Congenital Sucrase-Isomaltase Deficiency
Identification of a Glutamine to Proline Substitution That Leads to a Transport Block of Sucrase-Isomaltase in a Pre-Golgi Compartment

Joke Ouwendijk,* Catharina E.C. Moolenaar,† Wilma J. Peters,* Cornelis P. Hollenberg,‡ Leo A. Ginsel,* Jack A.M. Fransen,* and Hassan Y. Naim§

*Department of Cell Biology and Histology, University of Nijmegen, 6500 HB Nijmegen, The Netherlands; and †Institute of Microbiology, Heinrich-Heine University of Düsseldorf, D-40225 Düsseldorf, Germany

Abstract

Congenital sucrase-isomaltase deficiency is an example of a disease in which mutant phenotypes generate transport-incompetent molecules. Here, we analyze at the molecular level a phenotype of congenital sucrase-isomaltase deficiency in which sucrase-isomaltase (SI) is not transported to the brush border membrane but accumulates as a mannose-rich precursor in the endoplasmic reticulum (ER), ER-Golgi intermediate compartment, and the cis-Golgi, where it is finally degraded. A 6-kb clone containing the full-length cDNA encoding SI was isolated from the patient’s intestinal tissue and from normal controls. Sequencing of the cDNA revealed a single mutation, A/C at nucleotide 3298 in the coding region of the sucrase subunit of the enzyme complex. The mutation leads to a substitution of the glutamine residue by a proline at amino acid 1098 (Q1098P). The Q1098P mutation lies in a region that is highly conserved between sucrase and isomaltase from different species and several other structurally and functionally related proteins.

This is the first report that characterizes a point mutation in the SI gene that is responsible for the transport incompetence of SI and for its retention between the ER and the Golgi. (J. Clin. Invest. 1996. 97:633-641.) Key words: ERGIC-53 • intestine • processing • quality control • retention

Introduction

The highly organized cascade of processing and modification events along the biosynthetic pathway of secretory and membrane proteins has led to the general concept of compartmentalization of the cell into structurally and functionally distinct organelles (1-4). Conformational modifications of membrane and secretory proteins commence already during their translo-

Address correspondence to Dr. Hassan Y. Naim, Institute of Microbiology, Heinrich-Heine University of Düsseldorf, Univeritätsstrasse 1, Geb. 26.12, D-40225 Düsseldorf, Germany. Phone: 211-811-3733/3586; FAX: 211-811-3733/5370; E-mail: hassan.naim@uni-duesseldorf.de

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Abbreviations used in this paper: AMV, avian myeloblastosis virus; A2P, aminopeptidase N; CSID, congenital sucrase-isomaltase deficiency; Endo F/GF; Endo N-acetylglucosaminidase F/glycoprotease F; Endo H; Endo N-acetylglucosaminidase H; ER, endoplasmic reticulum; ERGIC, ER-Golgi intermediate compartment; LPH, lactase-phlorizin hydrolase; SI, sucrase-isomaltase.

1. Abbreviations used in this paper: AMV, avian myeloblastosis virus; A2P, aminopeptidase N; CSID, congenital sucrase-isomaltase deficiency; Endo F/GF; Endo N-acetylglucosaminidase F/glycoprotease F; Endo H; Endo N-acetylglucosaminidase H; ER, endoplasmic reticulum; ERGIC, ER-Golgi intermediate compartment; LPH, lactase-phlorizin hydrolase; SI, sucrase-isomaltase.
apical membrane and cleaved in situ by luminal pancreatic proteases to its two active subunits, sucrase and isomaltase (22, 23).

The molecular basis of CSID is still not elucidated. However, it has been proposed that different molecular defects or mutations in the SI gene are responsible for CSID (24). This hypothesis has been strongly supported by the analysis of several cases of CSID, which has led to the identification of five different phenotypes of SI (14, 17). Phenotypes I and II, for instance, are characterized by an intracellular accumulation of mannose-rich SI in the ER and the Golgi, respectively. In phenotype III an enzymatically inactive, but transport-competent, SI is expressed. Phenotype IV expresses a partially folded, mannose-rich SI molecule that is missorted to the basolateral membrane. Finally, phenotype V reveals an SI species that undergoes intracellular degradation leaving behind the isomaltase subunit that is correctly targeted to the brush border membrane. In this paper we analyzed a biopsy specimen of CSID phenotype II and demonstrated that SI is located in the ER, the intermediate compartment and the cis-Golgi. We show for the first time that a single point mutation is responsible for the impaired transport behavior of SI.

