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Deficiency of the human cysteine protease inhibitor cystatin M/E causes hypotrichosis and dry skin

Ellen H. J. van den Bogaard, PhD¹, Michel van Geel, PhD^{2,3,4},
Ivonne M. J. J. van Vlijmen-Willems, BSc¹, Patrick A. M. Jansen, PhD¹, Malou Peppelman, PhD¹,
Piet E. J. van Erp, PhD¹, Selma Atalay, MSc¹, Hanka Venselaar, PhD⁵, Marleen E. H. Simon, MD, PhD⁶,
Marieke Joosten, MD, PhD⁷, Joost Schalkwijk, PhD¹ and Patrick L. J. M. Zeeuwen, PhD¹

Purpose: We aimed to assess the biological and clinical significance of the human cysteine protease inhibitor cystatin M/E, encoded by the *CTS6* gene, in diseases of human hair and skin.

Methods: Exome and Sanger sequencing was performed to reveal the genetic cause in two related patients with hypotrichosis. Immunohistochemical, biophysical, and biochemical measurements were performed on patient skin and 3D-reconstructed skin from patient-derived keratinocytes.

Results: We identified a homozygous variant c.361C>T (p.Gln121*), resulting in a premature stop codon in exon 2 of *CST6* associated with hypotrichosis, eczema, blepharitis, photophobia and impaired sweating. Enzyme assays using recombinant mutant cystatin M/E protein, generated by site-directed mutagenesis, revealed that this p.Gln121* variant was unable to inhibit any of

its three target proteases (legumain and cathepsins L and V). Three-dimensional protein structure prediction confirmed the disturbance of the protease/inhibitor binding sites of legumain and cathepsins L and V in the p.Gln121* variant.

Conclusion: The herein characterized autosomal recessive hypotrichosis syndrome indicates an important role of human cystatin M/E in epidermal homeostasis and hair follicle morphogenesis.

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INTRODUCTION

The importance of regulated proteolysis in epithelia and execution of protease-mediated processes^{1–4} is well illustrated by the identification of pathogenic variants in proteases and their inhibitors in a variety of human cornification/hair diseases (Supplementary Table S1 online).^{5–11} These include a number of cysteine proteases of the lysosomal cathepsin family and their cognate inhibitors of the cystatin family. In addition, studies in mice have shown that misregulation of proteolysis disturbs epidermal homeostasis, as witnessed by for example cathepsin L deficiency,^{12,13} targeted ablation of cathepsin D,¹⁴ and cystatin M/E deficiency.¹⁵

The protein encoded by the human *CST6* gene (MIM 601891) was originally discovered independently by two groups and named cystatin M or cystatin E (here further referred to as cystatin M/E).^{16,17} Cystatin M/E is an atypical

cystatin in that it was found to inhibit both lysosomal cysteine proteases such as cathepsin L (CTSL) and cathepsin V (CTSV), and the asparaginyl endopeptidase legumain (LGMN). We showed that cystatin M/E had an unusually tissue-specific expression pattern with high expression levels restricted to the skin.¹⁸ Our research on the function of this protein led to the discovery that the phenotype of the spontaneous *ichq* mouse mutant was caused by homozygosity for *Cst6* null alleles. The resulting cystatin M/E deficiency in these mice causes disturbed epidermal cornification, impaired barrier function, and neonatal lethality because of excessive transepidermal water loss.^{15,19,20} Biochemical evidence indicated that human and mouse cystatin M/E controls the activity of several proteases and may indirectly regulate activity of epidermal transglutaminases, which are key enzymes in skin cornification.²¹ Using double knockout

¹Department of Dermatology, Radboud Institute for Molecular Life Sciences (RIMLS), Radboud University Nijmegen Medical Center (Radboudumc), Nijmegen, The Netherlands;

²Department of Dermatology, Maastricht University Medical Centre, Maastricht, The Netherlands; ³Department of Clinical Genetics, Maastricht University Medical Centre, Maastricht, The Netherlands; ⁴GROW Research Institute for Oncology and Developmental Biology, Maastricht University Medical Centre, Maastricht, The Netherlands; ⁵Center for Molecular and Biomolecular Informatics, RIMLS, Radboudumc, Nijmegen, The Netherlands; ⁶Department of Medical Genetics, University Medical Centre Utrecht, Utrecht, The Netherlands; ⁷Department of Clinical Genetics, Erasmus MC, University Medical Center, Rotterdam, The Netherlands. Correspondence: Patrick L. J. M. Zeeuwen (Patrick.Zeeuwen@radboudumc.nl)

These authors contributed equally: Joost Schalkwijk and Patrick L.J.M. Zeeuwen.

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models in mice, we showed that a tightly regulated balance between cathepsin L and cystatin M/E is essential not only for tissue integrity of the epidermis, but also for maintenance of healthy hair follicles (HFs) and corneal epithelium.²² Our most recent data, in which we rescued the lethal skin phenotype of cystatin M/E-deficient mice by transgenic, epidermis-specific, re-expression of cystatin M/E under control of the human involucrin promoter, showed that cystatin M/E is not only a key molecule in a biochemical pathway that controls stratum corneum (SC) homeostasis but additionally plays an important role in maintaining HF integrity.²³

In the current study, we report for the first time a loss-of-function (LOF) variant in the human *CST6* gene, which causes a novel autosomal recessive hypotrichosis syndrome.

