

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Doi H, others, Matsumoto N, others.	Siblings with the adult-onset slowly progressive type of pantothenate kinase-associated neurodegeneration and a novel mutation, Ile346Ser, in PANK2: Clinical features and (99m)Tc-ECD brain perfusion SPECT findings.	J Neurol Sci	290 (1-2)	172-176	2010
Miyake N, others, Matsumoto N.	Loss of decorin dermatan sulfate impairing collagen bundle formation in a new type of Ehlers-Danlos syndrome.	Hum Mut	31(8)	966-974	2010
Kosho T, Miyake N, others, Matsumoto N.	A New Ehlers-Danlos Syndrome With Craniofacial Characteristics, Multiple Congenital Contractures, Progressive Joint and Skin Laxity, and Multisystem Fragility-related Manifestations.	Am J Med Genet	152A (6)	1333-1346	2010
Kimura S, others, Matsumoto N, Ishibashi M.	Rudimentary Claws and Pigmented Nail-like Structures on the Distal Tips of the Digits of <i>Wnt7a</i> Mutant Mice: <i>Wnt7a</i> Suppresses Nail-like Structure Development in Mice.	Birth Defects Res A Clin Mol Teratol	88(6)	487-496	2010
Kanazawa K, others, Matsumoto N	Choreo-ballistic movements in a case carrying a missense mutation in syntaxin binding protein 1 gene.	Mov Disord	25(13)	2265-2267	2010
Ng S, others, Matsumoto N, others.	Exome sequencing identifies <i>MLL2</i> mutations as a cause of Kabuki syndrome.	Nat Genet	42(9)	790-793	2010
Yamada-Okabe T, others, Matsumoto N.	Functional characterization of the zebrafish <i>WHSC1</i> -related gene, a homologue of human <i>NSD2</i> .	Biochem Biophys Res Commun	402(2)	335-339	2010
Miyake N, others	CHN1 mutations are not a common cause of sporadic Duane's retraction syndrome.	Am J Med Genet	152A(1)	215-217	2010

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Kato M, others	Frameshift mutations of the ARX gene in familial Ohtahara syndrome.	Epilepsia	51(9)	1679-1684	2010
Tohyama J, others, Matsumoto N.	Dandy-Walker malformation associated with heterozygous <i>ZIC1</i> and <i>ZIC4</i> deletion: Report of a new patient.	Am J Med Genet	155A(1)	130-131	2011
Furuichi T, others, Miyake N, Matsumoto N, others.	<i>CANT1</i> is also responsible for Desbuquois dysplasia, type 2 and Kim variant.	J Med Genet	48(1)	32-37	2011
Okada I, others, Miyake N, others, Matsumoto M, Saitsu H.	<i>SMOC1</i> is essential for ocular and limb development in humans and mice.	Am J Hum Genet	88(1)	30-41	2011

IV. 研究成果の刊行物・別刷

Loss-of-Function Mutations of *CHST14* in a New Type of Ehlers-Danlos Syndrome

Noriko Miyake,^{1*†} Tomoki Kosho,^{2†} Shuji Mizumoto,^{3†} Tatsuya Furuichi,⁴ Atsushi Hatamochi,⁵ Yoji Nagashima,⁶ Eiichi Arai,⁷ Kazuo Takahashi,⁸ Rie Kawamura,² Keiko Wakui,² Jun Takahashi,⁹ Hiroyuki Kato,⁹ Hiroshi Yasui,¹⁰ Tadao Ishida,¹⁰ Hirofumi Ohashi,¹¹ Gen Nishimura,¹² Masaaki Shiina,¹³ Hiroto Saito,¹ Yoshinori Tsurusaki,¹ Hiroshi Doi,¹ Yoshimitsu Fukushima,² Shiro Ikegawa,⁴ Shuhei Yamada,³ Kazuyuki Sugahara,³ and Naomichi Matsumoto^{1*}

¹Department of Human Genetics, Yokohama City University Graduate School of Medicine, Yokohama, Japan; ²Department of Medical Genetics, Shinshu University School of Medicine, Matsumoto, Japan; ³Laboratory of Proteoglycan Signaling and Therapeutics, Hokkaido University Graduate School of Life Science, Sapporo, Japan; ⁴Laboratory for Bone and Joint Disease, Center for Genomic Medicine, RIKEN, Tokyo, Japan; ⁵Department of Dermatology, Dokkyo Medical University, School of Medicine, Tochigi, Japan; ⁶Department of Molecular Pathology, Yokohama City University Graduate School of Medicine, Yokohama, Japan; ⁷Department of Pathology, Saitama Medical University, Saitama, Japan; ⁸Department of Environmental Immuno-Dermatology, Yokohama City University Graduate School of Medicine, Yokohama, Japan; ⁹Department of Orthopedics, Shinshu University School of Medicine, Matsumoto, Japan; ¹⁰First Department of Internal Medicine, Sapporo Medical University, Sapporo, Japan; ¹¹Division of Medical Genetics, Saitama Children's Medical Center, Saitama, Japan; ¹²Department of Radiology, Tokyo Metropolitan Kiyose Children's Hospital, Tokyo, Japan; ¹³Department of Biochemistry, Yokohama City University Graduate School of Medicine, Yokohama, Japan

Communicated by Jürgen Horst

Received 17 March 2010; accepted revised manuscript 24 May 2010.

Published online 8 June 2010 in Wiley InterScience (www.interscience.wiley.com). DOI 10.1002/humu.21300

ABSTRACT: Ehlers-Danlos syndrome (EDS) is a heterogeneous connective tissue disorder involving skin and joint laxity and tissue fragility. A new type of EDS, similar to kyphoscoliosis type but without lysyl hydroxylase deficiency, has been investigated. We have identified a homozygous *CHST14* (carbohydrate sulfotransferase 14) mutation in the two familial cases and compound heterozygous mutations in four sporadic cases. *CHST14* encodes dermatan 4-O-sulfotransferase 1 (D4ST1), which transfers active sulfate from 3'-phosphoadenosine 5'-phosphosulfate to position 4 of the *N*-acetyl-D-galactosamine (GalNAc) residues of dermatan sulfate (DS). Transfection experiments of mutants and enzyme assays using fibroblast lysates of patients showed the loss of D4ST1 activity. *CHST14* mutations altered the glycosaminoglycan (GAG) components in patients' fibroblasts. Interestingly, DS of decorin proteoglycan, a key regulator of collagen fibril assembly, was completely lost and replaced by chondroitin sulfate (CS) in the patients' fibroblasts, leading to decreased flexibility of GAG chains. The loss of the decorin DS proteoglycan due to *CHST14* mutations may preclude proper collagen bundle formation or maintenance of collagen bundles while the sizes and shapes of collagen fibrils are unchanged as observed in the patients' dermal tissues. These findings indicate the important role of decorin DS

in the extracellular matrix and a novel pathomechanism in EDS.

Hum Mutat 31:966–974, 2010. © 2010 Wiley-Liss, Inc.

KEY WORDS: Ehlers-Danlos syndrome; EDS; *CHST14*; dermatan sulfate; dermatan 4-O-sulfotransferase 1; D4ST1; collagen bundle formation; decorin

Introduction

Ehlers-Danlos syndrome (EDS) is a heterogeneous connective tissue disorder characterized by joint and skin laxity and tissue fragility [Steinmann et al., 2002], affecting as many as 1 in 5,000 individuals. The pathomechanisms of EDS consist of dominant-negative effects of mutant procollagen α -chains, α -chain haploinsufficiency, and deficiency of collagen-processing enzymes [Mao and Bristow, 2001]. In a revised nosology [Beighton et al., 1998], EDS was classified into six major types as well as additional minor forms. We previously described two unrelated patients showing characteristic facial and skeletal features with partial similarity to kyphoscoliosis type EDS but without lysyl hydroxylase deficiency (EDS-VIB) [Kosho et al., 2005]. Through long-term clinical evaluation of them as well as additional four unrelated patients including one reported previously [Yasui et al., 2003], we confirmed that these patients represent a new type of EDS [Kosho et al., 2010]. The evidence that two of six probands were born between consanguineous parents (Fig. 1A) suggests that this disease is inherited in an autosomal recessive fashion. Thus, we performed homozygosity mapping to find the disease-causative gene and successfully identified pathological mutations in the carbohydrate sulfotransferase 14 (*CHST14*: GenBank reference sequence, NM_130468.3) gene, in all six probands. *CHST14* encodes dermatan 4-O-sulfotransferase 1 (D4ST1), which transfers active sulfate to the *N*-acetyl-D-galactosamine (GalNAc) residues of dermatan sulfate (DS). Furthermore, we

Additional Supporting Information may be found in the online version of this article.

[†]The first three authors contributed equally to this article.

*Correspondence to: Noriko Miyake, Department of Human Genetics, Yokohama City University Graduate School of Medicine, 3-9 Fukuura, Kanazawa-ku, Yokohama 236-0004, Japan. E-mail: nmiyake@yokohama-cu.ac.jp or Naomichi Matsumoto, Department of Human Genetics, Yokohama City University Graduate School of Medicine, 3-9 Fukuura, Kanazawa-ku, Yokohama 236-0004, Japan. E-mail: naomat@yokohama-cu.ac.jp

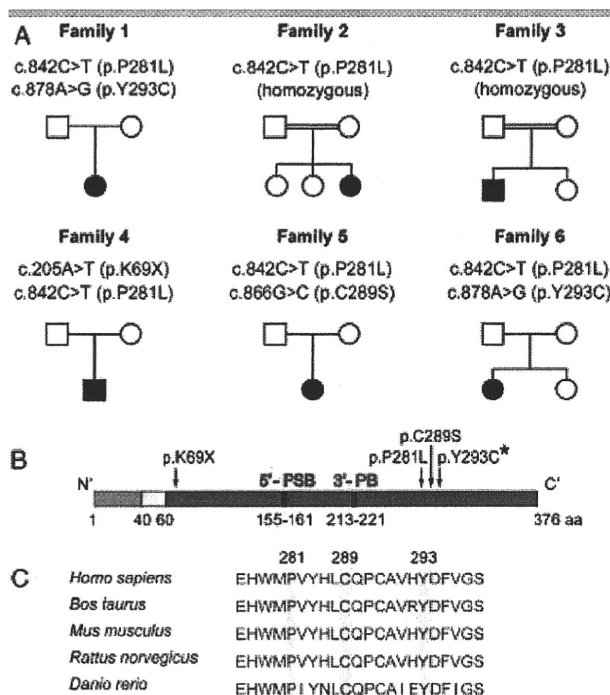


Figure 1. *CHST14* mutations in the patients. **A:** Pedigrees of the patients. Mutations in both alleles were found in all. **B:** A schematic representation of D4ST1 encoded by *CHST14*. Arrows indicate the position of mutations found in the patients. The red and blue boxes indicate a 5'-phosphosulfate binding site (5'-PSB) and a 3'-phosphate binding site (3'-PB), respectively. Light blue, yellow, and purple boxes denote cytoplasmic, transmembrane, and luminal regions, respectively. *p.Y293C (c.878A>G) is the same missense mutation identified in the Japanese ATCS sibs reported [Dündar et al., 2009]. **C:** D4ST1 amino acid alignment for three missense mutations evolutionarily conserved.

conducted pathological and glycochemical investigations to reveal the pathomechanism of this disease.

Materials and Methods

Subjects

We analyzed six Japanese patients clinically diagnosed as showing a specific type of EDS [Kosho et al., 2005, 2010] in this study. Briefly, they clinically resemble the kyphoscoliosis type EDS characterized by joint laxity, progressive scoliosis, tissue fragility with atrophic scars, easy bruising, arterial rupture, and Marfanoid habitus. However, the lysyl hydroxylase deficiency, which is the reliable diagnostic test for the kyphoscoliosis type, was not observed in the probands [Kosho et al., 2005, 2010]. One of the authors (T.K.) evaluated all cases. This study was approved by the institutional review boards of Yokohama City University School of Medicine, Shinshu University School of Medicine, and Hokkaido University Graduate School of Advanced Life Science. Informed consent was obtained from all subjects involved in this study.