Methods

Antibodies. Four epitope-specific mAbs directed against sucrase, isomaltase, or SI were studied (25). These antibodies were products of the following hybridomas: HBB 1/691, HBB 2/614, HBB 2/219, and HBB 3/705. mAb anti-human lactase-phlorizin hydrolase (mAb anti-LPH) and mAb anti-human aminopeptidase N (mAb anti-ApN) were products of hybridomas HBB 1/909 and HBB 3/153, respectively (25). mAb anti-ER-Golgi intermediate compartment-53 (anti-ERGIC-53) was a product of hybridoma G1/93 (26). All mAbs were generously provided by Dr. H. P. Hauri (Biocenter, Basel, Switzerland) and Dr. E. E. Sterchi (University of Bern, Bern, Switzerland).

Cell lines. Caco-2 cells (ATCC HTB-37) were cultured in Dulbecco's modified Eagle's medium (DME) supplemented with 20% fetal calf serum, 2 mM L-glutamine, 100 IU/ml penicillin, 100 mg/ml streptomycin, and 1.25 mg/ml amphotericin B. Cells were grown at 37°C in a humidified 5% CO₂ incubator.

Processing of biopsy samples. The patient with CSID had a lifelong history of abdominal pain, cramps, and osmotic diarrhea upon ingestion of sucrose. CSID was assessed by the breath hydrogen test following an overnight fast with 50 g sucrose. CSID was confirmed by enzyme activity measurements of sucrase and isomaltase in intestinal biopsy samples from CSID and controls. Total RNA isolation and cDNA synthesis were performed as described by Moolenaar et al. (31). Total RNA was isolated from postconfluent Caco-2 cells using the guanidinium isothiocyanate method (32). Randomly primed cDNA was synthesized from 10 μg total RNA by avian myeloblastosis virus (AMV) reverse transcriptase (AMV-RT; Boehringer Mannheim Corp., Indianapolis, IN). mRNA was incubated for 60 min at 37°C in 50 mM Tris/HCl (pH 8.3), 40 mM KCl, 6 mM MgCl₂, 2.5 mM DTT, 1 mM dNTP (Boehringer Mannheim Corp.) in the presence of 200 ng random primers and 25 U AMV-RT. The negative control underwent the same procedure, only AMV-RT was excluded from the reaction mixture. For each PCR reaction 1/10 (2 μl) of the reaction mixture was used as a template. PCR reactions were performed as described by Innis et al. (33) with minor modifications. Briefly, cDNA templates were amplified in a total volume of 100 μl containing, 10 mM Tris/HCl (pH 8.3), 25 mM KCl, 1.5 mM MgCl₂, 140 mM of each dNTP, 700 nM of primer, and 2 U of Taq polymerase (Boehringer Mannheim Corp.). Double-stranded DNA was dissociated at 94°C for 45 s, the annealing step was at 55°C for 1 min, and the extension step was at 70°C for 1.5 min. The reactions were subjected to 30 cycles of amplification. Some of the primers used in the PCR reactions were designed based on the published sequences of the rabbit and human SI cDNA (34, 35). Table I shows the position of these primers within the SI cDNA. The products of two independent PCR reactions were directly or after cloning into pGEM4Z sequenced with a DNA sequencing kit according to the instructions of the manufacturer (Sequenase V.2.0; United States Biochemical Corp., Cleveland, OH).

The various DNA fragments of wild-type SI cDNA were ligated together and cloned into the expression vector pCB7 (36) to generate the plasmid pHSI encoding wild-type pro-SI. Sequencing of the DNA fragments obtained by PCR from the CSID tissue revealed a single mutation A/C at nucleotide 3298. This mutation was found in the PCR product encompassing nucleotides 2967–4201. A 619-bp (Nco–Apal) fragment containing this mutation was purified and exchanged with its counterpart in