MATERIALS AND METHODS

Patients

We ascertained a consanguineous Turkish family with hypotrichosis, including three affected and four unaffected individuals (Fig. 1a). All patients (PII:1, PII:2, and PIII:2) were offspring of parents who were first cousins (in generation I and generation II). Saliva samples for DNA extraction were collected from all individuals. Biophysical measurements were performed on the right upper ventral forearm skin of PII:2 (father) and two healthy age-matched male controls. Skin biopsies from these individuals were taken from the same body location. Photographs were taken from PII:2 (father) and PIII:2 (daughter, index patient). The parents did not give permission to publish frontal facial photographs nor was it allowed to take biopsies from the index patient (PIII:2). We obtained informed consent from all subjects using protocols approved by the local ethics committee (CMO area Arnhem-Nijmegen), which comply with the Declaration of Helsinki principles.

Infinium_CytoSNP_850K genotyping array analysis

DNA amplification, tagging, and hybridization were performed according to the manufacturer's protocol (Illumina). Details of the procedure can be found in Supplementary Materials and Methods.

Exome sequencing

Exome sequencing was performed on genomic DNA extracted from the blood of the index patient (PIII:2). Details of the procedure can be found in Supplementary Materials and Methods.

Sequencing of the candidate gene

The pathogenic variant was confirmed in the patient by Sanger sequencing (BigDye terminator v1.1 sequencing kit, 3730 DNA analyzer, Applied Biosystems) using primers flanking exon 2 of *CST6* (sense: 5'-AAGTGAGGAT-CAAGGGTCAAGAC-3', antisense: 5'-CAGTTCAGGATG-GACAACCT-3'). Sequence data was visualized with Mutation Surveyor (version 4.0.7, Softgenetics). Segregation analysis of

the pathogenic variant in all available family members was performed.

Biophysical measurements

Details of the biophysical measurements (transepidermal water loss, subcutaneous [SC] hydration, skin redness and pigmentation) can be found in Supplementary Materials and Methods online.

Development of 3D human epidermal equivalents

Primary human keratinocytes were isolated from biopsies of family members PII:2 and II:3 and six healthy controls (wild-type *CST6*, *CST6*^{+/+}) and expanded according to the Rheinwald-Green protocol.²⁴ Passage 1–2 keratinocytes were used to generate 3D epidermal equivalents as described previously.²⁵ The quality and phenotype of the epidermal equivalents was investigated by morphology, protein expression (immunohistochemistry), and gene expression (quantitative polymerase chain reaction [qPCR]). The reconstructed epidermis was used to generate cell extracts to study protease activity (enzymology) and cystatin M/E protein expression (western blotting).

RNA isolation and qPCR analysis

A brief description of the procedure can be found in Supplementary Materials and Methods. An overview of the qPCR primer pairs used is shown in Table S2.

Morphological and immunohistochemical analysis

A brief description of the procedure can be found in Supplementary Materials and Methods. An overview of the antibodies used is shown in Table S3.

Protein extraction from epidermal equivalents

Proteins were extracted from the 3D epidermal equivalents. A brief description of the procedure can be found in Supplementary Materials and Methods.

Western blotting

Proteins were separated by SDS-PAGE using the NuPAGE electrophoresis system and precast 12% Bis-Tris polyacrylamide gels under reducing conditions (Invitrogen). A brief description of the procedure can be found in Supplementary Materials and Methods.

Production of recombinant cystatin M/E variant p.Gln121*

Cystatin M/E variant p.Gln121* was obtained using the recombinant plasmid pGEX2T-cystatin M/E and Quick-change site-directed mutagenesis kit (Stratagene). Primers used for the generation of the p.Gln121* variant were 5'-GGCAGCAGGGGCGCAGTAGGAGAAGCTGCGC-3' (sense) and 5'-GCGCAGCTTCTCCTACTGCGCCCTGCTGCC-3' (antisense). Sequence analysis was performed to check for the desired pathogenic variants in the complementary DNA (cDNA). Production and purification of recombinant wild-type and mutant forms of human