Mapping and Mutation Analysis

We performed the whole genome linkage analysis using Affymetrix Human Mapping single nucleotide polymorphism (SNP) 10K XbaI 142 2.0 array (Affymetrix, St. Clara, CA) in two affected probands (patients 2 and 3) and six unaffected

members from two consanguineous families. Haplotype analysis was performed using seven microsatellite markers (*D15S1002*, *D15S1007*, *D15S1118*, *D15S1044*, *D15S214*, *D15S978*, and *D15S1117*) purchased from Applied Biosystems (Bedford, MA). These markers were designed based on the Marshfield genetic map (<http://research.marshfieldclinic.org/genetics>). We screened three affected individuals (patients 1, 2, and 3) for mutations in seven genes (*THBS1*, *FSIP1*, *VSP39*, *MEIS2*, *DLL4*, *CHAC1*, and *CHST14*) among 109 known genes within the 7.3-Mb candidate locus. After identifying a mutation, we only screened *CHST14* (NM_130468.3) in the remaining individuals (patients 4, 5, and 6). We amplified genomic DNA by PCR using four primer sets (sequences available on request). Nucleotide changes found in the patients were checked in 376 Japanese control samples (752 alleles). Compound heterozygosity was confirmed by direct sequencing of the patients' parents or allele specific sequencing after cloning of respective regions covering two different mutation sites of *CHST14*. Nucleotide numbering reflects cDNA numbering with +1 corresponding to the A of the ATG translation initiation codon in the reference sequence, according to journal guidelines (www.hgvs.org/mutnomen). The initiation codon is codon 1.

Primary Fibroblast Culture

We obtained skin fibroblasts from patient 1 at age 6 years, her mother at 27 years, and patient 3 at 29 years. Their age- and sex-adjusted normal controls (a 6-year-old girl: control 1 and a 36-year-old man: control 2) were purchased from Japan Health Sciences Foundation (<http://www.jhsf.or.jp/>). Cells were cultured in Dulbecco's modified Eagle's medium with 10% heat-inactivated fetal bovine serum (FBS), 100 U/ml penicillin, 100 U/ml streptomycin, and 2 mM L-glutamine (Invitrogen, Carlsbad, CA).

Sulfotransferase Assays

COS-7 cells transiently transfected with the N¹-V5-D4ST1 vectors using FuGENETM 6 (Roche Diagnostics, Indianapolis, IN) and human fibroblasts were lysed with 200 μ l of M-PER[®] mammalian protein extraction reagent (Thermo Fisher Scientific Inc., Waltham, MA). Sulfotransferase activities of each cell lysate toward dermatan, chemically desulfated DS were assayed as described before [Mikami et al., 2003]. The [³⁵S]sulfate incorporation was quantified by determination of the radioactivity in the flow-through fractions of the gel filtration chromatography by liquid scintillation counting.

Disaccharide Composition Analysis of Chondroitin (CS)/DS Chains Isolated from Fibroblasts

Cell lysates of fibroblasts cultured on 100-mm plates were digested with actinase E (Kaken Pharmaceutical Co., Ltd., Tokyo, Japan) and GAG-peptides were recovered by 80% ethanol as described [Uyama et al., 2006]. After being desalted using a centrifugal filter, Amicon[®] Ultra-4 (Ultracel-3k, Millipore Corp., Billerica, MA), the GAG-peptides were digested with CSase ABC from *Proteus vulgaris* (EC 4.2.2.20) (Seikagaku Corp., Tokyo, Japan) [Yamagata et al., 1968], a mixture of CSase AC-I from *Flavobacterium heparinum* (EC 4.2.2.5) (Seikagaku Corp.) [Yamagata et al., 1968], and AC-II from *Arthrobacter aureus* (EC 4.2.2.5) (Seikagaku Corp.) [Hiyama and Okada, 1975], or CSase B from *Flavobacterium heparinum* (EC 4.2.2.19) (IBEX Technologies, Kawasaki, Japan) [Michelacci and Dietrich, 1974]. The digests were labeled with a fluorophore 2-aminobenzamide (2AB) and aliquots

of the 2AB-derivatives of CS/DS oligosaccharides were analyzed by anion-exchange HPLC on a PA-03 column (YMC Co., Kyoto, Japan) as previously described [Kinoshita and Sugahara, 1999].

Immunoblotting

Each cell lysate of COS-7 cells expressing the recombinant N'-V5-D4ST1 was subjected to SDS-PAGE using a 10–20% SDS-polyacrylamide gradient gel. The serum-free conditioned medium from fibroblast cultures was collected and concentrated using Amicon Ultra-4 filters (Ultracel-30k). An aliquot of the sample was digested with CSase ABC, CSase AC, CSase B, or buffer alone, and each digest was subjected to SDS-PAGE using a 7.5% SDS-polyacrylamide gel. Immunoblotting was carried out using anti-V5 antibody (Invitrogen) for D4ST1-transfected cells or antihuman decorin antibody (clone 115402; R&D Systems, Minneapolis, MN) for secreted decorin DS proteoglycan from fibroblasts.

PAI1 and SMAD7 Expression Analysis with TGF- β 1 Stimulation

Fibroblasts were grown to 70–80% confluence in 24-well multiplates and transfected with 200 ng of empty, wild-type, or mutant *CHST14* expression vector, using FuGENE 6. At 24 hr after transfection, cells were transferred to low serum medium (0.2% FBS) and, after 8 hr treated with TGF- β 1 (1 ng/ml; PeproTech Inc., Rocky Hill, NJ) for 24 hr. Total RNA was extracted using the SV-Total RNA Isolation system (Promega, Madison, WI). Randomly primed cDNA was synthesized using the Taqman Multiscribe Reverse Transcriptase kit (Applied Biosystems). Real-time PCR was carried out on the StepOnePlus Real-Time PCR system (Applied Biosystems) using the QuantiTect SYBR Green PCR kit (Qiagen, Tokyo, Japan). The following primers were used for amplification: *CHST14*, 5'-CTATGAGAGGCTGGAGGCTG-3' and 5'-AGGCAAAGAGGGAGAAGTCC-3'; *PAI1*, 5'-CAGACCAAGAGCCCTCTCCAC3' and 5'-GACTGTTCC-TGTGGGGTTGT-3'; *SMAD7*, 5'-TTGCTGTGAATCTTACGGGA-3' and 5'-CCAGATAATTCGTTCCCCCT-3'; and *GAPDH*, 5'-ACCA-CAGTCCATGCCATCAC-3' and 5'-TCCACCACCCTGTGCTGTGA-3'.

Reporter Gene Assay

Fibroblasts were grown to 70–80% confluence in 24-well multiplates and transfected with plasmid DNA mixtures using FuGENE 6. The DNA mixture involved 100 ng of SBE4-luc vector, 200 ng of empty, wild-type, or mutant *CHST14* expression vector, and 25 ng of a reference vector, pRL-TK. The SBE4-Luc vector is a TGF- β -responsive reporter containing four tandem copies of a SMAD-binding element (SBE) linked to luciferase [Zawel et al., 1998]. At 24 hr after transfection, cells were transferred to low serum medium (0.2% FBS) and, after 8 hr, treated with TGF- β 1 (1 ng/ml) for a further 24 hr. Luciferase activities were measured using the PG-DUAL-SP reporter assay system (Toyo Ink., Tokyo, Japan) and a Lumat LB 9507 luminometer (Berthold Technologies GmbH & Co. KG, Bad Wildbad, Germany). Relative luciferase activity was calculated by normalizing the transfection efficiency of Renilla luciferase activity against the reference vector.

SMAD2 Phosphorylation with TGF- β 1 Stimulation

Fibroblasts were grown to 70–80% confluence in six-well multiplates and were transfected with 1.5 μ g of empty, wild-type, or mutant *CHST14* expression vector, using FuGENE 6. At 24 hr after transfection, cells was transferred to low serum medium

(0.2% FBS) and, after 8 hr, were treated with TGF- β 1 (1 ng/ml) for 30 min. Cells were lysed using M-PER protein extraction kits (Pierce, Rockford, IL) containing a protease inhibitor cocktail (Roche Diagnostics). Proteins in the cell lysate were separated on SDS-PAGE gels and electrophoretically transferred to PVDF membranes. After blocking with 5% nonfat dry milk in PBS-Tween, the membranes were incubated first with antibodies against phospho-SMAD2 or SMAD2 (Cell Signaling Technology, Danvers, MA) and then with goat polyclonal antibodies against rabbit IgG conjugated with horseradish peroxidase (Cell Signaling Technology). Band intensities were measured using ImageJ software (<http://rsbweb.nih.gov/ij/>).

Pathology

Skin specimens were obtained from the upper arms of patients 5 and 6. For light microscopy using a BX51 microscope (Olympus, Tokyo, Japan), skin specimens were fixed with 20% buffered neutral formalin solution and 3–5- μ m sections were stained with H&E. For transmission electron microscopy, skin specimens were fixed in 2.5% glutaraldehyde for 2 hr, postfixed in 1% osmium tetroxide for 2 hr, dehydrated in a graded ethanol series, and embedded in epoxy resin (Epon 812, TAAB, Berks, UK). Semithin sections (4 μ m) were stained with toluidine blue. Ultrathin sections (100 nm) were stained with uranyl acetate and lead citrate and examined with a transmission electron microscope (JEM-1011, JEOL Ltd., Tokyo, Japan).

Statistical Analyses

All values are described as mean \pm SEM. Where appropriate, we assessed between-groups effects by unpaired *t* tests for two groups and ANOVA with Dunnett's adjustment for more than two groups using GraphPad Prism 5 for Windows, version 5.02 (www.graphpad.com).

The detailed methods are described in the Supp. Methods.

Results

Genetic Analysis

We performed homozygosity mapping of two independent consanguineous families (families 2 and 3) and identified the largest 8.15-Mb homozygous region at chromosome 15q14–q15.3 with the maximum LOD Score 2.885, and by using additional microsatellite markers narrowed it down to a 7.3-Mb region (Supp. Fig. S1). Among 109 known genes within this region, the *CHST14* gene (encoding D4ST1) harbored the same homozygous missense mutation (c.842C>T: p.P281L) in the two families and compound heterozygous mutations in the other four (Fig. 1A and B). Mutations include one nonsense (c.205A>T: p.K69X) and three missense mutations (c.842C>T: p.P281L, c.866G>C: p.C289S, and c.878A>G: p.Y293C) occurring at evolutionally conserved amino acids (Fig. 1C). They were absent in 376 Japanese normal controls.

