Small-intestinal dysfunction accompanies the complex endocrinopathy of human proprotein convertase 1 deficiency


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We have previously described the only reported case of human proprotein convertase 1 (PC1) deficiency, in a female (Subject A) with obesity, hypogonadism, hypoadrenalism, and reactive hypoglycemia. We now report the second case of human PC1 deficiency (Subject B), also due to compound heterozygosity for novel missense and nonsense mutations. While both subjects shared the phenotypes of obesity, hypoadrenalism, reactive hypoglycemia, and elevated circulating levels of certain prohormones, the clinical presentation of Subject B was dominated by severe refractory neonatal diarrhea, malabsorptive in type. Subsequent investigation of Subject A revealed marked small-intestinal absorptive dysfunction, which was not previously clinically suspected. We postulate that PC1, presumably in the enteroendocrine cells, is essential for the normal absorptive function of the human small intestine. The differences in the nature and severity of presentation between the two cases cannot readily be explained on the basis of allelic heterogeneity, as the nonsense and missense mutations from both subjects had comparably severe effects on the catalytic activity of PC1. Despite Subject A’s negligible PC1 activity, some mature ACTH and glucagon-like peptide 17-36amide were detectable in her plasma, suggesting that the production of these hormones, at least in humans, does not have an absolute dependence on PC1. The presence of severe obesity and the absence of growth retardation in both subjects contrast markedly with the phenotype of mice lacking PC1 and suggest that the precise physiological repertoire of this enzyme may vary between mammalian species.

to have more distinct functions, evidenced by tissue-specific expression. For example, corticotropes, which express only PC1, cleave proopiomelanocortin (POMC) to adrenocorticotropic hormone (ACTH), while melanotropes, which additionally express PC2, form α-melanocyte stimulating hormone (αMSH) and β endorphin (4). Similarly, proglucagon is processed to glucagon-like peptides 1 and 2 (GLP-1 and GLP-2) in intestinal L cells, which express PC1, and to glucagon in pancreatic islet α cells, which express PC2 (see Figure 3a) (5, 6).

The functions of mammalian PC1 and PC2 have been further clarified through the generation, by homologous recombination, of mice deficient in one or the other of these enzymes. PC2-null animals have fasting hypoglycemia due to a lack of production of mature glucagon from the pancreatic α cell, and PC1-null mice, while sharing some elements of the phenotype reported in humans, uniquely have severely reduced linear growth attributed to failure of growth hormone releasing hormone production (7–9). It is as yet unclear whether these enzymes have an identical range of actions in humans and to what extent dysfunction of these enzymes can contribute to human metabolic and endocrine disease.

We previously reported a woman (Subject A) with marked childhood obesity, hypogonadotropic hypogonadism, postprandial hypoglycemia, hypocortisolism, and evidence of impaired processing of ACTH (9, 10). She is a compound heterozygote for PC1 mutations: Gly593Arg (GenBank X64810), which causes failure of maturation of the inactive propeptide form of PC1 (pro-PC1) and its retention in the ER, and A→C+4 in the donor splice site C+4 in the donor splice site of intron 5, resulting in exon skipping and a frameshift and premature stop codon in the catalytic domain. We now describe the second case of congenital PC1 deficiency, in a patient (Subject B) in whom the presenting features were dominated by severe small-intestinal dysfunction. Re-evaluation of Subject A showed that she too had small-intestinal dysfunction, but of a lesser degree, which prompted us to examine whether dissimilarities between the subjects were due to differences in function of the mutant enzymes. Finally, in Subject A, we further evaluated circulating prohormones and products reportedly associated with PC1 expression and examined the role of this enzyme in prohormone processing events in humans.