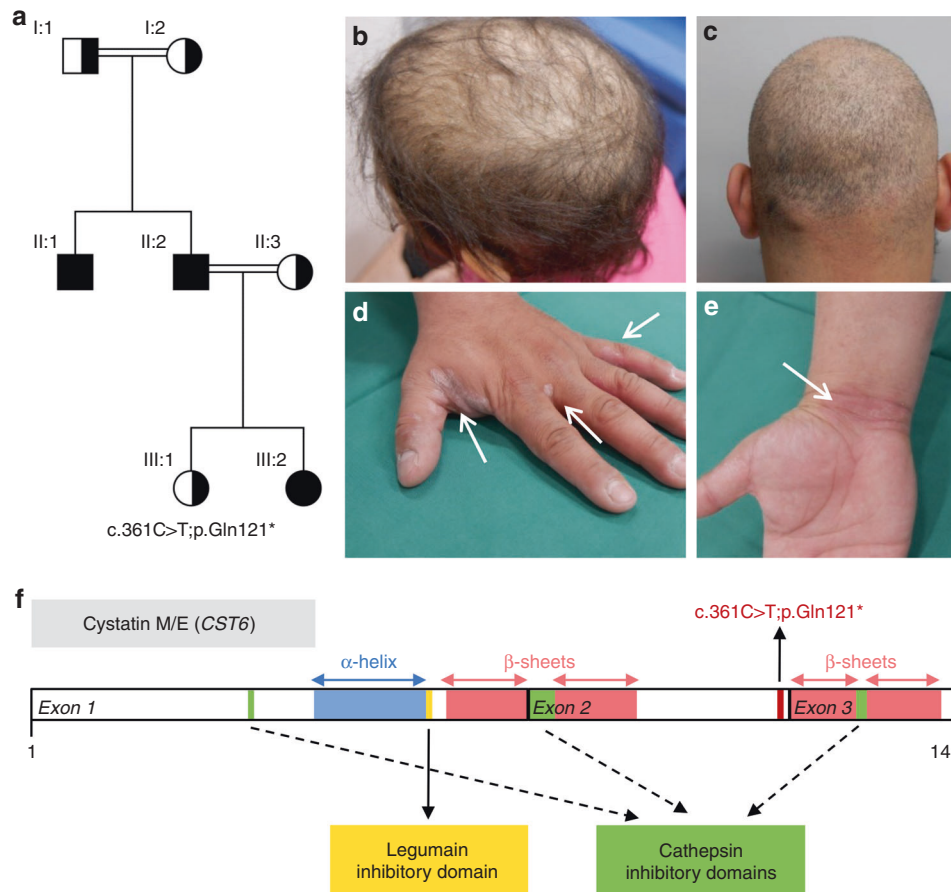


Fig. 1 Pedigree, clinical features, and schematic representation of *CST6* loss-of-function (LOF) variant. (a) Pedigree and genotypes of the family described. The homozygous LOF variant *c.361C>T* (p.Gln121*) was identified in individuals II:1, II:2, and III:2 (index patient), who were all born to consanguineous parents. Index patient is indicated by the arrowhead. Squares, males; circles, females; black filled symbols, affected; the half-filled symbols indicate heterozygous carriers of the *CST6* variant. (b) The index patient (III:2) and (c) her father (II:2) showed hypotrichosis due to progressive hair loss from three years of age. (d,e) The father developed eczematous lesions later in life. (f) The homozygous nonsense variant *c.361C>T* (p.Gln121*) results in a premature stop codon in exon 2 of the *CST6* gene. The cystatin M/E (*CST6*) protein is 149 amino acids in size and the positions of the α -helix, the β -sheets, and the cathepsin and legumain inhibitory domains are indicated.

cystatin M/E were performed as described previously.²¹ pGEX2T plasmids containing the wild-type or mutated cystatin M/E cDNAs were transformed into *Escherichia coli* BL21 Star (DE3) bacteria (Invitrogen). Cultures were grown from one single colony overnight at 37 °C in LB medium with ampicillin, followed by induction with isopropyl-1-thio- β -D-galactopyranoside (Roche) at a final concentration of 0.5 mM for 4 hours at 30 °C. Cells were lysed by four cycles of freezing in liquid nitrogen and thawing. Glutathione Sepharose 4B beads (Amersham Biosciences) were used for purification of glutathione S-transferase fusion protein from the lysate. Cleavage of the fusion protein into glutathione S-transferase and cystatin M/E occurred overnight at 4 °C using thrombin (Sigma) at a thrombin to fusion protein ratio of 1:500 in a buffer containing 50 mM Tris-HCl (pH 7.5), 150 mM NaCl, and 2.5 mM CaCl₂. Recombinant cystatin M/E was purified from glutathione S-transferase using glutathione Sepharose 4B beads and dialyzed against phosphate-buffered saline with a YM3 filter (Millipore). Protein concentration

was determined using a bicinchoninic acid (BCA) protein assay (Pierce).

Fluorimetric enzyme assays

Protease inhibitor activity of recombinant human wild-type cystatin M/E and the p.Gln121* variant against recombinant human legumain,²⁶ human CTSV, and human CTSL (both from R&D Systems) was determined by measuring inhibition of the proteases using fluorogenic synthetic substrates Z-Ala-Ala-Asn-MCA (Peptides International) for legumain, and Z-Leu-Arg-AMC (R&D Systems) for CTSV and CTSL. Protease inhibition was measured after an incubation period of 5 minutes (CTSL) or 10 minutes (legumain and CTSV) at room temperature. The buffer that was used in the legumain assay contained 0.1 M phosphate (pH 5.7), 2 mM EDTA, 1 mM dithiothreitol, 2.67 mM L-cysteine, and 5 μ g/ μ l bovine serum albumin. CTSV and CTSL assays were performed in a buffer (pH 5.5) containing 0.1 M acetate, 1 mM ethylenediaminetetraacetate (EDTA), 2 mM dithiothreitol, and 5 μ g/ μ l bovine serum albumin. Cathepsin and legumain protease activity was

determined in protein extracts by measuring the hydrolysis of fluorescent substrates as described above.