Sulfotransferase Activity

D4ST1 transfers a sulfate group from 3'-phosphoadenosine 5'-phosphosulfate to position 4 of the *N*-acetyl-D-galactosamine (GalNAc) residues of DS, which is abundantly expressed in skin, aortic wall, tendon, and bone [Penc et al., 1998]. Mutant D4ST1 proteins showed significantly decreased sulfotransferase activity towards dermatan (Fig. 2A) regardless of similar expression levels (Fig. 2B). The sulfotransferase activity was also measured in the

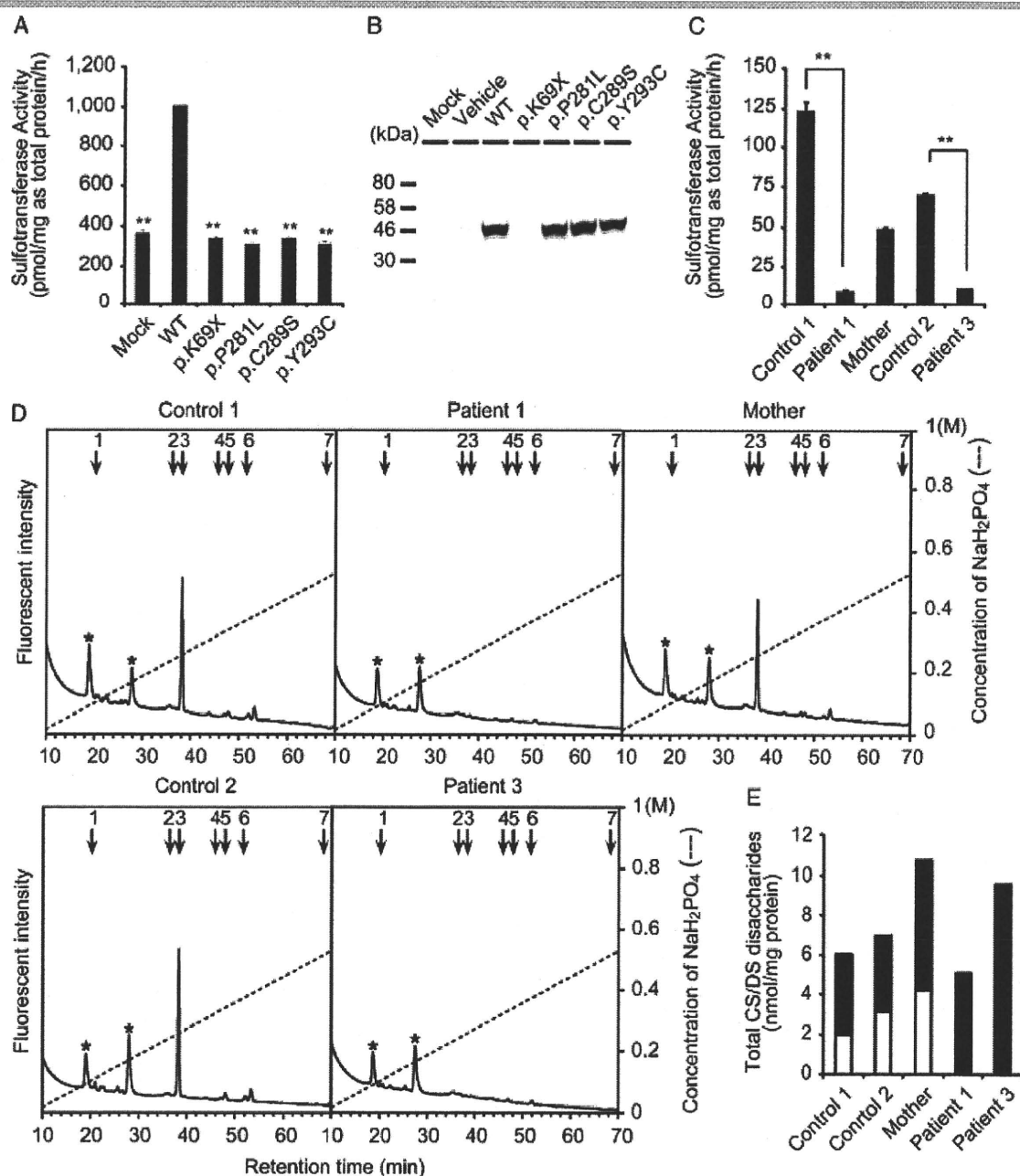


Figure 2. Glycobiological studies. **A:** Sulfotransferase activity of the recombinant D4ST1 of a control and four mutant forms: p.K69X, p.P281L, p.C289S, and p.Y293C. The activity is shown as the incorporation of [³⁵S]SO₄ from the donor [³⁵S]PAPS into the acceptor dermatan. Values are the mean ± SEM (*n* = 3) ***P* < 0.0001 versus WT by one-way ANOVA with Dunnett's adjustment. **B:** Immunoblotting for the exogenous D4ST1 expression in COS-7 cell using the anti-V5 antibody. The mutant p.K69X was too small to be detected (predicted to be approximately 11 kDa). **C:** Sulfotransferase activity toward dermatan in the skin fibroblasts derived from patients 1 and 3, mother of patient 1, and sex- and age-matched healthy controls (controls 1 and 2 for patients 1 and 2, respectively) (mean ± SEM, *n* = 3). ***P* < 0.0001 by two-tailed unpaired *t* test. **D:** Anion-exchange chromatograms of DS disaccharides obtained by CSase B digestion of GAG-peptides from the skin fibroblasts. The elution positions of authentic 2AB-labeled disaccharide standards are indicated by numbered arrows. 1, ΔHexUA-GalNAc; 2, ΔHexUA-GalNAc(6S); 3, ΔHexUA-GalNAc(4S); 4, ΔHexUA(2S)-GalNAc(6S); 5, ΔHexUA(2S)-GalNAc(4S); 6, ΔHexUA-GalNAc(4S,6S); and 7, ΔHexUA(2S)-GalNAc(4S,6S). Asterisks indicate impurities. **E:** The total amounts of CS and DS derived from skin fibroblasts. The total disaccharide contents of CS (black box) and DS (white box) were calculated based on the peak areas in the chromatograms of the digests with CSase AC and CSase B, respectively.

lysates of primary fibroblasts. The mean activity of the fibroblast lysates of the patients was significantly decreased to 6.7% (patient 1) and 14.5% (patient 3) of each age- and sex-matched control 1 and 2, respectively (Fig. 2C). These data indicate mutations in this study result in loss of function.

Glycosaminoglycan (GAG) Chain Analysis

The disaccharide compositions of DS and chondroitin sulfate (CS) in fibroblasts derived from patients and controls were analyzed by anion-exchange HPLC after digestion with

chondroitinase (CSase) B and a mixture of CSases AC-I and AC-II (CSase AC), respectively. CSase B cleaves the *N*-acetyl-D-galactosaminidic linkage in the GalNAc(4S)-L-iduronic acid (IdoUA) (\pm 2S) sequences and yields unsaturated disaccharides, Δ HexUA (4,5-unsaturated hexuronic acid)-GalNAc(4S) and Δ HexUA(2S)-GalNAc (4S) [Sugahara and Mikami, 2007; Yoshida et al., 1993], where 2S and 4S represent 2-*O*-sulfate and 4-*O*-sulfate, respectively. The 4-*O*-sulfation of GalNAc residues is essential for recognition by CSase B. In contrast, CSase AC does not act on IdoUA-containing sequences but degrades the *N*-acetyl-D-galactosaminidic linkages in the GalNAc (\pm 4S, \pm 6S)-D-glucuronic acid (GlcUA) sequences [Linhardt et al., 2006], where 6S stands for 6-*O*-sulfate. When digested with CSase B, 4-*O*-sulfated unsaturated disaccharides were detected only in the controls but not in the patients (Fig. 2D and Supp. Table S1), suggesting that D4ST1 is the major enzyme for 4-*O*-sulfation of DS in normal human skin fibroblasts and cannot be compensated functionally either by chondroitin 4-*O*-sulfotransferase 1 or 2.

To investigate whether the loss-of-function mutations in D4ST1 lead to an increase in nonsulfated dermatan, the GAG fraction prepared from the fibroblasts of the patients and controls were digested with CSase ABC, which cleaves all the *N*-acetyl-D-galactosaminidic linkages in CS/DS chains including the nonsulfated GalNAc-IdoUA sequence that is resistant to CSase B [Yoshida et al., 1993]. Notably, the proportion of the nonsulfated disaccharide Δ HexUA-GalNAc in the patients was very low (Supp. Table S1), suggesting that nonsulfated dermatan was negligible. Conversion of GlcUA to IdoUA most likely occurs even in the patients. However, the epimerization reaction is reversible that favors the GlcUA formation, and D4ST1 probably functions as the “4-*O*-sulfation lock” of IdoA [Malmström, 1984]. Hence, the defect in D4ST1 probably allowed back epimerization reactions converting IdoUA to GlcUA to form CS in the patients. The total amount of CS/DS disaccharides did not show a significant difference between the patients and controls (Fig. 2E). CS disaccharides produced by digestion with CSase AC increased markedly in the cell lysates of fibroblasts from the patients compared with those from the controls, suggesting the upregulation of CS chain biosynthesis (diverted from DS) in the patients.

Glycosaminoglycan of Decorin

We then examined decorin as a major DS proteoglycan in skin. Decorin directly binds to collagen via its core protein and the GAG side chains aggregate and function as interfibrillar bridges [Scott, 1996, 2003]. Decorin purified from the fibroblasts of the patients was resistant to CSase B, indicating that its GAG side chain consisted of only CS but no DS disaccharides, while decorin in the controls contained a hybrid CS/DS chain (Fig. 3A). Actually, the GAG side chain of decorin from the controls was mainly composed of DS disaccharides (approximately 95%), IdoUA-GalNAc(4S), and IdoUA(2S)-GalNAc(4S), but contained a small portion of CS disaccharide, GlcUA-GalNAc(4S) (Fig. 3B). In contrast, no DS disaccharides were detected in the decorin proteoglycan from the patients (Fig. 3B). Its GAG side chain most likely consisted of only CS disaccharides including 4-*O*-sulfated and 6-*O*-sulfated disaccharide units. These data and those shown in Figure 2E suggest that the DS chain of decorin proteoglycan in the patients has been replaced by CS.

TGF- β Signaling Is Unaltered in Fibroblasts Derived from Patients

DS proteoglycan is known to be related to TGF- β signaling [Hocking et al., 1998; Tiedemann et al., 2005; Yamaguchi et al., 1990]. As decorin neutralizes TGF- β 1 activity [Yamaguchi et al., 1990], we examined whether *CHST14* aberrations would affect TGF- β 1 signal transduction. We overexpressed wild-type or mutant *CHST14* cDNA in the fibroblasts from patients in which endogenous *CHST14* expression was negligible and found that the levels of *CHST14* expression from transgenes were approximately 300-fold those of endogenous mutant *CHST14* expression at 56 hr after transfection (Supp. Fig. S2A). To investigate whether *CHST14* overexpression would affect TGF- β signaling in the patient fibroblasts, we performed three different experiments. First, we measured the expression of *PAI1* and *SMAD7*, which are direct downstream targets of TGF- β . The expression of *PAI1* and *SMAD7* was significantly upregulated at 24 hr after TGF- β 1 treatment. However, the degree of upregulation was not different between the cells transfected with empty or *CHST14* expression

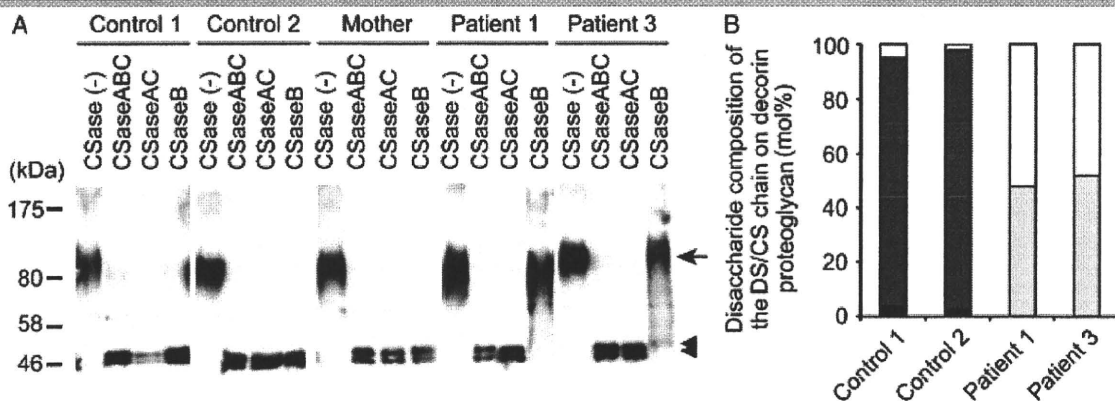


Figure 3. Disaccharide composition analysis of CS/DS chains of decorin. **A:** Immunoblot of decorin proteoglycans. Each serum-free conditioned medium of skin fibroblasts was digested with CSase ABC, CSase AC, CSase B, or buffer alone (–) and then subjected to Western blotting using an antihuman decorin antibody. The arrow and arrowheads indicate the decorin proteoglycans and the core protein of decorin without a CS/DS side chain, respectively. **B:** Proportion of the disaccharide units in the CS/DS hybrid chain in decorin proteoglycans secreted by the fibroblasts. White, light gray, dark gray, and black boxes are GlcUA-GalNAc(4S), GlcUA-GalNAc(6S), IdoUA-GalNAc(4S), and IdoUA(2S)-GalNAc(4S), respectively.

vectors (Supp. Fig. S2B). Second, TGF- β 1 transmits signals to the nucleus via phosphorylation of SMAD2/3 proteins. Therefore, a reporter gene assay was performed using SBE4-luc vector, a TGF- β -responsive reporter containing four tandem copies of the SMAD-binding element (SBE). *CHST14* overexpression did not affect the reporter activity upregulated by TGF- β 1 treatment (Supp. Fig. S2C). Third, we examined the level of phosphorylated SMAD2 proteins by Western blot analysis. Although TGF- β 1 treatment clearly stimulated the phosphorylation of SMAD2, there were no significant differences between the cells transfected with empty or *CHST14* vectors (Supp. Fig. S2D). The same results from these three experiments were obtained using fibroblasts from another patient (data not shown). Furthermore, we confirmed that overexpression of the mutant D4ST1 enzymes identified in patients did not affect TGF- β signaling in normal fibroblasts (data not shown). These results imply that TGF- β signaling might not be changed in the fibroblasts from patients with a D4ST1 deficit.

