoda, Y.; Gak, I.A.; Weisswange, I.; Mansfeld, J.; Buchholz, F.; Hyman, A.A.; Mann, M. A human interactome in three quantitative dimensions organized by stoichiometries and abundances. *Cell* 2015, 163, 712-723. doi: 10.1016/j.cell.2015.09.053
46. Nguyen, D.; Stutz, R.; Schorr, S.; Lang, S.; Pfeffer, S.; Freeze, H.F.; Förster, F.; Helms, V.; Dudek, J.; Zimmermann, R. Proteomics reveals signal peptide features determining the client specificity in human TRAP-dependent ER protein import. *Nat. Commun.* 2018, 9, 3763. doi: 10.1038/s41467-018-06188-z

47. Schorr, S.; Nguyen, D.; Haßdenteufel, S.; Nagaraj, N.; Cavalié, A.; Greiner, M.; Weissgerber, P.; Loi, M.; Paton, A.W.; Paton, J.C.; Molinari, M.; Förster, F.; Dudek, J.; Lang, S.; Helms, V.; Zimmermann, R. Proteomics identifies signal peptide features determining the substrate specificity in human Sec62/Sec63-dependent ER protein import. *FEBS J.* 2020, 287, 4612-4640. doi: 10.1111/febs.15274

48. Klein, M.-C.; Lerner, M.; Nguyen, D.; Pfeffer, S.; Dudek, J.; Förster, F.; Helms, V.; Lang, S.; Zimmermann, R. TRAM1 protein may support ER protein import by modulating the phospholipid bilayer near the lateral gate of the Sec61 channel. *Channels* 2020, 14, 28-44. doi: 10.1080/19336950.2020.1724759

49. Vizcaíno, J.A.; Csordas, A.; del-Toro, N.; Dianes, J.A.; Griss, J.; Lavidas, I.; Mayer, G.; Perez-Riverol, Y.; Reisinger, F.; Ternent, T.; Xu, Q.-W.; Wang, R.; Hermjakob, H. 2016 update of the PRIDE database and related tools. *Nucleic Acids Res.* 2016, D447-D456. doi: 10.1093/nar/gkw880

50. Cichocki, B.A.; Krumpe, K.; Vitali, D.G.; Rapaport, D. Pex19 is involved in importing dually targeted tail-anchored proteins to both mitochondria and peroxisomes. *Traffic* 2018, 19, 770-785. doi: 10.1111/tra.12604

51. Exner, T.; Romero-Brey, I.; Yifrach, E.; Rivera-Monroy, J.; Schril, B.; Zouboulis, C.C.; Stremmel, W.; Honsho, M.; Bartenschlager, R.; Zalckvar, E.; Füllekrug, J. An alternative membrane topology permits lipid droplet localization of peroxisomal fatty acyl-CoA reductase 1. *J. Cell Sci.* 2021, 132, jcs223016. doi: 10.1242/jcs.223016

52. Agarwal, A.K.; Sukumaran, S.; Cortés, V.A.; Tunison, K.; Mizrahi, D.; Sankella, S.; Gerard, R.D.; Horton, J.D.; Garg, A. Human 1-acylglycerol-3-phosphate O-acyltransferase isoforms 1 and 2: biochemical characterization and inability to rescue hepatic steatosis in Agpat2(-/-) gene lipodystrophic mice. *J. Biol. Chem.* 2011, 286, 37676-37691. doi: 10.1074/jbc.M111.250449

53. He, W.; Shi, Q.; Hu, X.; Yan, R. The membrane topology of RTN3 and its effect on binding of RTN3 to BACE1. *J. Biol. Chem.* 2007, 282, 29144-29151. doi: 10.1074/jbc.M704181200

54. Mayerhofer, P.U.; Bano-Polo, M.; Mingarro, I.; Johnson, A.E. Human peroxin PEX3 is co-translationally integrated into the ER and exits the ER in budding vesicles. *Traffic* 2016, 17, 117-130. doi: 10.1111/tra.12350

55. Jan, C.H.; Williams, C.C.; Weissman, J.S. Principles of cotranslational translocation revealed by proximity-specific ribosome profiling. Science 2014, 346, 6210. doi: 10.1126/science.1257521

56. Pfeiffer, N.V.; Dirndorfer, D.; Lang, S.; Resenberger, U.K.; Restelli, L.M.; Hemion, C.; Miesbauer, M.; Frank, S.; Neutzner, A.; Zimmermann, R.; Winklhofer, K.F.; Tatzelt, J. Structural features within the nascent chain regulate alternative targeting of secretory proteins to mitochondria. *EMBO J.* 2013, 32, 1036-1051. doi: 10.1038/embj.2013.46