Structural modeling

The structure of human cystatin M/E in complex with legumain has been solved experimentally and can be found in Protein Data Bank (PDB) file 4N6O.²⁷ This PDB file was superposed on the structure of cystatin A in complex with cathepsin V, which can be found in PDB file 3KFQ (not yet published). Superposition of the assemblies and subsequent analysis was performed using the MOTIF-alignment option in the YASARA & WHAT IF Twinsit.^{28,29}

RESULTS

Clinical evaluation and genetic analysis

A 9-year-old girl (individual III:2 in Fig. 1a, index patient) was referred to the clinical geneticist by a dermatologist because of progressive hair loss resulting in hypotrichosis. Her father (PII:2) and her uncle (II:1) showed great similarities with regard to the clinical phenotype. The girl's mother (II:3) and 6-year-old sister (III:1) were unaffected. The girl's parents and her paternal grandparents had consanguineous marriages. The paternal grandparents (I:1 and I:2) did not have any symptoms of hypotrichosis. The clinical features of the index patient and her father are summarized in Table S4. From birth both suffered from a dry skin, blepharitis, and photophobia. After three years the clinical phenotype of hypotrichosis appeared (Fig. 1b, c). Overall body hair was sparse, very slowly growing, and both patients had only a few eyelashes (not shown as no permission was given to publish frontal facial photographs). Later in life the father developed eczematous lesions on hands (Fig. 1d, e) and feet (not shown). Both patients were not able to sweat, except after extreme physical exercise.

High-resolution single-nucleotide polymorphism (SNP) array did not show any pathogenic copy-number variants. After

pretest genetic counseling and consent of the index patient's parents, we performed exome sequencing. This analysis revealed a homozygous nonsense pathogenic variant c.361C>T (p. Gln121*) resulting in a premature stop codon in exon 2 of the *CST6* gene (Fig. 1f). We confirmed this pathogenic variant with Sanger sequencing in the index patient and other family members. The results were consistent with autosomal recessive inheritance of c.361C>T as the causative pathogenic variant. The father of the index patient, who had very similar symptoms, was also found to be homozygous for the variant. All clinically unaffected family members were heterozygous.

Epidermal acanthosis and abnormal distribution of stratum corneum water content

To investigate the biology underlying the phenotype of cystatin M/E deficiency in this family, we first performed several noninvasive biophysical measurements on the mid-ventral forearm skin of patient PII:2 (father) to detect possible abnormalities. Transepidermal water loss measurements in this patient were comparable with healthy controls, indicating no difference of inside-out skin barrier function (Table S5). Interestingly, spectrophotometric determination of skin redness and pigmentation showed that the a^* value was increased in the patient, demonstrating the presence of erythema. Pigmentation (b^* value) was normal. In vivo reflectance confocal microscopy (RCM) measurements showed a thickened epidermis (acanthosis) as well as a thickened SC in the cystatin M/E-deficient patient (Table S5). The water content of the patient's SC was comparable with healthy controls (Fig. 2a). However, the water content pattern of the patient was more heterogeneous compared with the controls (Fig. 2b, c). This finding indicates a different distribution of water content in the SC, which can explain the dry skin.

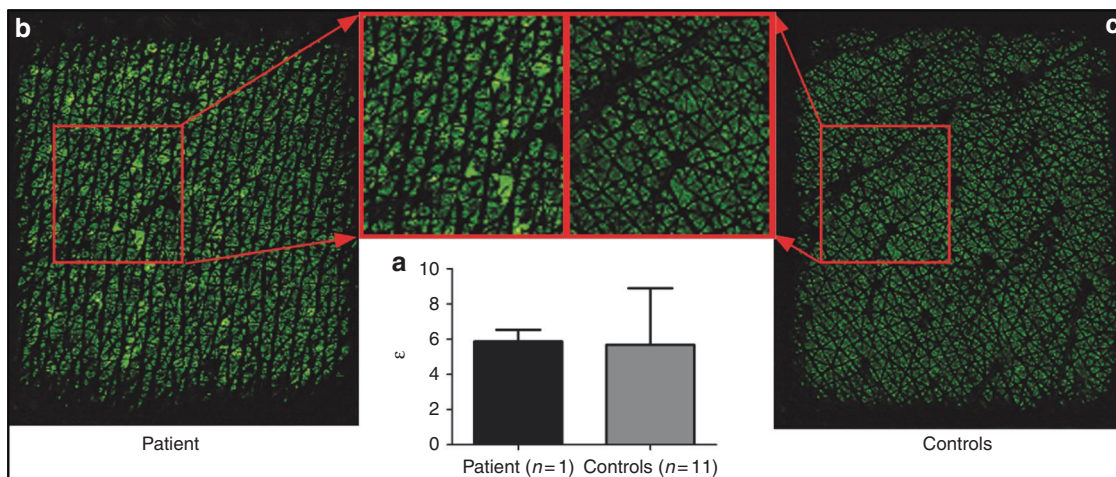


Fig. 2 Heterogeneous stratum corneum hydration in patient skin. (a) Permittivity measurements (bars represent mean \pm SD) of the patient and the healthy controls, demonstrating no overall quantitative difference between the patient and healthy controls. (b) Epsilon subcutaneous (SC) hydration image of the patient skin, showing a more heterogeneous water content pattern depicted by the larger black spaces (low SC hydration, representing the absence of water) between the green islets (high SC hydration, indicating the presence of water). (c) Epsilon SC hydration image of a healthy control, with small black areas between the homogeneous green islets, indicating a homogeneous distribution of the water content in the SC.