Histopathological Examination

Hematoxylin and eosin (H&E)-stained light microscopy on patients' skin specimens showed that fine collagen fibers were

predominant in the reticular to papillary dermis and normally thick collagen bundles were markedly reduced (Fig. 4). Electron microscopy showed that the collagen fibrils were dispersed in the reticular dermis, compared with regularly and tightly assembled ones observed in the control, whereas each collagen fibril was smooth and round, not varying in size and shape, similar to each fibril of the control (Fig. 4).

Discussion

In this study, we identified a total four *CHST14* mutations (three missense and one nonsense) in six Japanese patients presenting with a new type of EDS [Kosho et al., 2010]. This disorder represents all hallmarks of EDS: skin hyperextensibility, joint hypermobility, and tissue fragility affecting skin, ligaments, joints, blood vessels, and internal organs [Steinmann et al., 2002]. Tentatively we categorized patients 1 and 2 before as EDS VIB, a subtype of kyphoscoliosis type without lysyl hydroxylase deficiency, based on cutaneous (hyperextensibility, bruisability, fragility with atrophic scars) and skeletal (generalized joint laxity, kyphoscoliosis, Marfanoid habitus) features as well as mild delay of motor development with hypotonia in infancy [Kosho et al.,

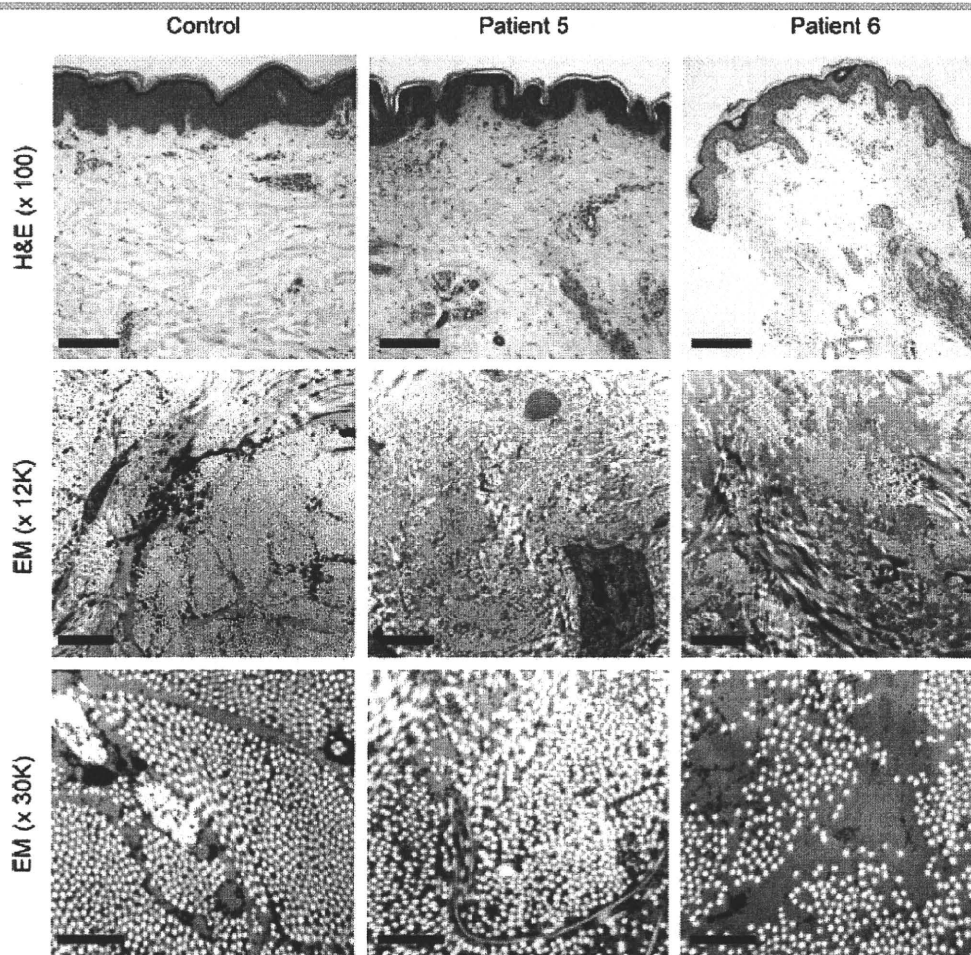


Figure 4. Pathological examination of dermal tissues from patients. H&E-stained light microscopy (upper picture) on skin specimens of patients 5 and 6 ($\times 100$) shows that fine collagen fibers were present predominantly in the reticular to papillary dermis with marked reduction of normally thick collagen bundles. Electron microscopy (EM) ($\times 12,000$ middle picture; $\times 30,000$ lower picture) showed that collagen fibrils were dispersed in the reticular dermis, compared with the regularly and tightly assembled ones observed in the control subject. However, each collagen fibril was smooth and round, not varying in size, similar to the control. Scale bars indicate 500 μ m (upper picture), 2 μ m (middle), and 1 μ m (lower).

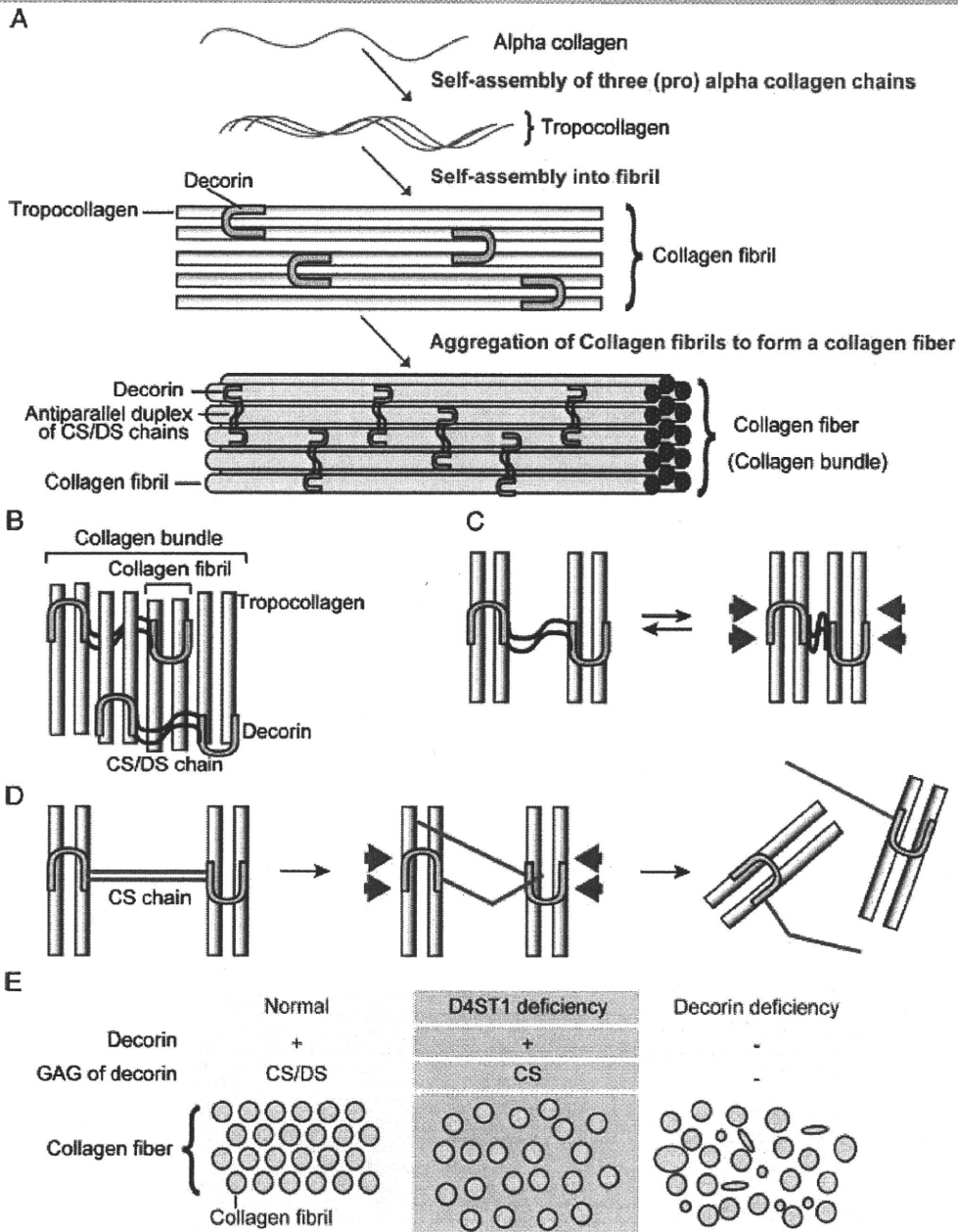


Figure 5. Putative model for abnormal collagen bundle assembly in this disease. **A:** The scheme of the collagen biosynthesis. Three alpha collagen chains are self-assembled into form a tropocollagen unit. Tropocollagens are packed via decorin to form a collagen fibril. Decorin core protein directly binds to collagen's particular amino sequence. Collagen fibrils assemble to form the collagen fiber, also called a collagen bundle when thick. **B:** The relationship between collagen fibril and decorin proteoglycan. Collagen fibrils are assembled into a collagen bundle by the antiparallel complex of the CS/DS hybrid GAG chains of decorin proteoglycan like a bridge to keep the space between fibrils and make collagen fiber tighter. **C, D:** The structural alteration of collagen fibers by mechanical compression in normal and affected states, respectively. CS/DS hybrid chains are able to bend against mechanical compression and rebound to form the original structure, because GAG chains function as suspension (C). D4ST1 defects result in absence of CS/DS hybrid chains in decorin (D). Replaced CS chains might not form collagen bundles properly. Even if they can form bundles, decorin CS chains cannot resist mechanical stresses and collagen fibrils get scattered after repetitive stresses irreversibly, as observed in patients (D). Red arrows indicate mechanical pressures. **E:** Comparisons of collagen fiber formation in normal (left), D4ST1 deficiency (middle), and decorin (core protein) deficiency (right). Collagen fibrils normally aggregate in line and form collagen fibers. The fibrils are round and uniform. In D4ST1 deficiency, collagen fibrils are scattered, although the shape and size of collagen fibrils are unchanged. Irregular shapes and sizes of collagen fibrils were seen in decorin null mice [Danielson et al., 1997], being apparently different from those in D4ST1 deficiency.

2005]. Patients in our series [Kosho et al., 2010] shared many clinical features with a Pakistani sister and brother reported by Steinmann et al. [1975], including down-slanting palpebral

fissures, high palate, talipes equinovarus, progressive talipes vagus and planus, joint laxity, scoliosis, skin hyperextensibility, bruisability, and fragility with atrophic scars; hyperalgesia to pressure,

radiologically identified tall vertebral bodies and diaphysal narrowing of phalanges, metacarpals, and metatarsals; and delayed motor development. The sibs have been classified into EDS VIB [Steinmann et al., 2002], with the lysyl hydroxylase activity proved to be normal [Wenstrup et al., 1989]. Pathological findings were similar to those observed in our series: light microscopically, collagen bundles were interspersed with filamentous material that stained only faintly; and, electron microscopically, a great proportion of collagen fibrils was not integrated into bundles but dispersed into the ground substance [Steinmann et al., 1975].