Table I. Oligonucleotides Used to Synthesize Various cDNAs by PCR

<table>
<thead>
<tr>
<th>DNA fragment (bp)*</th>
<th>Primers (5'–3' direction)</th>
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<tr>
<td>29–695</td>
<td>CCGGTTACCAGCCTTATCCAAGTCTG</td>
</tr>
<tr>
<td>652–1366</td>
<td>GCATGGTGCTCCCTATGGAAC</td>
</tr>
<tr>
<td>1299–2066</td>
<td>ATAGGTCGACGGCAATGGAAG</td>
</tr>
<tr>
<td>1952–3139</td>
<td>GATGAACTTGGGGGATTTCA</td>
</tr>
<tr>
<td>2967–4201</td>
<td>GTCTGGGATTACCCACAT</td>
</tr>
<tr>
<td>4071–4923</td>
<td>TCCAGACGTCTGAGCTTTC</td>
</tr>
<tr>
<td>4876–5555</td>
<td>TATCCAGAAGTTCTTTATGG</td>
</tr>
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*bp numbering according to Green et al. (35).
the wild-type phSI plasmid. The generated plasmid was denoted phSIa/c.

Transfection and metabolic labeling of COS-1 cells. COS-1 cells were either transfected by the DEAE-dextran method as described by Naim et al. (29) or via electroporation (Bio-Rad Laboratories, Richmond, CA): a 0.4-cm cuvette containing 200 µl ice-cold PBS with 1.5 million cells and 10 µg DNA was exposed to 0.3 kV 300 µF. Transiently transfected COS-1 cells were metabolically labeled with 80 µCi [35S]methionine (ICN Biomedicals, Inc.) as described by Naim et al. (29). After the labeling period, the cells were washed and stored at −70°C until use.

Immunoprecipitation. Homogenates of biosynthetically labeled biopsy specimens were lysed on ice by addition of Triton X-100 and sodium deoxycholate to final concentrations of 0.5% each (28). Bio synthetically labeled transfected COS-1 cells were lysed at 4°C for 1 h in lysis buffer (25 mM Tris-HCl, pH 8.0, 50 mM NaCl, 0.5% Triton X-100, 0.5% sodium deoxycholate, and a mixture of protease inhibitors containing 1 mM PMSF, 1 µg/ml pepstatin, 5 µg/ml leupeptin, 17.4 µg/ml benzamidine, and 1 µg/ml apro tin). Usually 1 ml ice-cold lysis buffer was used for each 100-mm culture dish (~2–4 × 10^6 cells). Detergent extracts of cells or biopsy samples were centrifuged for 1 h at 100,000 g at 4°C and the supernatants were immunoprecipitated as described by Naim et al. (17, 22, 28, 29). Four epitope-specific mAbs directed against sucrase, isomaltase, or SI were used (25). These antibodies were used in the form of ascites. Usually, 5–10 µl ascites was used for each immunoprecipitation. For epitope mapping studies with four mAbs directed against SI the extracts of biopsy samples were divided into equal aliquots and each aliquot was immunoprecipitated with a different antibody.

Immunofluorescence. Cellular localization of expressed proteins in COS-1 cells was studied with cells grown on coverslips. Cells were fixed with 3% paraformaldehyde and immunolabeling was carried out after 0.1% Triton X-100 permeabilization, using mAb anti-SI (HSB1/614) (1:1000) and mAb anti-ERGIC-53 (G1/93) (1:100) as the primary antibodies. The secondary antibodies used FITC-conjugated goat anti-mouse or anti-rabbit IgG (1:100), Texas red-conjugated goat anti-mouse or anti-rabbit IgG (1:200) (all were from Boehringer Mannheim Corp.). Cell surface localization of proteins was assessed in transfected cells that were incubated on ice with the primary antibody before fixation and without incubation with Triton X-100. The cells were visualized with a confocal laser microscope using a double channel for FITC and Texas red and/or with a routine fluorescence microscope.

Immunoelectron microscopy. Biopsy samples were cut into small pieces and fixed in a mixture of 2% formaldehyde and 0.1% glutaraldehyde for 1 h at room temperature. They were then stored in 2% formaldehyde until use. Cryosectioning and immunolabeling with anti-SI, anti-ERGIC-53, and protein A-gold were performed as described previously (26, 37).

Other procedures. SDS-PAGE was performed according to the method of Laemmli (38). High molecular weight markers from Sigma Immunohemicals were used as a reference. Treatment of immunoprecipitates with Endo-N-acetylglucosaminidase H (Endo H), Endo-N-acetylglucosaminidase F (Endo F), also known as PNGase F) (both from Boehringer Mannheim Corp.) was done as described before (28).