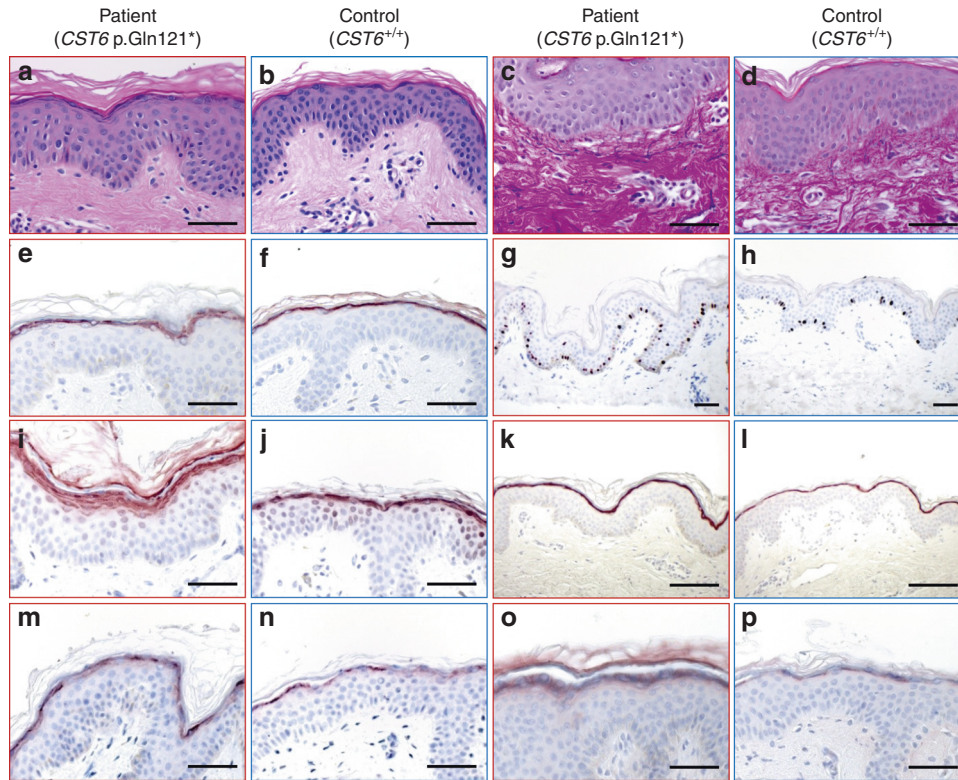


Fig. 3 Morphological and histological analysis of *CST6* loss-of-function (LOF) skin. Midventral forearm skin biopsies obtained from patient PII:2 (1st and 3rd column) and two healthy controls (2nd and 4th column) were examined for morphology and protein expression. Hematoxylin and eosin staining showed acanthosis and dense collagen fibers in the papillary dermis (**a, b**), which was confirmed by Elastica van Gieson staining (**c, d**). Cystatin M/E expression was present in the stratum granulosum (**e, f**). Induced expression was found in the patient skin for the proliferation marker Ki67 (**g, h**), epidermal differentiation proteins filaggrin (**i, j**), loricrin (**k, l**) and involucrin (**m, n**), and the crosslinking enzyme transglutaminase 1 (**o, p**). Scale bar represents 400 μm .

Abnormal epidermal proliferation and differentiation, and atypical dermal collagen fiber density

For in-depth phenotypical characterization we microscopically analyzed the skin biopsies. Hematoxylin and eosin (H&E) staining confirmed the RCM findings that the epidermal and SC thickness of the patient was increased as compared with six healthy controls (Fig. 3a, b). H&E staining also showed stronger eosinophilic collagen fibers in the papillary dermis of the patient's dermis compared with healthy control dermis. This was even more prominent in an Elastica van Gieson stained section, which showed densely packed collagen fibers in the patient's papillary dermis (Fig. 3c, d). Protein expression analysis revealed the presence of cystatin M/E in the upper granular layer of the epidermis (Fig. 3e, f) suggesting that the truncated protein was expressed. The increased number of proliferating cells, represented by Ki67 staining (Fig. 3g, h), was consistent with the increased epidermal thickness. We observed an increased protein expression of epidermal differentiation genes like filaggrin (FLG), loricrin (LOR), and involucrin (IVL) in the patient's skin (Fig. 3i–n). The expression of transglutaminase 1 (TGM1), an enzyme responsible for crosslinking structural proteins of the cornified envelope in epidermal keratinocytes, was clearly induced in the upper epidermal layers of the patient (Fig. 3o, p). The increased expression levels of *Ki67*,

TGM1, and the three differentiation genes were confirmed at the messenger RNA (mRNA) level (Supplementary Figure S1). To explain the observed increased redness (erythema) of the patient's skin, we evaluated CD31 expression (endothelial cell marker) as a marker for angiogenesis but no differences were found when compared with healthy controls (data not shown).

Disrupted protease binding sites in cystatin M/E p.Gln121*

The presence of cystatin M/E protein in the epidermis of the patient (Fig. 3e, f) indicated that the c.361 C>T pathogenic variant did not preclude translation of a (truncated) protein. Based on the known protease binding sites of cystatin M/E (Fig. 1f), we hypothesized that truncation of the protein would lead to a compromised inhibitory activity of cystatin M/E against cathepsins L and V. To study the functional consequences of the c.361 C>T pathogenic variant, we generated recombinant truncated cystatin M/E (p.Gln121*) by site-directed mutagenesis. Wild-type and mutant cystatin M/E were tested for inhibitory activity against several cathepsins and legumain. K_i values were determined using an Easson–Stedman plot as previously described.²¹ When the K_i could not be determined in case of very low affinity, the ratio of protease activity in the presence (V_i) and absence (V_0) of inhibitor was determined, and the remaining protease