CHST14 has recently been demonstrated as the causative gene for adducted thumb–clubfoot syndrome (ATCS; MIM# 601776), another autosomal recessive disorder [Dündar et al., 2009]. ATCS has been categorized as a new type of arthrogryposis, based on characteristic clinical pictures from birth to early childhood, including adducted thumbs and clubfoot as well as craniofacial dysmorphism (broad or bossed forehead, brachycephaly, late-closing large fontanelle, hypertelorism, downslanting palpebral fissures, blue sclerae, low-set posteriorly rotated or dysplastic ears, high or cleft palate, short neck), arachnodactyly with tapering fingers, cryptorchidism, inguinal hernia, atrial septal defect, kidney defects, cranial ventricular enlargement, and psychomotor retardation [Dündar et al., 1997, 2001; Janecke et al., 2001; Sonoda and Kouno, 2000]. In a recent study by Dündar et al. [2009], ATCS has been again categorized as a connective tissue disorder, based on additional clinical pictures from childhood to adolescence, including skin fragility, bruisability, and translucency; joint laxity, and osteopenia. In 11 ATCS patients from four families identified to date [Dündar et al., 2009], 5 died in early infancy or childhood: a male from a Austrian family (p.R213P) died shortly after birth due to respiratory failure [Janecke et al., 2001]; a female from a Turkish family (p.V49X) died at age 6 years [Dündar et al., 1997], and a female and two males from a Turkish family (p.[R135G]+[L137Q]) died before 4 months of age [Dündar et al., 2001]. ATCS patients may have wider and more severe manifestations than our series, implicating roles for DS not only in connective tissue maintenance but also in embryonic development [Dündar et al., 2009]. Furthermore, a skin specimen from a patient was interpreted as showing normal structure and ultrastructure [Dündar et al., 2009]. To date, it would be difficult to delineate whether a new type of EDS we have proposed and ATCS would be distinct clinical entities or a single clinical entity (D4ST1 deficiency) with variable inter- and intrafamilial expressions and with different presentations depending on the patients' ages at diagnosis. Longitudinal clinical information of ATCS patients would solve this issue.

Collagen bundles are composed of many collagen fibrils linked via antiparallel CS/DS chain complexes (Fig. 5A and B). The property of reversible deformation of proteoglycan comes from antiparallel anionic GAG chain formation [Scott, 2003]. Interestingly, the structure of DS chains is flexible, whereas that of CS chains is rigid, because L-IdoUA residues in DS can switch readily between almost equi-energetic 1C_4 , 2S_0 , and 4C_1 conformers; by comparison GlcUA in CS adopts purely the 4C_1 conformation [Casu et al., 1988; Catlow et al., 2008]. Transition from the CS/DS hybrid chain of decorin to a CS chain probably decreases the flexibility of the GAG chain and break the GAG antiparallel complex after compression stresses (Fig. 5C and D). This irreversible event could explain the progressive course of this disease. In the patients, the size and shape of the collagen fibrils seemed normal, whereas the collagen bundles were not properly organized. In decorin null mice, dermal collagen fibrils showed huge varieties of size and shape [Danielson et al., 1997]. These

findings suggest that the core protein of decorin is important for collagen fibril formation, and that the CS/DS hybrid chain of decorin proteoglycan regulates the space between the collagen fibrils and forms collagen bundles as reported previously (Fig. 5E) [Scott, 1995]. Decorin is able to neutralize the growth-stimulatory activity of TGF- β 1 in Chinese hamster ovary cells through the interaction of TGF- β 1 with decorin via its core protein, not via a GAG chain [Yamaguchi et al., 1990]. In our transfection experiments with *CHST14* cDNAs, no significant effects on TGF- β signaling between wild-type and mutants were detected, implying the importance of the decorin core protein (not GAG) for TGF- β signaling (Supp. Fig. S2).

In conclusion, we have detected *CHST14* mutations causing a new type of EDS with distinct craniofacial characteristics, multiple congenital contractures, progressive joint and skin laxity, and multisystem fragility related manifestations. Abnormal collagen bundle formation would be a main pathology associated with a decorin GAG abnormality. Decorin GAG side chains in the patients consist of only CS but no DS disaccharides. CS/DS hybrid chains are more flexible than CS chains. Collagen bundles bound by CS chains instead of CS/DS chains in patients should be more fragile than those in controls. These findings underscore the important aspects of decorin proteoglycans in the extracellular matrix and provide new insights for human connective tissue disorders.

Acknowledgments

We thank all the patients and their families for participating in this work. We also thank Ms. Y. Yamashita, Ms. T. Taniguchi, and Dr. S. Tominaga for their technical assistance, Drs. Y. Igawa, T. Miyahara (Shinshu University School of Medicine), Y. Seta, Y. Toki, K. Ono, T. Kosuda, A. Inoue (Tomioka General Hospital, Tomioka), T. Miyamoto (Meido Eye Clinic), K. Arai (Komachi Clinic), T. Shimizu (Gunma Spine Center), F. Miura, H. Ikei (Saku Central Hospital), N. Hoshino (Minami-Aiki-Mura Clinic), T. Muneta (Tokyo Medical and Dental University), N. Kurosawa (Tsuchiura Kyodo General Hospital), M. Nagasaka, M. Kato (Aichi Prefectural Colony Central Hospital), M. Kohyama (JA Hiroshima General Hospital), and T. Hattori (Aichi Children's Health and Medical Center) for providing clinical information of the patients. Dr. A. Kosaku (Dokkyo Medical University) kindly performed the electron microscopy. Grant sponsor: the Ministry of Education, Culture, Sports, Science and Technology; grant numbers: 21790341 (to N. Miyake), B-20390019 (to K.S.) and A-21249024 (to N. Matsumoto). Grant sponsor: the Japan Science and Technology Agency; grant number: 2014 (to N. Matsumoto). Grant sponsor: the Ministry of Health, Labour and Welfare, Japan; grant numbers: 2141039040 (to T.K. and N. Matsumoto), 20-S-3 (to S.I.). Grant sponsors: the Shinshu Association for the Advancement of Medical Sciences (to T.K.); a Grant-in-Aid for Exploratory Research of Young Scientists, Shinshu University (to T.K.); the Takeda Science Foundation (to N. Miyake); the Yokohama Foundation for Advancement of Medical Science; grant number: W2105 (to N. Miyake). Grant sponsor: the Hayashi Memorial Foundation for Female Natural Scientists; grant number: 09R198 (to N. Miyake). Grant sponsor: the Human Frontier Science Program; grant number: RGP0018/2005-C (to K.S.). This work has been done at Advanced Medical Research Center, Yokohama City University.

References

- Beighton P, De Paepe A, Steinmann B, Tsipouras P, Wenstrup RJ. 1998. Ehlers-Danlos syndromes: revised nosology, Villefranche, 1997. Ehlers-Danlos National Foundation (USA) and Ehlers-Danlos Support Group (UK). *Am J Med Genet* 77:31–37.
- Casu B, Petitou M, Provasoli M, Sinay P. 1988. Conformational flexibility: a new concept for explaining binding and biological properties of iduronic acid-containing glycosaminoglycans. *Trends Biochem Sci* 13:221–225.

- Catlow KR, Deakin JA, Wei Z, Delehedde M, Fernig DG, Gherardi E, Gallagher JT, Pavao MS, Lyon M. 2008. Interactions of hepatocyte growth factor/scatter factor with various glycosaminoglycans reveal an important interplay between the presence of iduronate and sulfate density. *J Biol Chem* 283:5235–5248.
- Danielson KG, Baribault H, Holmes DF, Graham H, Kadler KE, Iozzo RV. 1997. Targeted disruption of decorin leads to abnormal collagen fibril morphology and skin fragility. *J Cell Biol* 136:729–743.
- Dündar M, Demiryilmaz F, Demiryilmaz I, Kumandas S, Erkilic K, Kendirci M, Tuncel M, Ozyazgan I, Tolmie JL. 1997. An autosomal recessive adducted thumb-club foot syndrome observed in Turkish cousins. *Clin Genet* 51:61–64.
- Dündar M, Kurtoglu S, Elmas B, Demiryilmaz F, Candemir Z, Ozkul Y, Durak AC. 2001. A case with adducted thumb and club foot syndrome. *Clin Dysmorphol* 10:291–293.
- Dündar M, Muller T, Zhang Q, Pan J, Steinmann B, Vodopjuz J, Gruber R, Sonoda T, Krabichler B, Utermann G, Baenziger JU. 2009. Loss of dermatan-4-sulfotransferase 1 function results in adducted thumb-clubfoot syndrome. *Am J Hum Genet* 85:873–882.
- Hiyama K, Okada S. 1975. Amino acid composition and physicochemical characterization of chondroitinase from *Arthrobacter aurescens*. *J Biochem* 78:1183–1190.
- Hocking AM, Shinomura T, McQuillan DJ. 1998. Leucine-rich repeat glycoproteins of the extracellular matrix. *Matrix Biol* 17:1–19.
- Janecke AR, Unsinn K, Kreczy A, Baldissera I, Gassner I, Neu N, Utermann G, Muller T. 2001. Adducted thumb-club foot syndrome in sibs of a consanguineous Austrian family. *J Med Genet* 38:265–269.
- Kinoshita A, Sugahara K. 1999. Microanalysis of glycosaminoglycan-derived oligosaccharides labeled with a fluorophore 2-aminobenzamide by high-performance liquid chromatography: application to disaccharide composition analysis and exosequencing of oligosaccharides. *Anal Biochem* 269:367–378.
- Kosho T, Miyake N, Hatamochi A, Takahashi J, Kato H, Miyahara T, Igawa Y, Yasui H, Ishida T, Ono K, Kosuda T, Inoue A, Kohyama M, Hattori T, Ohashi H, Nishimura G, Kawamura R, Wakui K, Fukushima Y, Matsumoto N. 2010. A new Ehlers-Danlos Syndrome with craniofacial characteristics, multiple congenital contractures, progressive joint and skin laxity, and multisystem fragility-related manifestations. *Am J Med Genet A* [Epub ahead of print].
- Kosho T, Takahashi J, Ohashi H, Nishimura G, Kato H, Fukushima Y. 2005. Ehlers-Danlos syndrome type VIB with characteristic facies, decreased curvatures of the spinal column, and joint contractures in two unrelated girls. *Am J Med Genet A* 138A:282–287.
- Linhardt RJ, Avci FY, Toida T, Kim YS, Cygler M. 2006. CS lyases: structure, activity, and applications in analysis and the treatment of diseases. *Adv Pharmacol* 53:187–215.
- Malmström A. 1984. Biosynthesis of dermatan sulfate. II. Substrate specificity of the C-5 uronosyl epimerase. *J Biol Chem* 259:161–165.
- Mao JR, Bristow J. 2001. The Ehlers-Danlos syndrome: on beyond collagens. *J Clin Invest* 107:1063–1069.
- Michelacci YM, Dietrich CP. 1974. Isolation and partial characterization of an induced chondroitinase B from *Flavobacterium heparinum*. *Biochem Biophys Res Commun* 56:973–980.
- Mikami T, Mizumoto S, Kago N, Kitagawa H, Sugahara K. 2003. Specificities of three distinct human chondroitin/dermatan N-acetylgalactosamine 4-O-sulfotransferases demonstrated using partially desulfated dermatan sulfate as an acceptor: implication of differential roles in dermatan sulfate biosynthesis. *J Biol Chem* 278:36115–36127.
- Penc SF, Pomahac B, Winkler T, Dorschner RA, Eriksson E, Herndon M, Gallo RL. 1998. Dermatan sulfate released after injury is a potent promoter of fibroblast growth factor-2 function. *J Biol Chem* 273:28116–28121.
- Scott JE. 1995. Extracellular matrix, supramolecular organisation and shape. *J Anat* 187(Pt 2):259–269.
- Scott JE. 1996. Proteodermatan and proteokeratan sulfate (decorin, lumican/fibromodulin) proteins are horseshoe shaped. Implications for their interactions with collagen. *Biochemistry* 35:8795–8799.
- Scott JE. 2003. Elasticity in extracellular matrix “shape modules” of tendon, cartilage, etc. A sliding proteoglycan-filament model. *J Physiol* 553(Pt 2):335–343.
- Sonoda T, Kouno K. 2000. Two brothers with distal arthrogyposis, peculiar facial appearance, cleft palate, short stature, hydronephrosis, retentio testis, and normal intelligence: a new type of distal arthrogyposis? *Am J Med Genet* 91:280–285.
- Steinmann B, Gitzelmann R, Vogel A, Grant ME, Harwood R, Sear CH. 1975. Ehlers-Danlos syndrome in two siblings with deficient lysyl hydroxylase activity in cultured skin fibroblasts but only mild hydroxylysine deficit in skin. *Helv Paediatr Acta* 30:255–274.
- Steinmann B, Royce PM, Superti-Furga A. 2002. The Ehlers-Danlos syndrome. In: Royce PM, Steinmann B, editors. *Connective tissue and heritable disorders*. Hoboken, NJ: John Wiley & Sons Inc. p 431–523.
- Sugahara K, Mikami T. 2007. Chondroitin/dermatan sulfate in the central nervous system. *Curr Opin Struct Biol* 17:536–545.
- Tiedemann K, Olander B, Eklund E, Todorova L, Bengtsson M, Maccarana M, Westergren-Thorsson G, Malmstrom A. 2005. Regulation of the chondroitin/dermatan fine structure by transforming growth factor-beta1 through effects on polymer-modifying enzymes. *Glycobiology* 15:1277–1285.
- Uyama T, Ishida M, Izumikawa T, Trybala E, Tufaro F, Bergstrom T, Sugahara K, Kitagawa H. 2006. Chondroitin 4-O-sulfotransferase-1 regulates E disaccharide expression of chondroitin sulfate required for herpes simplex virus infectivity. *J Biol Chem* 281:38668–38674.
- Wenstrup RJ, Murad S, Pinnell SR. 1989. Ehlers-Danlos syndrome type VI: clinical manifestations of collagen lysyl hydroxylase deficiency. *J Pediatr* 115:405–409.
- Yamagata T, Saito H, Habuchi O, Suzuki S. 1968. Purification and properties of bacterial chondroitinases and chondrosulfatases. *J Biol Chem* 243:1235–1523.
- Yamaguchi Y, Mann DM, Ruoslahti E. 1990. Negative regulation of transforming growth factor-beta by the proteoglycan decorin. *Nature* 346:281–284.
- Yasui H, Adachi Y, Minami T, Ishida T, Kato Y, Imai K. 2003. Combination therapy of DDAVP and conjugated estrogens for a recurrent large subcutaneous hematoma in Ehlers-Danlos syndrome. *Am J Hematol* 72:71–72.
- Yoshida K, Arai M, Kohno Y, Maeyama KI, Myazono H, Kikuchi H, Morikawa K, Tawada A, Suzuki S. 1993. Activity of bacterial eliminases towards dermatan sulphates and dermatan sulphate proteoglycan. In: Scott JE, editor. *Dermatan sulfate proteoglycans: chemistry, biology, chemical pathology*. London: Portland Press. p 55–80.
- Zawel L, Dai JL, Buckhaults P, Zhou S, Kinzler KW, Vogelstein B, Kern SE. 1998. Human Smad3 and Smad4 are sequence-specific transcription activators. *Mol Cell* 1:611–617.