Results

Initial assessment of CSID in the patient was achieved by the breath hydrogen test. Subsequently a biopsy specimen was taken from the upper jejunum and enzymatic activity measurements of the disaccharidasises, SI, and LPH were performed. Here, no activity of sucrase or isomaltase could be detected, while the activity of LPH was in the normal range of the control tissue (16–32 IU/mg protein). Likewise, normal activity of another control brush border enzyme, ApN, was revealed in the patient’s biopsy sample. Therefore, the results pointed to an enzymatic defect that is restricted to SI. Previous analyses of several cases of SI deficiency have demonstrated that SI is synthesized as in normal controls, but is not correctly processed or transported along the secretory pathway (14, 17). We therefore set out to analyze the biosynthesis and processing of SI in the patient’s tissue. Metabolic labeling of biopsy samples and immunoprecipitation of the detergent extracts with four different epitope-specific monoclonal anti-SI antibodies revealed after 5 min of labeling an Endo H–sensitive 210-kD polypeptide that corresponds to mannosine-rich pro-SI (Fig. 1, lanes 1–3). This polypeptide remained the predominantly labeled species after 4 and 18 h of labeling (Fig. 1, lanes 4–9). In the control biopsy specimen, by contrast, the Endo H–sensitive mannosine-rich pro-SI (Fig. 1, lanes 10–12) was processed to a complex glycosylated Endo H–resistant form after 4 h of labeling (Fig. 1, lanes 13–15). When comparing the metabolic labeling intensity of the patient’s material after 4 and 18 h of labeling, no increase is observed as compared with the control (Fig. 1, lanes 4–6 and 7–9 vs. lanes 10–12 and 13–15). From these data we conclude that no processing of the patient’s SI in the Golgi apparatus has occurred and that the protein is finally degraded.

To determine whether the impaired processing is the consequence of general cellular defect or is restricted to SI, the transport of other brush border proteins was investigated. ApN and LPH were synthesized and processed in the patient’s intestinal cells in a fashion similar to control cells (Fig. 2, lanes 1–8). Thus, within 1 h of chase ApN was detected in the patient’s biopsy sample as an Endo H–sensitive mannosine-rich 130-kD polypeptide, which was converted into an Endo H-resistant 160-kD polypeptide after 20 h of chase (Fig. 2, lanes 3 and 4). Similarly no changes in the biosynthetic pattern of LPH could be observed. Here, LPH was synthesized as a 215-kD mannosine-rich species that was cleaved intracellularly to the mature 160-kD polypeptide (Fig. 2, lanes 7 and 8). Therefore, the failure of mannosine-rich SI to mature to an Endo H resistant species is indeed restricted to this molecule and is not the result of a general cellular defect since two control proteins, ApN and LPH, were processed to their mature forms and transported normally in the patient’s tissue.

Figure 1. Molecular forms of SI in CSID and in normal control. Biopsy samples from a patient with CSID [lanes 1–9] and normal control [lanes 10–13] were biosynthetically labeled for the indicated times with [35S]methionine. The specimens were homogenized, solubilized, and immunoprecipitated with monoclonal anti-SI antibodies. The immunoprecipitates were subjected to SDS-PAGE on 5% slab gels without treatment (lanes 1, 4, 7, 10, and 13) or after Endo H (lanes 2, 5, 8, 11, and 14) or Endo F (lanes 3, 6, 9, 12, and 15) treatment. Gels were analyzed by fluorography.

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Since SI is synthesized at almost normal levels in the patient's tissue but does not mature, we wanted to determine the subcellular localization of SI in the patient's epithelial cells. Here, ultrathin cryosections of the patient's tissue were incubated with monoclonal anti-SI antibodies followed by protein A–gold. The subcellular localization was finally assessed by electron microscopy (Fig. 3). While most of the labeling in the control intestinal cells was confined to the brush border membrane, no labeling of the brush border membrane in the patient's tissue could be detected (Fig. 3, A and B). Instead, intense labeling of the ER and Golgi cisternae was observed in the patient's tissue (Fig. 3 D), while the Golgi apparatus and the ER of the control were weakly labeled (Fig. 3 C). The latter case reflects the normal steady state situation in biopsy samples (14). Finally, both specimens, the patient's as well as the control's, did not show any labeling of the basolateral membrane.