Methods

Genetic studies in Subject B and construction of vectors expressing mutant PC1

The 14 exons of PC1 and their intronic boundaries in Subject B were bidirectionally sequenced (BigDye Terminator cycle sequencing; Roche Molecular Biochemicals, Mannheim, Germany) using PCR amplified genomic DNA (10). Construction of the Gly593Arg mutant PC1 cDNA has been described previously (10). A FLAG epitope tag (Asp-Tyr-Lys-Asp-Asp-Asp-Asp-Lys) was introduced between the propeptide and the catalytic domain by insertion mutagenesis using Altered Sites II in vitro Mutagenesis Systems (Promega Corp., Madison, Wisconsin, USA). PC1-Ala213del and all other mutations were created using a QuikChange site-directed mutagenesis kit (Stratagene, La Jolla, California, USA). So that the propeptide could be labeled metabolically with 35S-methionine, Ile86 was replaced by a Met in the co-immunoprecipitation study. WT PC1 and splice-site variant PC1 (PC1-ssv) minigenes were constructed using an approximately 7-kb genomic fragment of PC1 spanning exons 4–6, which was amplified by PCR and cloned in pGEM-T Easy (Promega Corp.). A 4-kb KpnI-KpnI fragment from intron 5 was deleted by digestion before insertion into the above cDNA. This construct and one with the splice-site mutation inserted were cloned into pcDNA3 (Invitrogen Life Technologies, Groningen, The Netherlands) and checked by sequencing.

Cell lines, DNA transfection, and protein analysis

Medium, serum, and supplements were obtained from Invitrogen Life Technologies, and cells were cultured as previously described (11). CHO-K1 and HEK-293T cells were transfected with FuGENE (Roche Molecular Biochemicals), AtT-20 cells with Lipofectamine (Invitrogen Life Technologies), and αTC1-6, βTC3, and Neuro2A cells with Lipofectamine 2000 (Invitrogen Life Technologies). Cell labeling, lysis, immunoprecipitation, SDS-PAGE, and Western blotting were performed as previously described (11) using anti-PC1 antibody (Alexis Corp., Läufelfingen, Switzerland) or anti-FLAG M2 (Sigma-Aldrich, Bornem, Belgium).

Assay of PC1 activity

Recombinant PC1 was immunopurified from cells and medium as previously described (11), except for the lysis buffer (0.5% Triton X-100, 10 μM E-64, 1 μM leupeptin, 10 μM pepstatin, and 100 μM tosyl phenylalanyl chloromethyl ketone; all from Sigma-Aldrich). Recombinant PC1 was selectively detected in cells expressing endogenous PC1 by means of the FLAG epitope tag. To normalize the quantity of enzyme used in activity assays, PC1 expression was determined by overnight labeling of cells to steady-state levels (11) followed by SDS-PAGE analysis of 10% of the immunoprecipitate with measurement of band intensities using Image Station 440 (Eastman Kodak Co., New Haven, Connecticut, USA). The enzymatic activity of immunoprecipitates was assayed using fluorogenic substrate p-Glu-Arg-Thr-Arg-Arg-amino methylcoumarin (Peptides International Inc., Louisville, Kentucky, USA) at 37°C (12).

Analysis of circulating prohormones and their products

Proglucagon. Plasma from Subject A was collected during fasting and at 20-minute intervals after the start of a test meal (4MJ: 52% carbohydrate, 41% fat, 7% protein). Proglucagon-derived peptides were measured by RIA.
using antisera specific to free C-terminal glucagon, mid-sequence glucagon, and amidated C-terminal GLP-1 (5). The results of these assays were consolidated by further RIAs using antisera specific to mid-sequence glucagon (antiserum 4304), free C-terminal glucagon (antiserum 4305), mid-sequence GLP-1 (antiserum 2135), amidated C-terminal GLP-1 (antiserum 89390), and free N-terminal GLP-2 (antiserum 92160) (13). To identify the proglucagon products contributing to the measured immunoreactivities, plasma from Subject A and controls was subjected to size-exclusion gel chromatography, as previously described, prior to RIA of eluted peptides using the above assays (13, 14). In addition, eluted N-terminal GLP-17-36 and mid-sequence GLP-2 immunoreactivities were measured by RIA (antisera 93242 and RAS 7167, respectively; Peninsula Laboratories Europe LTD, St. Helens, United Kingdom). GLP-1 forms in plasma were also identified using reverse-phase HPLC with RIA of eluted amidated C-terminal GLP-1 (5). In both systems, peaks were defined by migration relative to standards and by the RIA specificity.

Progastrin. Progastrin and gastrin forms were measured in the plasma collected for proglucagon analysis, by RIA using antisera to C-terminal progastrin uncleaved at Arg94,95 (antiserum L289), amidated C-terminal G17 and G34 gastrins (antiserum L2), intact G17 (antiserum L6), and free N-terminal G17 (antiserum 1295) (15–17). Gastrins were also measured in controls (five male and five female, mean ages 39 and 34 years, respectively) using antisera L289 and L2.