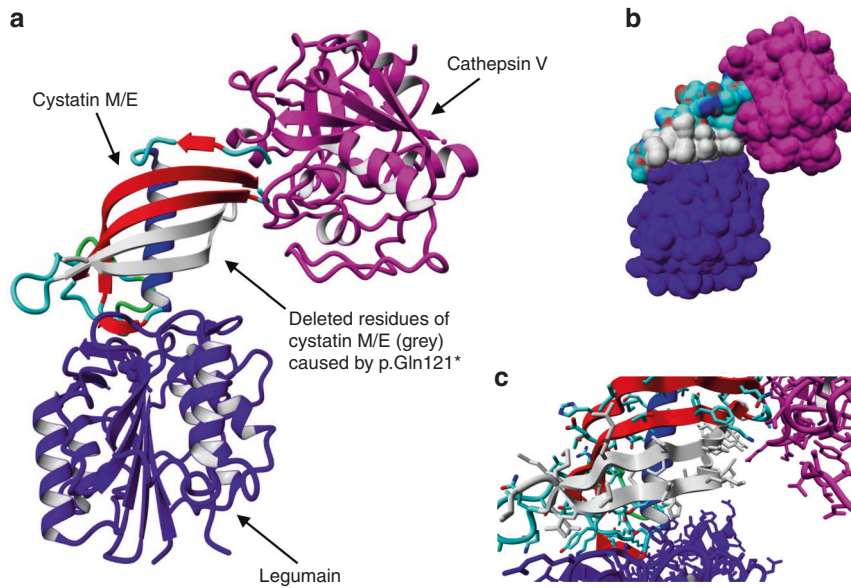


Fig. 4 3D visualization of the cystatin M/E–legumain–cathepsin V complex. The visualization was made by superposing Protein Data Bank (PDB) files 4N6O and 3KFQ. Legumain and cathepsin V are shown in purple and magenta respectively. Cystatin M/E is colored by element (β -strands = red, α -helix = blue, loops = cyan/green). The residues deleted after truncation of the protein are colored gray. **(a)** Overview of the complex in ribbon presentation. **(b)** Overview of the complex with protein surface shown. **(c)** Close-up of the protein–protein interaction sites with side chains of the residues shown.

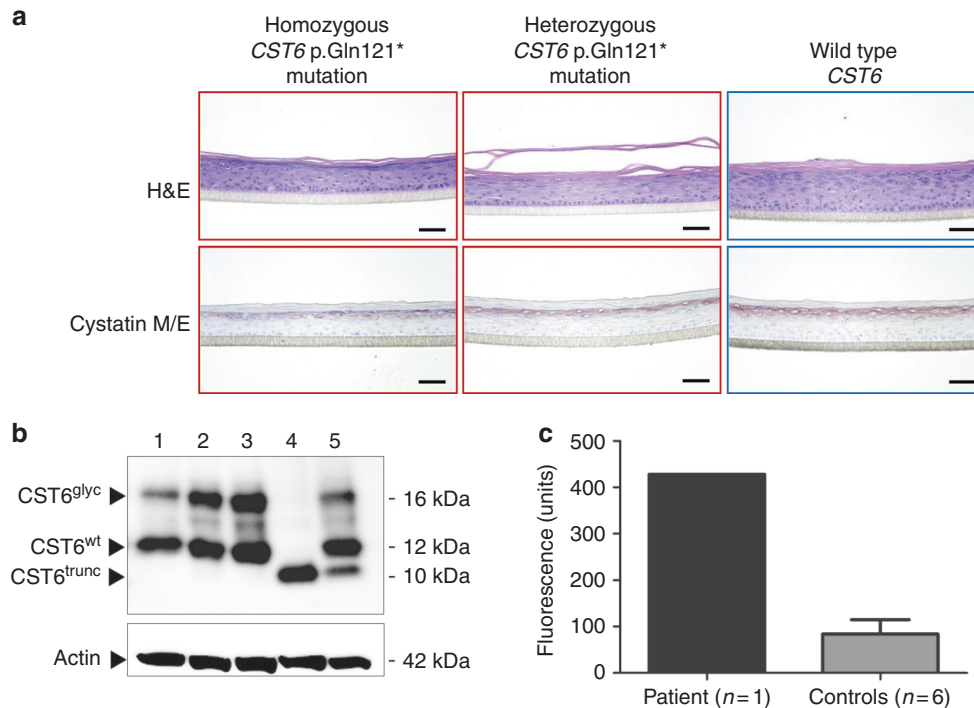


Fig. 5 Analysis of CST6 loss-of-function (LOF) 3D epidermal equivalents. **(a)** Hematoxylin and eosin (H&E) staining showed no specific morphological differences between epidermal constructs generated with primary keratinocytes that harbor a homozygous or heterozygous *CST6* p.Gln121* variant compared with the wild-type control(s). Cystatin M/E protein is expressed in the stratum granulosum of all constructs. Scale bar represents 400 μ m. **(b)** Western blot analysis of cell extracts derived from normal healthy skin biopsies (lane 1) and from epidermal equivalents from healthy controls (lanes 2 and 3), the patient (PII:2, father) with the homozygous *CST6* p.Gln121* LOF variant (lane 4), and the healthy mother (II:3) who is heterozygous for the variant (lane 5). *CST6*^{trunc} = truncated cystatin M/E, *CST6*^{glyc} = glycosylated cystatin M/E, *CST6*^{wt} = wild-type cystatin M/E. **(c)** Conversion of the fluorogenic z-A-A-N-AMC substrate. Cell extract of the patient-derived construct contains more free legumain activity compared with controls.