57. Nuebel, E.; Morgan, J.T.; Fogarty, S.; Winter, J.M.; Lettlova, S.; Berg, J.A.; Chen, Y.-C.; Kidwell, C.U.; Maschek, J.A.; Clowers, K.J.; Argyriou, C.; Chen, L.; Wittig, I.; Cox, J.E.; Roh-Johnson, M.; Braverman, N.; Bonkowsky, J.; Gygi, S.P.; Rutter, J. The biochemical basis of mitochondrial dysfunction in Zellweger spectrum disorder. *EMBO rep.* 2021, 22, e51991. doi: 10.15252/embr.202051991

58. Chen, C.; Li, J.; Qin, X.; Wang, W. Peroxisomal membrane contact sites in mammalian cells. Front. *Cell Dev. Biol.* 2020, 8, 512. doi: 10.3389/celldev.2020.00512

59. Saito, K.M.; Chen, F.; Bard, S.; Chen, H.; Zhou, D.; Woodley, R.; Polischuk, R.; Schekman, R.; Malhotra, V. TANGO1 facilitates cargo loading at endoplasmic reticulum exit sites. *Cell* 2009, 136, 891-902. doi: 10.1016/j.cell.2008.12.025

60. Raote, I.; Ortega-Bellido, M.; Santos, A.J.M.; Foresti, O.; Zhang, C.; Garcia-Parajo, M.F.; Campelo, F.; Malhotra, V. TANGO1 builds a machine for collagen export by recruiting and spatially organizing COPII, theters and membranes. *eLIFE* 2018, 7, e32723. doi: 10.7554/eLIFE.32723

61. Raote, I.; Ernst, A.M.; Campelo, F.; Rothman, J.; Pincet, F.; Malhotra, V. TANGO1 membrane helices create a lipid diffusion barrier at curved membranes. *eLIFE* 2020, 9, e57822. doi: 10.7554/eLIFE.57822
Pataki, C.I.; Rodrigues, J.; Zhang, L.; Qian, J.; Efron, B.; hastie, T.; Elias, J.E.; Levitt, M.; Kopito, R.R. Proteomic analysis of monolayer-integrated proteins of lipid droplets identifies amphipathic interfacial α-helical membrane anchors. *Proc. Natl. Acad. Sci. USA* **2018**, *115*, E8172-E8180. doi: 10.1073/pnas.1807981115

Pinto, M.P.; Grou, C.P.; Fransen, M.; Fransen, M.; Sa-Miranda, C. The cytosolic domain of PEX3, a protein involved in the biogenesis of peroxisomes, binds membrane lipids. *Biochem. Biophys. Acta* **2009**, *1793*, 1669-1675. doi: 10.1016/j.bbamcr.2009.08.007

Poulos, A.; Christodoulou, J.; Chow, C.W.; Goldblatt, J.; Paton, B.C.; Orii, T.; Suzuki, Y.; Shimozawa, N.; Peroxisomal assembly defects: clinical, pathologic, and biochemical findings in two patients in a newly identified complementation group. *J. Pediatr.* **1995**, *127*, 596–599. doi: 10.1016/s0022-3476(95)70121-4

Rappsilber, J.; Mann, M.; Ishihama, Y. Protocol for micro-purification, enrichment, pre-fractionation and storage of peptides for proteomics using StageTips. *Nat. Protoc.* **2007**, *2* 1896-1906. doi: 10.1038/nprot.2007.261

Nagaraj, N.; Kulak, N.A.; Cox, J.; Neuhauser, N.; Mayr, K.; Hoerning, O.; Vorm, O.; Mann, M. System-wide perturbation analysis with nearly complete coverage of the yeast proteome by single-shot ultra HPLC runs on a bench top Orbitrap. *Mol. Cell. Proteom.* **2012**, *11*, M111.013722. doi: 10.1074/mcp.M111.013722

Cox, J.; Mann, M. MaxQuant enables high peptide identification rates, individualized p.p.b.-range mass accuracies and proteome-wide protein quantification. *Nat. Biotechnol.* **2008**, *26*, 1367-1372. doi: 10.1038/nbt.1511

Hyunsoo, K.; Golub, G.H.; Park, H. Missing value estimation for DNA microarray gene expression data: local least squares imputation. *Bioinform.* **2005**, *21*, 187-198. doi: 10.1039/bioinformatics/btk053