POMC. To study the diurnal and stimulated secretion of POMC, ACTH, and cortisol, we sampled peripheral venous blood throughout a 24-hour period and 20, 60, and 120 minutes after i.v. administration of 100 µg corticotropin-releasing hormone (CRH). Cortisol was measured by radioimmunometric binding assay using transcortin (18), and ACTH was measured by immunoradiometric assay (IRMA), (ELSA-ACTH; CIS bio international, Gif-sur-Yvette, France). POMC cross-reactivity was such that 7,300 U/ml gave a signal corresponding to less than 6 pg/ml ACTH. The plasma POMC IRMA has been described previously (19). One unit of POMC corresponded to 1 pg of POMC protein, and the detection limit was 60 U/ml. Twenty minutes after administration of CRH, corticotropin-like intermediate lobe peptide (CLIP) and ACTH were identified in plasma by extraction on a C18 Sep-Pak cartridge (Waters Associates, Milford, Massachusetts, USA) followed by reverse-phase HPLC on a 0.46 cm × 25 cm C8 column (Aqua- pore RP 300; Brownlee Laboratory, Applied Biosystems, Foster City, California, USA) (20). Fractions were lyophilized and reconstituted in plasma buffer or ACTH C-terminal RIA, in which CLIP and phosphorylated and nonphosphorylated ACTH showed equal immunoreactivity. Forms of β endorphin, β lipotropin, and POMC in plasma were identified using size-exclusion gel chromatography prior to β endorphin RIA (Peninsula Laboratories Inc., Belmont, California, USA) and POMC IRMA (19). Sephadex G-50 Superfine (Pharmacia, Amersham Biosciences UK Ltd., Little Chalfont, Buckinghamshire, United Kingdom) in a 0.9 cm × 60 cm column was equilibrated and developed in 1% acetic acid with 0.1% BSA. The column was loaded with 0.8 ml of plasma and eluted at 2.5 ml/h. Dinitrophenylalanine and 125I-labeled IgG were added to the plasma sample as markers of the total and void volumes respectively, in order to calculate fractional elution volume. One-milliliter fractions were lyophilized and reconstituted in assay buffer prior to immunoassay.

Subject B’s plasma was examined by direct immunoassay of cortisol, ACTH, and precursors (POMC and proACTH) in blood collected at 0900 hours (21).

Proinsulin. Insulin forms in Subject B’s plasma were assayed by direct immunometric assay, and those in Subject A and her teenage children were analyzed by reverse-phase HPLC with immunoassay of eluted peptides as previously described (9).

Procalcitonin. Serum procalcitonin was assayed in Subject A in triplicate using an immunometric assay (Brahms Diagnostica, Berlin, Germany) with antisera to C-terminal procalcitonin and the centrally located calcitonin sequence (22). This assay cross-reacted with conjoined calcitonin and C-terminal peptide. Procalcitonin was also measured by N-terminal RIA, in which free N-terminal fragment (aminoprocalcitonin) cross-reacts (22, 23).

Prorenin. We sampled plasma before the subjects rose in the morning and after they had ambulated for an hour; then we measured active renin in the plasma by IRMA (24). In the same samples, total renin (prorenin plus renin) was determined as the active renin level after trypsinization (25).

Results

Clinical phenotype of Subject B

Subject B was a female infant with a normal 46XX karyotype, the first child of non-consanguineous Caucasian parents, who was born after 37 weeks of uncomplicated gestation. Diarrhea started on the third postnatal day and persisted despite oral feeding with a variety of whole protein-, hydrolysate-, and amino acid-based infant formula feeds with differing contents of carbohydrate (including lactose, glucose polymer, glucosce, and fructose) and fat (predominantly long-chain triglyceride and medium-chain triglyceride). Diarrhea continued even when she was fed nutritionally inadequate fat-free glucose- and amino acid-based formula feeds. She was found to be hypocortisolemic, but hydrocortisone replacement did not affect the diarrhea. She became grossly obese despite a parenteral calorie intake of less than 50% of Recommended Daily Allowance. When 18 months old, she suffered a fatal cardiopulmonary arrest of uncertain cause.