The SI labeling in the patient's cells was visible on one side of the Golgi (Fig. 3 D, arrows) and corresponds most likely to the cis-Golgi and ERGIC. This is based on the labeling pattern obtained when biopsy samples are labeled with antibodies against ERGIC-53, a marker of the cis-Golgi as well as ERGIC (Fig. 4). It should be noted that in the patient's biopsy material more label was observed for ERGIC-53 than in the control biopsies (compare Fig. 4, A and B). In addition, ERGIC-53 was also observed in the cisternae on one side of the Golgi apparatus in contrast to control biopsies where only a tubulo-vascular structure on one side of the Golgi was labeled (Fig. 4 A; reference 26). Based on morphological criteria we conclude that the labeling of ERGIC-53 in the patient's biopsy coincides with the label of SI (compare Figs. 3 D and 4 B), although a direct double labeling of SI and ERGIC-53 proteins could not be performed. The antibodies used to detect these proteins are both mAbs and these are difficult to combine in immunogold double staining experiments. Control experiments using antibodies against the brush border glycoproteins LPH and ApN demonstrated localization of these proteins in the apical membrane (not shown).

Altogether, the data demonstrate that in the patient's biopsy the SI molecule has egressed the ER and is arrested in an ERGIC-53-positive, cis-Golgi compartment where it is finally degraded. The phenotype therefore resembles phenotype II of CSID described previously (17).

It is likely that the transport block of SI in the ERGIC, cis-Golgi, and ER as well as the persistence as a mannose-rich polypeptide are the consequence of structural alterations in the SI molecule. To examine this possibility, we performed epitope mapping using four different epitope-specific monoclonal anti-SI antibodies, HBB 2/614, HBB 2/219, HBB 1/691, and HBB 3/705. Except for mAb HBB1/691 all other antibodies were able to recognize epitopes on mutant SI (not shown). The results imply that the epitope of HBB 1/619 did not mature in the patient's SI and was altered. Drastic alterations in the protein folding of SI did not presumably take place, since three out of four antibodies have recognized the mutant SI.

To analyze the molecular defects in this phenotype, we cloned the full-length cDNA encoding wild-type SI from Caco-2 cells and from the patient's biopsy sample. Construction of full-length cDNA encoding wild-type SI was performed by combining two approaches. The first used conventional screening of a λgt-11 expression library made from the colon carcinoma cell line, Caco-2. Here, a clone comprising 1486 bp from nucleotides 550 to 2036 (base pair numbering according to Green et al. [35]) and covering almost 50% of the isomaltase subunit was isolated. This clone was denoted IM21.1 (Lacey, S.W., unpublished observations). In the second approach we made use of the published sequences of rabbit and human SI (20, 34). Many oligonucleotides were designed and used to amplify SI DNA fragments using RT-PCR on RNA extracted from Caco-2 cells or biopsy samples. These fragments and the IM21.1 clone were ligated together to generate the full-length pro-SI cDNA. Sequencing of this clone revealed two nucleotide differences versus the published sequence (34). These differences are not due to PCR artifacts, as assessed by sequencing of PCR products of two independent PCR reactions. One difference is A/G at nucleotide 4001 which represents a silent mutation. The second variation, a C/G at nucleotide 3612, leads to a glutamic acid instead of glutamine at amino acid 1203 in the sucrase subunit. Comparison with the amino acid sequences of rat and rabbit and our patient's (see below) sucrase subunits revealed also a glutamic acid residue at this particular position. Moreover, the human and rabbit isomaltase also contained a glutamic acid in the corresponding homologous stretch. Therefore, it is likely that the glutamine/glutamic acid exchange represents a polymorphism in the published sequence. To test the full-length cDNA, it was cloned into the pCB7 expression vector and transiently expressed in COS-1 cells. Metabolic labeling of these cells showed that SI is processed as in normal control samples (Fig. 5, lanes 1–3, compare with Fig. 1, lanes 10–15). Also, when expressed in COS-1 cells, cell surface expression could be observed (Fig. 6, A and B).