A New Ehlers–Danlos Syndrome With Craniofacial Characteristics, Multiple Congenital Contractures, Progressive Joint and Skin Laxity, and Multisystem Fragility-Related Manifestations

Tomoki Kosho,^{1*} Noriko Miyake,² Atsushi Hatamochi,³ Jun Takahashi,⁴ Hiroyuki Kato,⁴ Teruyoshi Miyahara,⁵ Yasuhiko Igawa,⁶ Hiroshi Yasui,⁷ Tadao Ishida,⁷ Kurahito Ono,⁸ Takashi Kosuda,⁹ Akihiko Inoue,¹⁰ Mohei Kohyama,¹¹ Tadashi Hattori,¹² Hirofumi Ohashi,¹³ Gen Nishimura,¹⁴ Rie Kawamura,¹ Keiko Wakui,¹ Yoshimitsu Fukushima,¹ and Naomichi Matsumoto²

¹Department of Medical Genetics, Shinshu University School of Medicine, Matsumoto, Japan

²Department of Human Genetics, Yokohama City University Graduate School of Medicine, Yokohama, Japan

³Department of Dermatology, Dokkyo Medical University, School of Medicine, Mibu, Japan

⁴Department of Orthopedics, Shinshu University School of Medicine, Matsumoto, Japan

⁵Department of Ophthalmology, Shinshu University School of Medicine, Matsumoto, Japan

⁶Department of Urology, Shinshu University School of Medicine, Matsumoto, Japan

⁷First Department of Internal Medicine, Sapporo Medical University, Sapporo, Japan

⁸Department of Orthopedics, Tomioka General Hospital, Tomioka, Japan

⁹Department of Pediatrics, Tomioka General Hospital, Tomioka, Japan

¹⁰Department of Surgery, Tomioka General Hospital, Tomioka, Japan

¹¹Department of Surgery, JA Hiroshima General Hospital, Hiroshima, Japan

¹²Department of Orthopedics, Aichi Children's Health and Medical Center, Oobu, Japan

¹³Division of Medical Genetics, Saitama Children's Medical Center, Saitama, Japan

¹⁴Department of Radiology, Tokyo Metropolitan Kiyose Children's Hospital, Kiyose, Japan

Received 27 January 2010; Accepted 13 April 2010

We previously described two unrelated patients showing characteristic facial and skeletal features, overlapping with the kyphoscoliosis type Ehlers–Danlos syndrome (EDS) but without lysyl hydroxylase deficiency [Kosho et al. (2005) *Am J Med Genet Part A* 138A:282–287]. After observations of them over time and encounter with four additional unrelated patients, we have concluded that they represent a new clinically recognizable type of EDS with distinct craniofacial characteristics, multiple congenital contractures, progressive joint and skin laxity, and multisystem fragility-related manifestations. The patients ex-

hibited strikingly similar features according to their age: *craniofacial*, large fontanelle, hypertelorism, short and downslanting palpebral fissures, blue sclerae, short nose with hypoplastic columella, low-set and rotated ears, high palate, long philtrum, thin vermilion of the upper lip, small mouth, and micro-retrognathia in infancy; slender and asymmetric face with protruding jaw from adolescence; *skeletal*, congenital contractures of fingers, wrists, and hips, and talipes equinovarus with anomalous insertions of flexor muscles; progressive joint laxity with recurrent dislocations; slender and/or cylindrical fingers and

Additional supporting information may be found in the online version of this article.

Grant sponsor: Research on Intractable Diseases, Ministry of Health, Welfare, and Labor, Japan; Grant Number: #2141039040; Grant sponsor: Shinshu Association for the Advancement of Medical Sciences; Grant sponsor: Grant-in-Aid for Exploratory Research of Young Scientists, Shinshu University.

*Correspondence to:

Tomoki Kosho, M.D., Department of Medical Genetics, Shinshu University School of Medicine, 3-1-1 Asahi, Matsumoto 390-8621, Japan.

E-mail: ktomoki@shinshu-u.ac.jp

Published online 14 May 2010 in Wiley InterScience

(www.interscience.wiley.com)

DOI 10.1002/ajmg.a.33498

progressive talipes valgus and cavum or planus, with diaphyseal narrowing of phalanges, metacarpals, and metatarsals; pectus deformities; scoliosis or kyphoscoliosis with decreased physiological curvatures of thoracic spines and tall vertebrae; *cutaneous*, progressive hyperextensibility, bruiseability, and fragility with atrophic scars; fine palmar creases in childhood to acrogeria-like prominent wrinkles in adulthood, recurrent subcutaneous infections with fistula formation; *cardiovascular*, cardiac valve abnormalities, recurrent large subcutaneous hematomas from childhood; *gastrointestinal*, constipation, diverticula perforation; *respiratory*, (hemo)pneumothorax; and *ophthalmological*, strabismus, glaucoma, refractive errors. © 2010 Wiley-Liss, Inc.

Key words: a new type Ehlers–Danlos syndrome; craniofacial characteristics; multiple congenital contractures; joint laxity; talipes deformities; kyphoscoliosis; skin laxity; multisystem fragility; recurrent subcutaneous hematomas

INTRODUCTION

The Ehlers–Danlos syndrome (EDS) is a heterogeneous group of heritable connective tissue disorders affecting as many as 1 in 5,000 individuals, characterized by joint and skin laxity, and tissue fragility [Steinmann et al., 2002]. The fundamental mechanisms of EDS are known to consist of dominant-negative effects or haploinsufficiency of mutant procollagen α -chains and deficiency of collagen-processing-enzymes [Mao and Bristow, 2001]. In a revised nosology, Beighton et al. [1998] classified EDS into six major types: (1) classical type (OMIM#130000) (causative gene, *COL5A1* or *COL5A2*; affected protein, $\alpha 1(V)$ or $\alpha 2(V)$ procollagen), (2) hypermobility type (OMIM#130020) (*TNXB*; tenascin-XB, in a small subset of cases), (3) vascular type (OMIM#130050) (*COL3A1*; $\alpha 1(III)$ procollagen), (4) kyphoscoliosis type (OMIM#225400) (*PLOD*; lysyl hydroxylase), (5) arthrochalasia type (OMIM#130060) (*COL1A1* or *COL1A2*; $\alpha 1(I)$ or $\alpha 2(I)$ procollagen), and (6) dermatospraxis type (OMIM#225410) (*ADAMTS2*; procollagen I N-proteinase). Additional minor variants of EDS have been identified with molecular and biochemical abnormalities: Brittle cornea syndrome (OMIM#229200) (*ZNF469*) [Abu et al., 2008], EDS-like syndrome due to tenascin-XB deficiency (OMIM#606408) (*TNXB*; tenascin-XB) [Schalkwijk et al., 2001], progeroid form (OMIM#130070) ($\beta 4GALT7$; xylosylprotein 4-beta-galactosyltransferase) [Kresse et al., 1987], cardiac valvular form (OMIM#225320) (*COL1A2*; $\alpha 2(I)$ procollagen) [Schwarze et al., 2004], and EDS-like spondylocheirodysplasia (OMIM#612350) (*SLC39A13*; a membrane-bound zinc transporter) [Giunta et al., 2008].

We previously described two unrelated patients showing characteristic facial and skeletal features, with similarities to kyphoscoliosis type EDS but without lysyl hydroxylase deficiency [Kosho et al., 2005]. After observations of them over time and encounter with four additional unrelated patients including one reported by Yasui et al. [2003], we have concluded that they represent a new clinically recognizable type of EDS characterized by distinct craniofacial features, multiple congenital contractures, progressive

How to Cite this Article:

Kosho T, Miyake N, Hatamochi A, Takahashi J, Kato H, Miyahara T, Igawa Y, Yasui H, Ishida T, Ono K, Kosuda T, Inoue A, Kohyama M, Hattori T, Ohashi H, Nishimura G, Kawamura R, Wakui K, Fukushima Y, Matsumoto N. 2010. A new Ehlers–Danlos syndrome with craniofacial characteristics, multiple congenital contractures, progressive joint and skin laxity, and multisystem fragility-related manifestations.

Am J Med Genet Part A 152A:1333–1346.

joint and skin laxity, and progressive multisystem complications associated with tissue fragility including recurrent large subcutaneous hematomas. Here, we present detailed clinical courses of the six patients to delineate the disorder.