Investigations required for clinical management confirmed malabsorption of monosaccharides (both glucose and fructose) and fat. Exocrine pancreatic
function was preserved (normal fecal elastase). Serial small-intestinal biopsies demonstrated persistent, patchy normoplasic villous atrophy (villus/crypt ratio 1:1 to 4:1), inconsistent with the gross gastrointestinal dysfunction. Some biopsies had a mild increase in lamina propria inflammatory cells, including eosinophils. The surface epithelium was organized normally, with enterocytes and a brush border membrane of normal appearance as observed by routine morphological methods and periodic acid-Schiff and alkaline phosphatase stains. Lactase, sucrase, and trehalase activities were normal in enterocytes of morphologically normal villi. Anti-enterocyte antibodies were absent. In fasting plasma, concentrations of proinsulin (1,079 pmol/l, normal <7) and des-64,65 proinsulin (71 pmol/l, normally undetectable) were very high, but levels of insulin (14 pmol/l, normal <60) and des-31,32 proinsulin (25 pmol/l, normal <16) were relatively normal. This pattern resembled that reported previously in Subject A (9). Investigation of Subject A’s lifelong and frequent episodes of hypoglycemia revealed hypoketosis (30–80 nmol/l, normal >200) associated with low/normal ACTH (2–3 pmol/l, normal 1–11) and high precursor concentrations (104 pmol/l, normal 5–40). The impaired POMC processing and cleavage of proinsulin between the B and C chains, both known activities of PC1 (3, 26), suggested that PC1 was defective.

Detection of compound-heterozygous PC1 mutations in Subject B

Sequencing of PC1 in Subject B and her parents revealed that she was a compound heterozygote for Glu250stop (937G>T, GenBank X64810) and Ala213del due to deletion of GCA or CAG (Figure 1a). The former mutation was predicted to truncate the PC1 protein within the catalytic domain, while the latter mutation deletes a highly conserved alanine residue near the catalytically essential His208 (Figure 1b). Substitution of Glu250 results in a premature stop codon, which is predicted to truncate the protein at position 251, resulting in the loss of the catalytic domain and the entire enzyme. The Ala213 deletion results in the loss of a single alanine residue, which is highly conserved in PC1 homologs. This mutation is predicted to truncate the protein at position 212, resulting in the loss of the catalytic domain and the entire enzyme.

Intestinal phenotype of Subject A

The association of severe gastrointestinal disturbance with PC1 mutations in Subject B led us to re-evaluate the surviving 48-year-old PC1-deficient adult, Subject A. Upon questioning, she revealed a lifelong history of abdominal bloating and alternating diarrhea and constipation. Upper and lower gastrointestinal endoscopy revealed normal appearances, and a duodenal biopsy was morphologically normal. Celiac disease serology was negative. Fasting gastric pH was normal and responded appropriately to food. Normal fecal elastase levels indicated an absence of pancreatic exocrine insufficiency. However, there was evidence for gross abnormalities of small-intestinal absorptive function, with high-volume diarrhea, steatorrhea, bile salt malabsorption, and impaired vitamin B12 uptake even in the presence of administered intrinsic factor (Table 1).

Processing of enteroendocrine cell–derived prohormones in Subject A

In view of the death of Subject B, investigation of the impact of PC1 deficiency on prohormone processing in enteroendocrine cells focused on Subject A, using established chromatography and immunoassays to examine progastrin, proglucagon, and their products in plasma, during fasting and in response to food.

Progastrin. Progastrin (residues 1-101) is cleaved at Arg57,58 and Arg94,95 to release a peptide that becomes the C-terminally glycine-extended peptide G34gly. Carboxyamidation of G34gly forms G34, some of which is cleaved at Lys74,75 to release the mature heptadecapeptide gastrin G17 (Figure 2a) (15). While fasting plasma progastrin immunoreactivities in Subject A and controls were similar, postprandial concentrations were higher in the subject (Figure 2b). Similarly, carboxyamidated gastrin immunoreactivities were higher postprandially in Subject A than in controls (Figure 2c). To determine whether this arose from increased levels of amidated gastrin intermediates such as G34, G17 was measured with antisera L6, which reacts preferentially with G17 (16). Though the levels of mature G17 measured with L6 were indeed lower than those of total carboxyamidated gastrin measured with L2, the ratio of L2 to L6 immunoreactivity was within reference limits (2 to 5) (27), suggesting unimpeded cleavage of G34 to G17 at the Lys74,75 site. This view was supported by the finding that plasma free N-terminal G17 immunoreactivity (antisera 1295) was also within reference limits (data not shown) (16). Thus, despite the apparently normal secretion of mature gastrin, the elevated postprandial progastrin/gastrin ratio suggests some impairment of progastrin processing.