activity at a fixed high inhibitor (I) concentration (300 nM) was calculated. K_i values of wild-type human cystatin M/E and the p.Gln121* variant for interaction with human cathepsins and human legumain are shown in Table S6. Wild-type cystatin M/E is a high affinity inhibitor for cathepsin L, cathepsin V, and legumain as witnessed by their respective K_i values of 1.78 nM, 0.47 nM, and 0.25 nM,²¹ whereas cathepsin B is not inhibited by human cystatin M/E ($v_i/v_0 > 0.71$ at $[I] = 300$ nM). The K_i value of the p.Gln121* variant for interaction with cathepsins L and V could not be determined ($v_i/v_0 > 0.86$ and $v_i/v_0 > 0.98$ respectively at $[I] = 300$ nM), which indicates that the recombinant mutated variant was not able to inhibit any of these cathepsins. As the legumain inhibitory domain is positioned in the first exon (Fig. 1f) we anticipated that legumain inhibitory activity would be retained. This, however, was not the case as no inhibition was observed by the p.Gln121* variant ($v_i/v_0 > 1.09$ at $[I] = 300$ nM). As a control we used heat-inactivated wild-type recombinant cystatin M/E showing that this denatured protein was not able to inhibit both cathepsins and legumain.

Based on the position of the pathogenic variant (Fig. 1f) at the proximal end of exon 2 and just before two of the four β -sheets, we hypothesized that the protein structure of the truncated protein had changed, resulting in an inactive molecule. Indeed, 3D-modeling using the published X-ray crystal structures of the cystatin M/E–legumain (PDB file 4N6O) and cystatin A–cathepsin V (PDB file 3KFQ) complexes indicates that the p.Gln121* pathogenic variant will severely affect the protein structure and function (Fig. 4). According to the created in silico model of the complex, legumain and cathepsin V both seem to bind to their own distinct sites on cystatin M/E. However, both sites require interactions with residues located in the last two β -strands (direct), or with residues that are correctly positioned by those residues in the last two β -strands (indirect). As a result, the deletion of these β -strands caused by pathogenic variant p.Gln121* will affect these binding sites and thereby affect the interaction with other proteins.

Free legumain activity in cystatin M/E-deficient 3D skin equivalents

Because cystatin M/E is known to target three proteases—cathepsin L, cathepsin V, and legumain, all of which are expressed in human epidermis and HFs—we tried to assess the contribution of unrestricted activity of these proteases due to the p.Gln121* variant. Therefore, we generated 3D epidermal equivalents from the patient's keratinocytes (PII:2, homozygous *CST6* p.Gln121*), keratinocytes from the unaffected mother (II:3, heterozygous *CST6* p.Gln121*), and keratinocytes from healthy individuals ($N = 6$, homozygous *CST6* wild type). Similar to the in vivo biopsy material of PII:2, we observed cystatin M/E protein expression in the stratum granulosum of the homozygous *CST6* p.Gln121* epidermal equivalents (Fig. 5a). Western blot analysis revealed the expression of a truncated protein of

approximately 10 kDa in cell lysates of the p.Gln121* epidermal equivalent (Fig. 5b). The heterozygous p.Gln121* epidermal equivalent expressed both truncated and wild-type cystatin M/E as indicated by two bands, one at 12 kDa and one at 10 kDa (Fig. 5b). The 16-kDa bands indicate the presence of glycosylated cystatin M/E in the 3D epidermal equivalents. Clearly, the truncated cystatin M/E is not glycosylated as the presumed glycosylation site carrying an N-linked carbohydrate chain at position 137Asn is deleted due to the p.Gln121* variant.¹⁷ In contrast to in vivo skin we could not detect differences in the expression of specific genes/proteins involved in differentiation and proliferation (not shown). Protease activity measurements in the cell extracts of these epidermal equivalents indicated that cathepsin L and cathepsin V activity was low to undetectable in all epidermal equivalents (not shown). Strikingly, we found a high level of legumain activity in the patient-derived *CST6* p.Gln121* epidermal equivalent compared with six control *CTS6* wild-type epidermal equivalents (Fig. 5c). Although this finding did not allow unequivocal identification of the relevant protease responsible for the pathological alterations, it is consistent with the observation that legumain inhibitory activity is lost in the p.Gln121* variant.

DISCUSSION

Our previous genetic studies in mice and in vitro biochemical studies using human enzymes and inhibitors have provided information on the role of cystatin M/E in epidermal cornification (Figure S2A). Based on the phenotype of the cystatin M/E-deficient *ichq* mice, and the biochemical and cell biological properties of human cystatin M/E, we anticipated that deficiency in humans would cause a severe ichthyosis-like phenotype rather than a hair phenotype. For this reason we have previously sequenced the *CST6* gene in a large number (>100) of patients with autosomal recessive congenital ichthyosis who were negative for variants in known ichthyosis-associated genes. In none of these patients did we detect variants in the coding sequences of the *CST6* gene.³⁰ Although the patients described here do have a mild epidermal phenotype, their hypotrichosis is the most prominent and serious clinical feature, and led us to classify this syndrome as a heritable hair disease rather than an ichthyosis.