CLINICAL REPORTS

Patient 1

The patient is a now 16-year-old Japanese girl. Part of her history was described previously [Kosho et al., 2005]. She was the first child of a healthy mother and a healthy non-consanguineous father, both 19 years of age. She was born by normal vaginal delivery at 42 weeks of gestation. Her birth weight was 2,724 g (-1.3 SD), length 50.0 cm (-0.1 SD), and OFC 32.5 cm (-1.0 SD). She was admitted for the treatment of hypoglycemia, hyperbilirubinemia, and left talipes equinovarus (Fig. 1J). Her craniofacial features included a large fontanelle, hypertelorism, short and downslanting palpebral fissures, blue sclerae, a short nose with a hypoplastic columella, low-set and rotated ears, a high palate, a long philtrum, a thin upper lip vermilion, a small mouth, and micro-retrognathia (Fig. 1A). She had arachnodactyly, flexion-adduction contractures of bilateral thumbs, flexion contractures of the metacarpophalangeal (MP) and interphalangeal (IP) joints in the other fingers (Fig. 1E,F), and rigidity of bilateral hip joints. She suckled poorly, and was admitted again for the treatment of dehydration at age 1 month. Talipes equinovarus was treated with serial plaster casts, and was surgically corrected at age 2 years. Anomalous insertions of the flexor muscles were observed at the operation. Gross motor development was delayed: she sat at age 10 months and walked unassisted at age 2 years. Her skin was easily torn, but showed normal hemostasis in open wounds. Her face became longer with bushy and arched eyebrows and a pointed chin (Fig. 1B). At age 4 years, she developed a large subcutaneous hematoma over the occiput after falling, followed by acute hemorrhagic anemia that required admission and transfusion of hemostatic agents and packed red cells. During the admission, she was suspected to have EDS. At age 6 years, she developed a large subcutaneous hematoma over the temporoparietal region after falling, requiring admission and intravenous administration of hemostatic agents. She had recurrent dislocations of the shoulders, elbows, and knees.

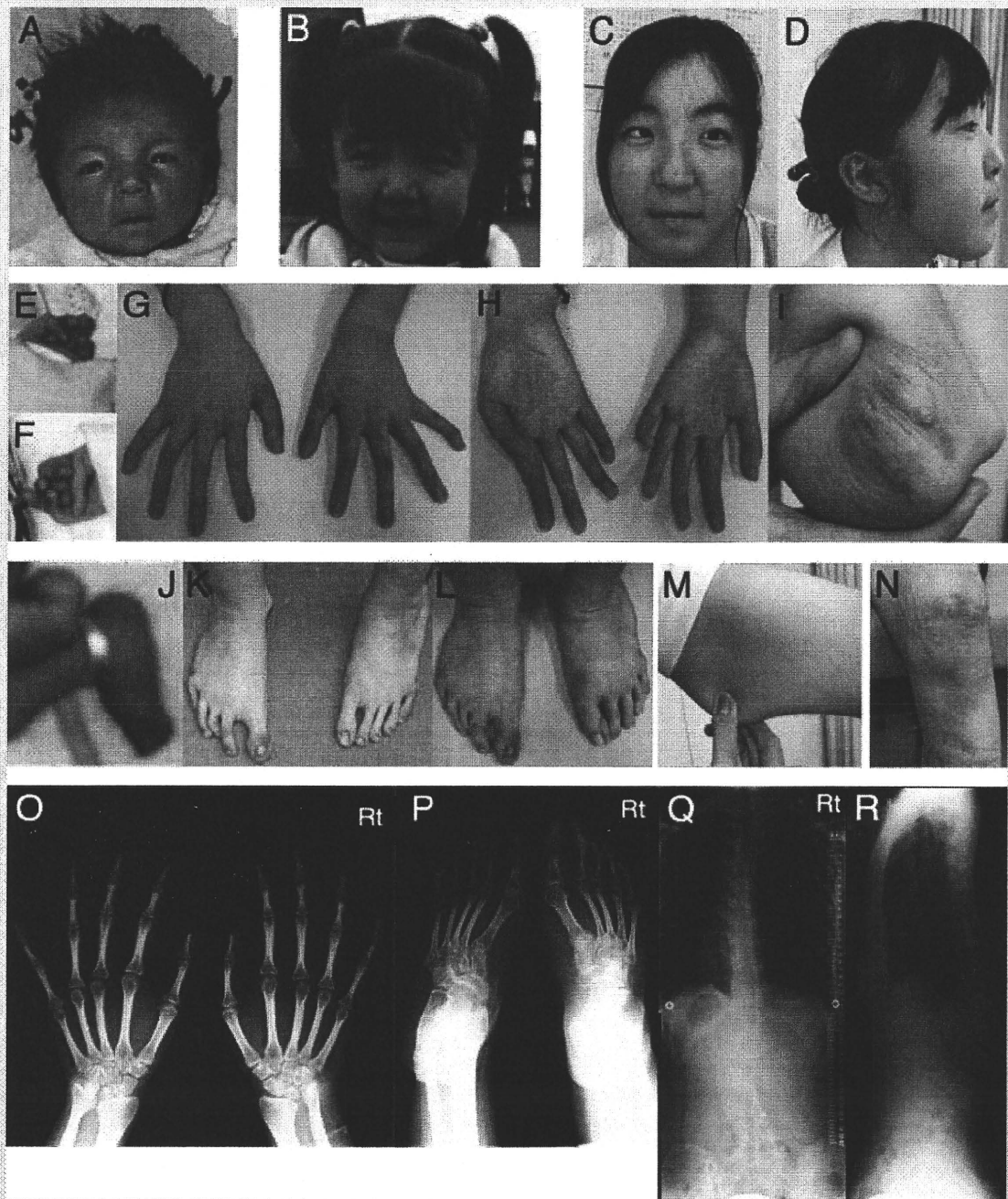


FIG. 1. Patient 1. Clinical photographs of the face at age 23 days (A), 3 years (B), and 16 years (C,D); the left (E) and the right (F) hands at age 23 days; the hands at age 16 years (G,H); the left elbow at age 16 years (I); the feet at birth (J), age 11 years (K), and 16 years (L); the skin on the left upper arm at age 16 years (M); and the left knee at age 16 years (N). Radiographs of the hands (O), the feet (P), and the spine (O,R) at age 16 years. [Color figure can be viewed in the online Issue, which is available at www.interscience.wiley.com.]

When first seen by us at age 7 years, she weighed 19.2 kg (-1.0 SD), height 123.8 cm ($+0.8$ SD), and OFC 51.5 cm (± 0 SD). She had generalized joint laxity, a straight back with scoliosis, and cylindrical and slender fingers. Her skin was hyperextensible, bruisable, and fragile with multiple atrophic scars. Hyperalgesia to pressure such as measuring blood pressure at the upper arms was

noted. Ophthalmological examinations showed microcornea and hyperopia. Otological examinations showed narrow middle ear spaces and hearing impairment of high-pitched sounds. Heart murmurs were not audible, and cardiac ultrasonography showed trivial mitral valve regurgitation. Her bladder was dilated with urinary retention and frequent cystitis, requiring manual pressure

voiding. She developed a large subcutaneous hematoma over the buttock after falling, requiring surgical drainage. Treatment with temporary intranasal administration of 1-desamino-8-D-arginine vasopressin (DDAVP) after injuries was started.

At age 11 years, she weighed 29.5 kg (-2 SD), height 148.1 cm ($+0.2$ SD), and OFC 54 cm ($+0.2$ SD). She had a slender face and a Marfanoid habitus with pectus excavatum and progressing talipes valgus and planus (Fig. 1K). Fine palmar creases and a ganglion on the left foot were noted. Hyperopia had improved and frequency of otitis media had decreased. Urinary retention persisted but manual pressure voiding was not necessary. Heart murmurs were audible, and cardiac ultrasonography showed moderate tricuspid valve regurgitation, prolapse of the tricuspid and mitral valves, and left-to-right shunt via a small atrial septal defect.

At age 12 years, menarche occurred. Kyphoscoliosis progressed with lumbago, necessitating a brace. The ganglion on the left foot was punctured and jellylike contents were suctioned, but soon it swelled again. Persistent urinary incontinence occurred, and urological examinations showed involuntary contractions and hypesthesia of the bladder with normal voiding function. She developed a large subcutaneous hematoma around the head and face after hitting the forehead on a door mirror, with a decline of hemoglobin concentration from 13.4 to 7.7 g/dl in several hours, and was admitted in an intensive care unit (ICU).

She had severe constipation (defecation once a week) and sometimes diarrhea (after having oily or watery foods), treated by medication of *Lactobacillus* and lactulose from age 15 years. Large bowel sounds resembling a frog-croak were frequently heard. At age 16 years, acute gastric ulcer occurred in the antrum, leading to massive hematemesis. Gastric obstruction soon progressed, and treatment with a proton pump inhibitor and intravenous hyperalimentation (IVH) was not effective. A large thrombus developed at the site of inserting an IVH catheter, and an inferior vena cava filter was placed. Distal gastrectomy with Billroth I reconstruction was performed, complicated by a massive hematoma in the rectus sheath necessitating intranasal administration of DDAVP and transfusion of packed red cells and fresh frozen plasma. She attended a class for handicapped children from her junior high school days due to her physical fragility and mild learning disability.

When last seen by us at age 16 years, she weighed 49.4 kg (-0.4 SD), height 159.2 cm ($+0.3$ SD), and OFC 55.8 cm ($+0.2$ SD). Her face was slender with a protruding jaw (Fig. 1C,D). She suffered from lumbago because of progressive kyphoscoliosis. The distal IP joints in bilateral index to little fingers and the IP joints in bilateral thumbs could hardly be flexed or extended (Fig. 1G,H). The proximal IP joints in bilateral index to little fingers and the MP joints in all fingers could be flexed and extended, but could not be moved separately and smoothly (see supporting information Video 1 which may be found in the online version of this article). She had chronic dislocations of bilateral distal radio-ulnar joints and radial heads. Bilateral talipes valgus and planus progressed with extremely soft subcutaneous tissues at the heels (Fig. 1L), resulting in difficulty in walking. The IP joints in bilateral toes could not be moved and metatarsophalangeal joints could only be moved slightly. She had skin redundancy (Fig. 1M) and fragility with atrophic scars (Fig. 1N), fine palmar creases (Fig. 1H), and recurrent subcutaneous infections at the elbows (Fig. 1I) and the buttocks with fistula formation.

Patient 2

The patient is a now 32-year-old Japanese woman. Part of her history was described previously [Kosho et al., 2005]. She was born at term as the third child of a healthy 27-year-old mother and a healthy 37-year-old father, who were first cousins once removed. Her birth weight was 2,500 g (-2 SD) and two elder sisters were healthy. She suckled poorly, and was gavage fed for the first week of life. At age 9 weeks, she was admitted for the treatment of multiple contractures. Her craniofacial features included a large anterior fontanelle, hypertelorism, strabismus, short and downslanting palpebral fissures, a short nose with a hypoplastic columella, low-set and rotated ears, a long philtrum, a thin upper lip vermilion, a high palate, and micro-retrognathia (Fig. 2A). Skeletal features included extension contractures with ulnar deviation of bilateral wrists, flexion-adduction contractures of bilateral thumbs, flexion contractures of the MP joints in bilateral middle to little fingers, extension contractures of the distal IP joints in bilateral index to little fingers (Fig. 2E,F), bilateral talipes equinovarus (Fig. 2I), and a congenital dislocation of the right hip. Active movement of her fingers was poor. Mild skin hyperextensibility was noted. Talipes equinovarus and finger-wrist contractures were treated with serial plaster casts.

At age 1 year and 2 months, she underwent surgical corrections of bilateral talipes equinovarus. Anomalous insertions of the flexor muscles (the tibialis posterior to the talus [normal, navicular and cuneiform bones], the flexor digitorum longus to the abductor hallucis muscle [normal, bases of distal phalanges of four lesser toes], and the flexor hallucis longus to the calcaneus [normal, base of distal phalanx of hallux]) (Fig. 3) were shown to cause talipes equinovarus and inability to flex toes. Tendon sheaths of the muscles were hypoplastic, and surrounding tissues were fragile and hyperextensible. Skin hyperextensibility became more evident with frequent bruises. At age 2 years, dislocation of the right hip was treated through overhead traction and manual reduction under general anesthesia. Her tentative clinical diagnosis was Freeman-Sheldon syndrome.