Proglucagon. Plasma free C-terminal glucagon immunoreactivity, which normally reflects glucagon, was 333 pg/ml 120 minutes postprandially, compared with 253 pg/ml in controls. However, mid-sequence glucagon immunoreactivity was 3,430 pg/ml, compared with 328 pg/ml in controls, raising the possibility of high plasma levels of proglucagon, glicentin, and/or oxyntomodulin, which also react in this assay (Figure 3a). In view of the reported dependence of PC1 of GLP-1 cleavage from proglucagon (5, 6), plasma immunoreactivity of the mature product, amidated C-terminal GLP-1, 60 minutes after food intake was unexpectedly high: 158 pg/ml, compared with 162 ± 53 pg/ml in controls. These findings were confirmed by the second panel of site-specific RIAs (data not shown). The molecular forms contributing to immunoreactivities were identified by chromatography with RIA of eluted peptides, which showed that the reportedly PC1-dependent products GLP-11-36amide, GLP-17-36amide (the bioactive form), GLP-2, and glicentin were all present in plasma and increased with feeding (Figure 3, b–d). Chromatography peaks consistent with proglucagon (glucagon, GLP-1, and GLP-2 immunoreactivities) and

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proglucagon minus the C-terminal GLP-2 peptide (glucagon and GLP-1 immunoreactivities only) confirmed observations of high–molecular weight proglucagon forms made with direct plasma assays (Figure 3, c and d, and data not shown). Thus, proglucagon processing was highly impaired despite the continued secretion of mature L cell–type products.

**Functional study of mutant PC1s**

**Expression of recombinant WT PC1, PC1-Gly593Arg, and PC1-Ala213del.** WT PC1 is synthesized as inactive pro-PC1. Cleavage of the propeptide occurs rapidly by an autocatalytic process in the ER, and the carboxyl terminus is cleaved later in the secretory pathway (28, 29). In order to study the biosynthesis, maturation, and secretion of mutant PC1 lacking Ala213, the deletion was introduced into PC1 cDNA and overexpressed in CHO-K1 cells, which have been shown to express and secrete active recombinant PC1 (10, 29). WT PC1 was detected intracellularly only in its processed form lacking the propeptide, consistent with rapid cleavage (Figure 4a, top panels). The majority of immunoreactive PC1, including all carboxy-terminally processed PC1, was secreted. Carboxy-terminal processing was similar to that of recombinant mouse PC1 expressed in CHO-DG44 cells (29). Differences in molecular weight and relative amounts are pertinent to human PC1 (data not shown). PC1-Gly593Arg from Subject A was only detected intracellularly as pro-PC1, as reported previously (10). Surprisingly, PC1-Ala213del was detected as both pro-PC1 and PC1, the latter being secreted to some extent. This indicated that propeptide processing of PC1-Ala213del was not entirely blocked. Similar results were obtained using βTC3 cells, which express endogenous PC1 and efficiently store mature and carboxy-terminally truncated PC1 (Figure 4a, bottom panels). Recombinant PC1 in these cells was selectively detected by means of the FLAG epitope tag.

**PC1-Gly593Arg and PC1-Ala213del do not process substrates in trans.** Since PC1-Ala213del displayed detectable autoprocessing, studies were performed to test for residual activity for substrates in trans. Neither recombinant mutant PC1 processed detectable amounts of a fluorogenic substrate (Figure 4b). These studies were initially performed using recombinant protein immunopurified from lysates of βTC3 cells, but similar results were obtained using cell lysates and conditioned medium from transfected CHO-K1 cells (data not shown).