To isolate a full-length SI cDNA encoding SI in the CSID tissue, a similar strategy was followed as for wild-type SI. The PCR products derived from two independent reactions were cloned and sequenced. The sequence analysis revealed one alteration relative to the SI cDNA cloned from Caco-2 cells. This alteration, A/C at nucleotide 3298, was found in the sequence encoding the S of the SI complex and resulted in a substitution of the glutamine 1098 by a proline (Q1098P). The mutation was confirmed by direct sequencing of several independent PCR products. The wild-type sequence could never

![Figure 2. Biosynthesis of the control brush border glycoproteins in CSID and normal control. After the immunoprecipitations with mAb anti-SI in Fig. 1, the supernatants depleted from SI were immunoprecipitated sequentially with mAb anti-ApN (lanes 1–4) or mAb anti-LPH (lanes 5–8). The immunoprecipitates were analyzed by SDS-PAGE on 5% slab gels without treatment (lanes 1, 3, 5, and 7) or after Endo H (lanes 2, 4, 6, and 8) treatment.](image-url)
be identified, suggesting that both alleles of the SI gene contain this mutation or, more likely, that one allele is not expressed.

The patient's cDNA was also cloned into the expression vector pCB7 and transiently expressed into COS-1 cells. Again, metabolically labeled COS-1 cells showed that mutant cDNA is processed as in the patient's biopsy (Fig. 5, lanes 4-6, compare with Fig. 1, lanes 1-9). In addition, localization studies showed that the patient's SI is not expressed at the cell surface of COS-1 cells, but instead accumulates intracellularly in the ER and ERGIC, as shown by immuno double-labeling on transiently transfected cells (Fig. 6, C and D).
Figure 4. Immunocytochemical localization of ERGIC-53 in CSID and normal control. Localization of ERGIC-53 was performed on ultrathin cryosections from a control biopsy (A) and from the patient's biopsy (B) using antibody Gl/93, followed by rabbit anti-mouse and protein A-10 nm gold staining. (A) In the control biopsy vesicular structures on the cis-side of the Golgi apparatus are stained; (B) in contrast, in the patient's biopsy the labeling is not only stronger, but also additional staining over cisternae on one side of the Golgi is observed. Morphologically the label coincides with that observed for SI (compare with Fig. 3 D). Bar, 0.25 μm.

<table>
<thead>
<tr>
<th>SI</th>
<th>Wt</th>
<th>Q1098P</th>
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<tbody>
<tr>
<td>Endo H</td>
<td>-</td>
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<td>Endo F</td>
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**Figure 5.** Expression of wild-type and mutant SI cDNA in COS-1 cells. COS-1 cells were transfected with phSI (lanes 1–3) or phSIa/c (lanes 4–6) plasmids and biosynthetically labeled with [35S]methionine for 6 h. The cell extracts were immunoprecipitated with mAb anti-SI. The immunoprecipitates were analyzed by SDS-PAGE on 5% gels without treatment (lanes 1 and 4) or after Endo H (lanes 2 and 5) or Endo F treatment (lanes 3 and 6).

Upon comparison, lysosomal α-glucosidase, *Schwanniomyces occidentalis* glucoamylase, sucrase, and isomaltase from various species share striking similarities in their amino acid sequences. This has led to the hypothesis that these molecules are derived from a common ancestral gene (39). Sequence comparison revealed that the mutation Q1098P in the S subunit is located in a stretch of 10 amino acids that is highly homologous among sucrase (amino acids 1093–1116), isomaltase (amino acids 219–243), lysosomal α-glucosidase (amino acids 239–262) and *Sch. occidentalis* glucoamylase (amino acids 196–220) (Fig. 7).

**Discussion**

CSID has been used to characterize various steps in the biosynthesis, transport, and sorting of SI as a model for cell surface membrane proteins. Several different phenotypes of CSID have been meanwhile described (14–17). In all of these CSID phenotypes, the transport incompetence, aberrant enzymatic function, or missorting to the basolateral membrane is exclusively restricted to the SI molecule, ruling out a possible general cellular defect.

While analyses of the current and previous phenotypes at the subcellular and protein levels lent strong support to the hypothesis that a point mutation in the cDNA of SI is responsible for the generation of CSID, the present data provide the first definite evidence that this is in fact the case. Here, we could characterize a point mutation in the region encoding the sucrase subunit of the SI complex which converts a glutamine to a proline (Q1098P).