The clinical skin phenotype of the two patients was less severe than we expected based on our mouse studies and in vitro knockdown experiments.^{15,22,23,30} The severe and lethal skin phenotype in *ichq* mice, where the protein is completely absent, might be due to larger quantities of body hair, compared with humans, because neonatal keratinization of the HFs and subsequent dehydration is a hallmark of these mice. In the current study we show that the patient's keratinocytes still produce a significant amount of truncated protein, which is unable to inhibit protease activity. However, this truncated protein may still have some hitherto unknown functions that are important for a proper development of the epidermis, as complete short hairpin RNA (shRNA)-mediated

knockdown of cystatin M/E in a human skin equivalent showed a more severe phenotype.³⁰

We herein show abnormal epidermal proliferation and differentiation, and atypical dermal collagen fiber density in skin biopsies of the patient, features that we did not or could not observe in the *in vitro* epidermal equivalents generated with the patient's keratinocytes. These differences between the *in vivo* and *in vitro* situation are likely due the fact that the 3D skin model used here only contains keratinocytes in a sterile environment, thereby lacking interactions with skin microbes, fibroblasts, and/or other dermal and epidermal cells. Potential microbe–host or immune cell–mediated interactions may cause the phenotypical characteristics seen in our patient. Of note, there is currently no *in vitro* model available to grow hairs from epidermal keratinocytes, which precluded investigation of the hair phenotype *in vitro*.

Despite the fact that skin biopsies were taken from macroscopically healthy skin, histopathology and immunostaining revealed epidermal hyperproliferation and mildly abnormal differentiation compared with normal skin from healthy volunteers taken from the same anatomical location. Obviously it has to be taken into account that we could only study histology of one single patient (the father but not the daughter), and the observed abnormalities may be within the normal range. The dry skin observed in the two patients and the distribution of the SC water content were clearly abnormal compared with what is observed in healthy controls. It has to be noted that both patients had atopic eczema-like lesions, which may be mechanistically related to their dry skin. Speculatively, cystatin M/E deficiency may yet be another example of an epidermal defect that predisposes to atopic eczema, as has been reported for variants in filaggrin and genes that lead to a disturbed protease/inhibitor balance resulting in an aberrant skin barrier function.^{31–33} Observations in rare skin barrier diseases, for which there are now a few examples, will lend further support to initiatives to prevent atopic disease by strengthening skin barrier function in early life.

We demonstrated that due to the c.361 C>T LOF pathogenic variant in the *CST6* gene, the resulting truncated cystatin M/E protein is unable to inhibit legumain and the cathepsins L and V. As all three proteases are expressed in the human HF and also colocalize with cystatin M/E in the distal part and infundibulum of the HF³⁴ (see Figure S2B), it is likely that unrestricted activity of these proteases is the cause of hypotrichosis in patients. Our previous animal studies in double knockout mice and transgenic mice revealed that unrestricted cathepsin L activity is responsible for ichthyosis and lethality in the *ichq* mouse, but not for the persistent alopecia and fibrotic changes.^{22,23} However, it should be noted that the cystatin M/E controlled biochemical pathway in mice is distinct from humans. Mice do not have a cathepsin V orthologue, and we showed that mouse cystatin M/E differs from the human protein at amino acid position 64 (Asn instead of Asp), which renders it inactive against legumain. Ablation of legumain in mice with a cystatin M/E-deficient

background (*Cst6*^{-/-}*Lgmn*^{-/-}) did not allow survival beyond the neonatal phase and mice displayed exactly the same phenotype as *ichq* mice. Human cystatin M/E, however, is a specific and high affinity inhibitor of human legumain. In the current study we found free legumain activity in an extract of the epidermal construct generated from patient keratinocytes, which was more than fourfold higher than in control keratinocytes (Fig. 5c). This suggests that cystatin M/E is a major inhibitor of epidermal legumain activity. We were unable to quantitatively assess the effect of cystatin M/E deficiency on cathepsin L and V activities because no activity could be detected in mutant or wild-type cell extracts. We surmise that activity of these enzymes is masked by the presence of other inhibitors in the extracts, such as hurpin and the cystatins A, B, and C.^{21,35} Due to the lack of material from the patient (e.g., biopsy material including HFs) and the limitations of our 3D model (no hair growth) we are, at this point, not able to dissect the mechanism in more detail or identify the key protease that causes the hypotrichosis.

Our previous animal work has also shown evidence that free cathepsin L activity is responsible for the eye pathology in cystatin M/E-deficient mice.²² These mice developed severe keratitis, thickening of the corneal stroma, and squamous metaplasia of the corneal epithelium. The patients presented in the current study also have clinically relevant eye problems, although milder than in the mice, such as dry eyes, blepharitis, and photophobia. If these problems are also caused by unrestrained cathepsin L activity this might provide therapeutic opportunities.

In conclusion, we have identified a new autosomal recessive hypotrichosis syndrome caused by a LOF pathogenic variant in the *CST6* gene. These findings provide another example of the importance of protease–antiprotease balance in the regulation of epidermal and hair follicle homeostasis.

ELECTRONIC SUPPLEMENTARY MATERIAL

The online version of this article (<https://doi.org/10.1038/s41436-018-0355-3>) contains supplementary material, which is available to authorized users.

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DISCLOSURE

The authors declare no conflicts of interest.

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