**Binding and lack of internal cleavage of the propeptide of PC1-Ala213del.** Since PC1-Ala213del displayed detectable autoprocessing in cis (autocatalytic propeptide processing) but not in trans, the possibility that the propeptide remained associated with the enzyme after cleavage was investigated. It has been shown that the propeptide of another member of the PC family, furin, remains associated until a second internal autocatalytic cleavage occurs in the more acidic environment of the trans-Golgi network (30, 31). This step is necessary for release of the propeptide and hence activation of furin. A similar internal cleavage site is present in the propeptide of PC1 (−P-R-R-S-R-R↓). To study the association of the propeptide with PC1, we replaced Ile86 with Met in both PC1 and PC1-Ala213del. The intact propeptide (∼10 kDa) co-immunoprecipitated with both constructs. However, the internally cleaved form (∼4 kDa) was detected only in PC1 and not in PC1-Ala213del. Since only the intact propeptide is a potent inhibitor of PC1 (32), the impaired internal cleavage of the propeptide in PC1-Ala213del provides further evidence for the deleterious effect of the mutation. Together, these results show that PC1-Gly593Arg and PC1-Ala213del do not have significant residual activity.

**Table 1**

Results of tests of small-intestinal absorptive function in Subject A

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
<th>Reference</th>
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<tr>
<td>3-Day fecal weight</td>
<td>1,516 g</td>
<td>&lt;600 g</td>
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<tr>
<td>Fecal fat content</td>
<td>59 mmol/24 h</td>
<td>&lt;18 mmol/24 h</td>
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<td>Seleno-homotaurocholate absorption test</td>
<td>0.8%</td>
<td>&gt;8%</td>
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<td>Serum B&lt;sub&gt;12&lt;/sub&gt;</td>
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<td>&gt;223 ng/l</td>
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<tr>
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<td>7.5%</td>
<td>&gt;12%</td>
</tr>
<tr>
<td>B&lt;sub&gt;12&lt;/sub&gt; absorption with intrinsic factor</td>
<td>7.5%</td>
<td>&gt;12%</td>
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PC1-ssv is incorrectly spliced. The second allele of Subject A contains a variant at position +4 of the intron 5 donor splice site (A→C), which causes skipping of exon 5 from mature mRNA, resulting in a frameshift, 14 ectopic amino acids, and a premature stop codon (10). Although no correctly spliced PC1 was found by RT-PCR, small amounts could have remained undetected. To test this possibility, we constructed WT and splice-site variant (PC1-ssv) minigenes containing introns 4 and 5. When expressed in CHO-K1 cells, the minigene that contained WT sequence was correctly spliced, resulting in full-length PC1, which was efficiently secreted (Figure 5a). Transfection of PC1-ssv, on the other hand, failed to result in the expression of any detectable full-length PC1. However, the two low–molecular weight (22- and 24-kDa) proteins that were detectable intracellularly were consistent with exon 5 skipping. The predicted molecular weight of the unglycosylated truncated protein lacking the signal peptide, but containing the propeptide and the ectopic sequences, is 19.574 kDa. These immunoreactive bands contain the same core protein but are heterogeneous in Asn-link glycosylation, as deglycosylation by endoglycosidase F results in a single protein of approximately 19.5 kDa (Figure 5a, right panel). A more sensitive way to detect trace amounts of correctly spliced PC1 is to measure processing activity using a synthetic substrate, but no activity was detected by this method either (Figure 5b). Residual activity was undetectable in conditioned medium and cell lysates from transfected HEK-293T, AtT-20, αTC1-6, βTC3, and Neuro2A cells as well (data not shown).

In summary, no processing activity could be detected in the PC1 variants of Subject A or the Ala213del protein of Subject B.