Therefore, the CSID malabsorption is another example of a clinical disorder or disease that ensues by single amino acid
mutation in the polypeptide chain leading ultimately to dramatic alterations in the structural features, biosynthesis, posttranslational processing, transport, and function of cell surface and secretory proteins. While the CSID phenotype usually ends up in mild malabsorption of sugars and starch, the symptoms of which are readily identifiable, other disorders, such as cystic fibrosis (40), collagen-based diseases, and familial hypercholesterolemia (41), just to name a few, are more severe (for review see reference 13).

A rather unusual mutant form of SI is generated in CSID phenotype II. Here, SI exits the ER and is transported to the medial and trans-Golgi where it undergoes degradation (17). To our knowledge, no mammalian cell surface membrane glycoprotein has been so far identified that has efficiently egressed the ER and arrived in the Golgi cisternae where its transport is blocked. This strongly suggests that there are other criteria, in addition to the acquisition of correct folding and quaternary structure in the ER, that may modulate and control protein transport within the Golgi and to the cell surface.

The phenotype of SI described in this paper is very much similar to phenotype II (17). Analyses of the structural features of SI in this phenotype by epitope mapping and biosynthetic labeling reveal that incomplete initial folding of the SI precursor protein is responsible for the observed inefficient transport of the molecule along the secretory pathway. An effect of the oligomeric state of SI could be excluded since human SI does not dimerize and exits the ER efficiently as a monomeric molecule (8).

The most interesting aspect of phenotype II is the ability of SI to egress the ER to the Golgi, while still in an incompletely or partially folded state. The subsequent arrest and final degradation of SI in a cis-Golgi compartment or in the Golgi is unusual and, except for SI, has not been demonstrated for an endogenous protein destined for the cell surface. This is surprising since membrane and secretory proteins are believed to acquire transport competence in the ER and leave this organelle en route to their final destination.

It is generally accepted that a quality control mechanism

Figure 6. Immunofluorescence of wild-type and mutant SI cDNA in COS-1 cells. COS-1 cells were transfected with control SI cDNA (A and B) and Q1098P mutant SI cDNA (C and D). (A) Control SI can be observed in the rough ER, Golgi, and cell surface of COS-cells; (B) cell surface staining of control SI, see Methods for details of the procedure. (C and D) Immunodouble-labeling of SI and ERGIC-53 in a COS cell transfected with mutant SI-cDNA and observed using a confocal scanning laser microscope equipped with a Kr/Ar laser; (C) immunostaining with HBB 2/614 anti-SI antibody in the ER and an ERGIC-53-positive compartment, note the absence of surface label (compare with A); (D) immunostaining with G1/93 anti-ERGIC-53 antibody, note the structure (arrow) that is labeled both for SI (C) and ERGIC-53 (D). Bar, 15 μm.
While these alterations could be tolerated by the quality control mechanism while the entire isomaltase subunit remains unaffected. A mutation in the sucrase subunit leads to alterations in its structure and it is reasonable to assume that the presence of the regions containing the active catalytic centers of these proteins have almost identical amino acid sequences (34). If all proteins have almost identical amino acid sequences (34). If the mutation Q1098P has created a signal or conformational change in the S subunit, but rather alters a subdomain of the ER, whereby the folding of some domains may be more essential or critical than others in the context of the overall conformation. SI is, sorted in the trans-Golgi and maturation in the siss-Golgi compartment, where it is finally degraded by an unknown quality control mechanism operating at a level beyond the ER.

Figure 7. Amino acid homology around the mutated region in the patient’s SI. The amino acid sequences of normal human sucrase (32), rabbit sucrase (20), rat sucrase (45), human isomaltase (32), rabbit isomaltase (20), human lysosomal α-glucosidase (43), and Sch. ocellatilis glucoamylase (39) are aligned. The numbering and arrow indicate the position in this highly conserved region where in the patient’s SI the Q1098P mutation is located.

Acknowledgments

We conclude that the simplest and most straightforward explanation for the phenotype described in this paper is that the mutation Q1098P has created a signal or conformational change in the sucrase subunit which retains the enzyme complex in a cis-Golgi compartment, where it is finally degraded by an unknown quality control mechanism operating at a level beyond the ER.

Acknowledgments

The first two authors on this paper have contributed equally to the work described.

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