Analysis of further circulating prohormones and products in Subject A

POMC. The high plasma POMC immunoreactivity and the near-normal ACTH and cortisol immunoreactivities seen previously in Subject A were confirmed and found to be physiologically regulated in terms of diurnal rhythm and response to exogenous CRH, despite the processing abnormality (Figure 6b) (9). The timing of the CRH response was consistent with secretion via the regulated pathway, and the cortisol response indicated release of bioactive peptide. However, analysis of processing products only in terms of immunoreactivity is potentially misleading, since epitopes may be common to otherwise distinct peptides, e.g., terminally extended forms. Therefore, to identify the POMC products with greater certainty, plasma peptides in Subject A were separated by chromatography prior to site-specific immunoassays. HPLC with RIA of eluted C-terminal ACTH showed that immunoreactive ACTH in plasma was indeed authentic ACTH 1-39 (phosphorylated and nonphosphorylated) (Figure 6c). Size-exclusion gel chromatography of plasma with β endorphin RIA and POMC IRMA revealed the presence of authentic β lipotropin and POMC (Figure 6d). CLIP and β endorphin, which are formed by the action of PC2 on POMC (33), were absent, suggesting that PC2 does not substitute for PC1 in processing POMC (Figure 6, c and d).

Proinsulin. In postprandial plasma, the four healthy heterozygous children of subject A had ratios of HPLC peaks of proinsulin to des-31,32 proinsulin (respectively PC1 substrate and product after subsequent carboxypeptidase activity) which were 1.65, 5.25, 0.75, and 1.00. Subject A had a ratio of 2.79 and controls 0.73 in controls 25.7. Thus simple heterozygosity for PC1 mutations did not significantly impair proinsulin processing.

Procalcitonin. Plasma procalcitonin in Subject A was 21 pg/ml by RIA, compared with 33 ± 16 pg/ml in controls, and was less than 300 pg/ml (normal) by immuno-chemiluminescent assay.
The evidence for small-intestinal failure in both subjects was strong. Subject B was extensively investigated for known causes of congenital diarrhea; this investigation included multiple small-intestinal biopsies. Fecal analysis revealed failure to absorb multiple dietary sugars and other nutrients. The intestinal failure was so severe and protracted as to require lifelong parenteral feeding. Subject A has had lifelong intermittent diarrhea and constipation, which had been overlooked because of the complexity of her endocrine and metabolic illness. Although the lower small intestine of Subject A was not biopsied, the combination of fat, bile salt, and vitamin B12 malabsorption with normal pancreatic exocrine function and gut transit times indicated an absorptive defect in the small intestine.

We hypothesized that PC1 deficiency caused intestinal malfunction through failure of maturation of propeptides within the enteroendocrine cells and nerves that express PC1 throughout the gut (34). Using established chromatographic and immunoassay systems, we studied circulating products of the gut peptides progastrin and proglucagon. The finding of elevated levels of progastrin and proglucagon...
provided in vivo evidence that, indeed, prohormone processing in enteroendocrine cells was abnormal.

Gastrin is cleaved from progastrin in the neuroendocrine G cells of the gastric antral and proximal duodenal mucosa. PC1 is a candidate for this role, because its protein and mRNA colocalize with gastrin expression (15, 34). Although in Subject A postprandial plasma levels of mature gastrin were within the reference range, progastrin levels were supranormal (Figure 2b), suggesting a role for PC1, but not a critical one. This is consistent with immunocytochemistry in humans, which revealed PC1 to be quantitatively and spatially less distinctly related to gastrin than is PC2 (34).

GLP-2, which is formed in the gut from proglucagon, was an obvious candidate for a role in the gut pathology, because it has trophic effects on small-intestinal epithelium (35). However, direct plasma assays and chromatography/RIA showed that GLP-2 was still present in plasma despite the undetectable activity of PC1 mutants and the reported essential role of PC1 (3, 5). Since GLP-1 is formed in amounts equimolar with those of GLP-2, the presence in plasma of mature GLP-17-36amide was also evidence of GLP-2 formation, and vice versa. Although the elevated precursor levels clearly revealed the importance of PC1 for proglucagon processing, the continued GLP-1 and GLP-2 secretion despite undetectable activity of PC1 mutants raised the possibility of a redundant PC1-like activity, e.g., PC5, which is widely expressed in the gut (34–38) and might become more active as a result of hyperplasia of L cells analogous to the A cell hyperplasia of PC2 deficiency (39). Interestingly, PC1-null mice too have a gastrointestinal phenotype (bulky moist stools), but GLP-2 was undetectable in gut extracts (8). However, because plasma levels were not measured in the mouse and we could not study the human gut directly, the relationship between the murine and human findings remains unresolved, although it is very likely that, in both species, small-intestinal dysfunction results from disordered prohormone processing within enteroendocrine cells. It is possible that the relative importance of gut neuropeptides and their functions and convertases differs between mice and humans such that the gut dysfunction in humans relates to deficiency of peptides other than GLP-2, e.g., promotilin and pro-PYY products. Plasma levels reflect contributions from multiple sources, so the rise in plasma glucagon after the test meal (Figure 3d) might conceivably have been related to A cells rather than abnormal processing in L cells. Interestingly, serum glucagon was also elevated (2.5-fold) in the PC1-null mouse (40).

Figure 5
Transfection of a PC1 minigene containing the splice-site variant resulted in undetectable full-sized PC1 protein and catalytic activity. (a) Western blot (left and middle panels) and immunoprecipitation analysis (right panel) of HEK-293T cells transfected with empty vector (C) or PC1 minigene containing WT or splice-site variant (SSV) sequences. Cells used for immunoprecipitation were pulse-labeled for 1 hour. Endo F, endoglycosidase F. (b) Fluorogenic assay of PC1 proteolytic activity using PC1 immunopurified from lysates of aTc1-6 cells transfected as in a. Open circles, empty vector; filled diamonds, PC1 minigene; open squares, PC1 splice-site variant minigene.
We wondered whether the differences between the two subjects in terms of survival and of severity of gastrointestinal phenotype were explicable on the basis of the degree and type of enzymatic dysfunction produced by their mutations. Results obtained with recombinant PC1-Gly593Arg and PC1-Ala213del strongly suggested that both mutations blocked enzymatic activity toward substrates in trans. PC1-Gly593Arg appeared to have lost autoprocessing activity as well. As a consequence, the pro-PC1 was retained in the ER (10), as inactive forms of furin and PC7 have been shown to be (41–43). PC1-Ala213del had reduced autoprocessing activity, but no activity for substrates in trans. Separation of these two activities has also been reported for furin mutants (44) and can be explained in two ways. Firstly, propeptide cleavage is a prerequisite for exit from the ER but does not generate an active enzyme per se. As has been shown for furin, a second internal cleavage is required for release of the inhibitory propeptide. This internal cleavage was found to be impaired in PC1-Ala213del. Secondly, the separation of the activities might be explicable in terms of differences between the kinetics of zero-order autocatalytic cleavage and first-order cleavage of substrates in trans.

Gly593 is conserved in all PCs of all species, and, based on crystal structures of mouse furin and yeast kexin (45, 46), it is predicted to be located at the end of a β-strand. Because of its tight fit, replacement with a bulky arginine is likely to disturb existing salt bridges and cause deformation of the β-sheet. Mutagenesis studies have recently indicated that even substitution by a small amino acid like alanine is not tolerated (47). Ala213 is predicted to be located within the same hydrophobic helix as the catalytic His208 (45, 46, 48, 49). Deletion of Ala213 shortens the helix, which will be pulled into the molecule. The charged Glu215 at one end of the helix may prevent movement there, forcing dislocation of His208 on the other end. This might be expected to reduce catalytic activity and destabilize the protein through distortion of the local structure. Indeed, both PC1-Gly593Arg and PC1-Ala323del, but not WT PC1, were seen in immunoprecipitations (data not shown) to bind BiP, an ER chaperone that binds to unfolded and misfolded proteins and prevents their export from the ER (50).

The splice-site mutation in intron 5 of Subject A was previously shown by RT-PCR to cause exon skipping of exon 5 (10). Here we have shown by a more robust method in multiple cell types that no correctly spliced mRNA, and hence no active PC1, was produced. Only a truncated protein, the predicted translation product of mRNA devoid of exon 5, was seen. This protein and the truncated protein encoded by the Glu250stop mutant allele in Subject B both lack the serine residue known to be essential for serine proteases and will therefore not be catalytically active. In conclusion, these in vitro data provide strong evidence for the complete absence of residual PC1 activity in both patients, and, therefore, differences in clinical presentation are likely to reflect the influence of genetic and environmental background.

The recent description of mice rendered null for PC1 by homologous recombination confirmed features of PC1 deficiency first reported in humans, such as defective proinsulin and POMC processing (9, 40). However,
 Subject A’s children, who were heterozygous for PC1 (effectively PC1 substrate and product, respectively) in...