Gross motor development was delayed: she raised her head at age 4 months, sat unassisted at 7 months, stood up assisted at 2 years and 6 months, cruised furniture at 3 years, and waked alone with short leg braces at 5 years. Generalized hypotonia and joint laxity became evident, and bilateral talipes valgus and cavus progressed (Fig. 2J). Her face became longer with drooping eyelids and bushy eyebrows (Fig. 2B). At age 6 years, dislocations of the left hip and the patella occurred, which were reduced manually. She also developed a subcutaneous hematoma around the left knee that required surgical drainage. She had constipation, and from incontinence and recurrent urinary tract infections associated with an atonic bladder and urinary retention. She had visual impairment from myopia and astigmatism, and impaired hearing for high-pitched sounds. At age 8 years, she developed a wound on the buttocks after falling, followed by bacterial infections and fistula formation, which lead to skin defects including decubitus necessitating plastic surgery. At age 11 years, she fell and developed a large subcutaneous hematoma over the head, requiring surgical drainage and transfusion of packed red cells. At age 12 years, she had dislocation of the left patella and left shoulder after minor injuries, which were

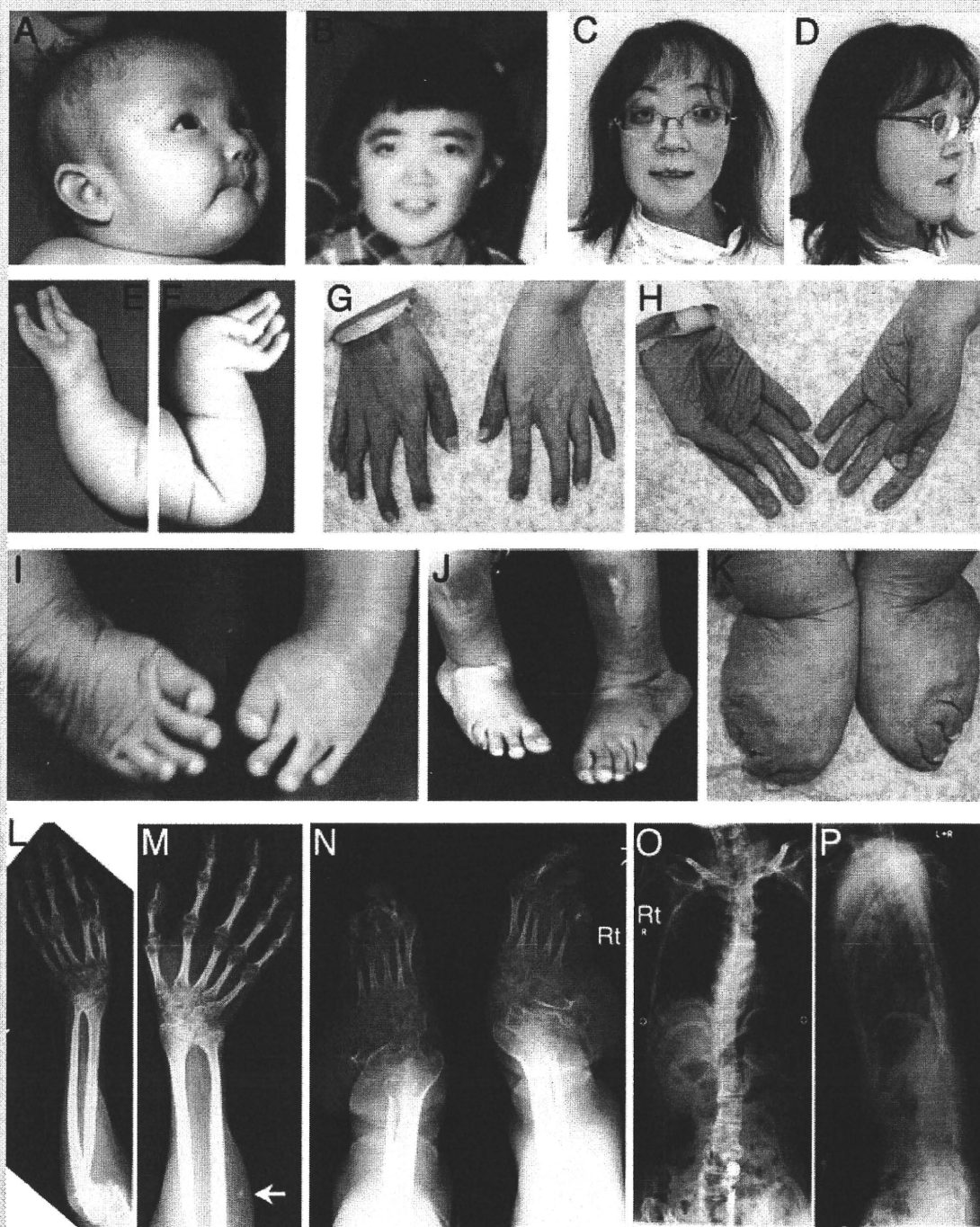


FIG. 2. Patient 2. Clinical photographs of the face at age 3 months [A], 8 years [B], and 28 years [C,D]; the right arm at age 2 months [E]; the left arm at age 3 months [F]; the hands at age 28 years [G,H]; and the feet at age 2 months [I], 6 years [J], and 28 years [K]. Radiographs of the left [L] and the right [M] arms, the feet [N], and the spine [O,P] at age 28 years. An arrow indicates ectopic calcification [M]. [Color figure can be viewed in the online issue, which is available at www.interscience.wiley.com.]

reduced manually. Her IQ was normal at 99. She attended a school for handicapped children because of limitations of daily activities bound to a wheelchair. Her clinical diagnosis was confirmed as EDS. At age 15 years, her height was 140 cm (-3.2 SD). She had marked muscle weakness with grip power 5 kg in the right and 2 kg in the

left. She became exhausted easily. Frequent dislocations of the shoulders and elbows were reduced manually by her. Scoliosis was also noted.

At age 24 years, she underwent surgery for colonic perforation associated with diverticulitis. At the operation, multiple colonic

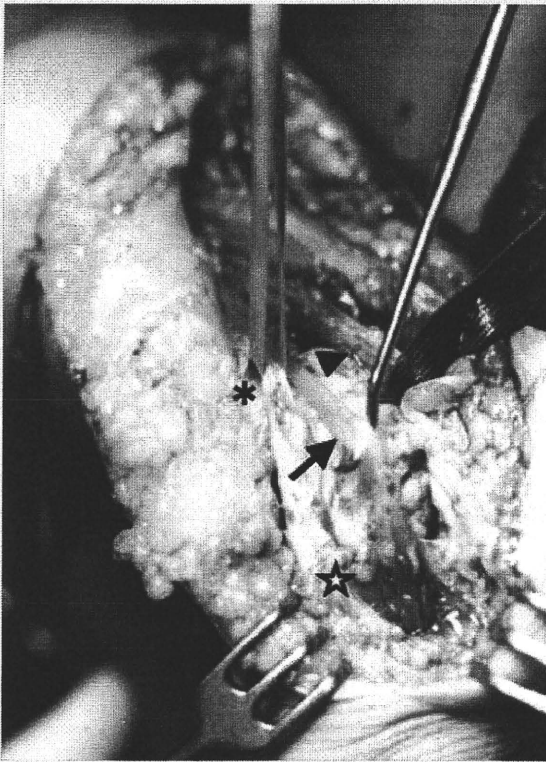


FIG. 3. An operative finding at postero-medial release in the right foot of Patient 2. Anomalous insertions of the flexor muscles are noted: the tibialis posterior muscle (*) to the talus, the flexor digitorum longus muscle (arrow) to the abductor hallucis muscle [blank star], and the flexor hallucis longus muscle (arrowhead) to the calcaneus.

diverticula were observed. At age 26 years, visual impairment progressed due to bilateral glaucoma with an elevation of intraocular pressure (IOP) to 20 mmHg in the right eye and 21 mmHg in the left eye, accompanied by a decreased visual field. At age 28 years, she suffered from a right pneumothorax, treated with chest tube drainage. Acute cystitis also occurred, and cystolithiasis was detected in the follow-up ultrasonography.

When seen by us at age 28 years, she was wheelchair-bound. She had a slender face with a protruding jaw (Fig. 2C,D). Her thorax was flat and thin. Her fingers were cylindrical. She had mild flexion contractures of the distal IP joints in bilateral index to little fingers, mild flexion-adduction contractures of bilateral thumbs (Fig. 2G,H), and chronic dislocations of bilateral distal radio-ulnar joints and the left radial head accompanied by ruptured tendon of the extensor pollicis longus muscle. She also had talipes valgus and cavus with extremely soft subcutaneous tissues at the heels (Fig. 2K). She could rarely flex or extend the left thumb or all toes. Her skin was redundant, bruisable, and fragile with atrophic scars. Prominent wrinkles in the palms showed acrogeria (Fig. 2H). She showed hyperalgesia to pressure such as measuring blood pressure at the upper arms. She had chronic subcutaneous abscesses with fistula formation at the elbows and buttocks. She suffered from

severe constipation (defecation once a week) requiring oral laxatives and *Lactobacillus*, but sometimes had diarrhea. Large bowel sounds resembling a frog-croak were frequently heard. Cardiac ultrasonography showed no structural or functional abnormalities in the mitral or aortic valve.

When last seen by us at age 31 years, she suffered from progressive visual field loss, although IOP was controlled within the range of 14–16 mmHg through topical administration of a prostaglandin F₂ α derivative (latanoprost) and a β -adrenergic receptor blocker (timolol).

Patient 3

The patient, now a 32-year-old Japanese man, was the first child of a healthy 31-year-old mother and a healthy 25-year-old father, who were first cousins. His younger sister was healthy. He was delivered by vacuum extraction at 40 weeks of gestation. His birth weight was 3,300 g (+0.5 SD), length 52.0 cm (+1.3 SD), and OFC 35.0 cm (+0.8 SD). He was admitted for the treatment of orthopedic complications. He had a triangular face with a large skull, a large anterior fontanelle, hypertelorism, short and downslanting palpebral fissures, strabismus, a short nose with a hypoplastic columella, low-set and rotated ears, a long philtrum, a thin upper lip vermilion, a small mouth, and micro-retrognathia (Fig. 4A). His skeletal features included extension contractures of bilateral wrists, flexion-adduction contractures of bilateral thumbs, flexion contractures of the MP joints in bilateral index to little fingers, of the proximal IP joints in the right index and middle fingers, and of the distal IP joint in the left middle finger; extension contractures of the other IP joints (Fig. 4D,E), rigidity of bilateral hip joints, and bilateral talipes equinovarus (Fig. 4H). His skin was redundant with a lot of creases (Fig. 4D,E). Widely spaced nipples and bilateral cryptorchidism were noted. He was diagnosed as arthrogryposis. Talipes equinovarus was treated with serial plaster casts, and was surgically corrected at age 1 year and 8 months. He also underwent tendon transplantations for defects of tendons to bilateral thumbs, and orchiopexy.

At age 6 years, he developed a subcutaneous hematoma over the buttocks after falling, followed by bacterial infections and fistula formation. At age 8 years, a large subcutaneous hematoma occurred spontaneously over the head and progressed acutely with loss of consciousness, treated with emergency surgical drainage and transfusion of packed red cells. Large subcutaneous hematomas occurred in the left elbow at age 11 years, on the right shin followed by a rupture of another vessel, which spread the hematoma from the thigh to the ankle and made him bedridden for a month at age 14 years; on the right ankle necessitating admission at age 19 years, on the left arm making him bedridden for 2 months at age 26 and 27 years, and on the right shin making him bedridden for 2 months at age 29 years. He also suffered from recurrent joint dislocations of the right knee twice at age 9–10 years and the left shoulder five times at age 22–27 years. He was admitted for the treatment of bronchitis, leading to hearing impairment. At age 29 years, a rupture of a small intestine diverticulum was treated with emergency surgery.

When referred to us at age 30 years, his weight was 45 kg (–1.7 SD) and height 178 cm (+1.2 SD). He could walk independently but used a wheelchair when he went out. He had a skull with a

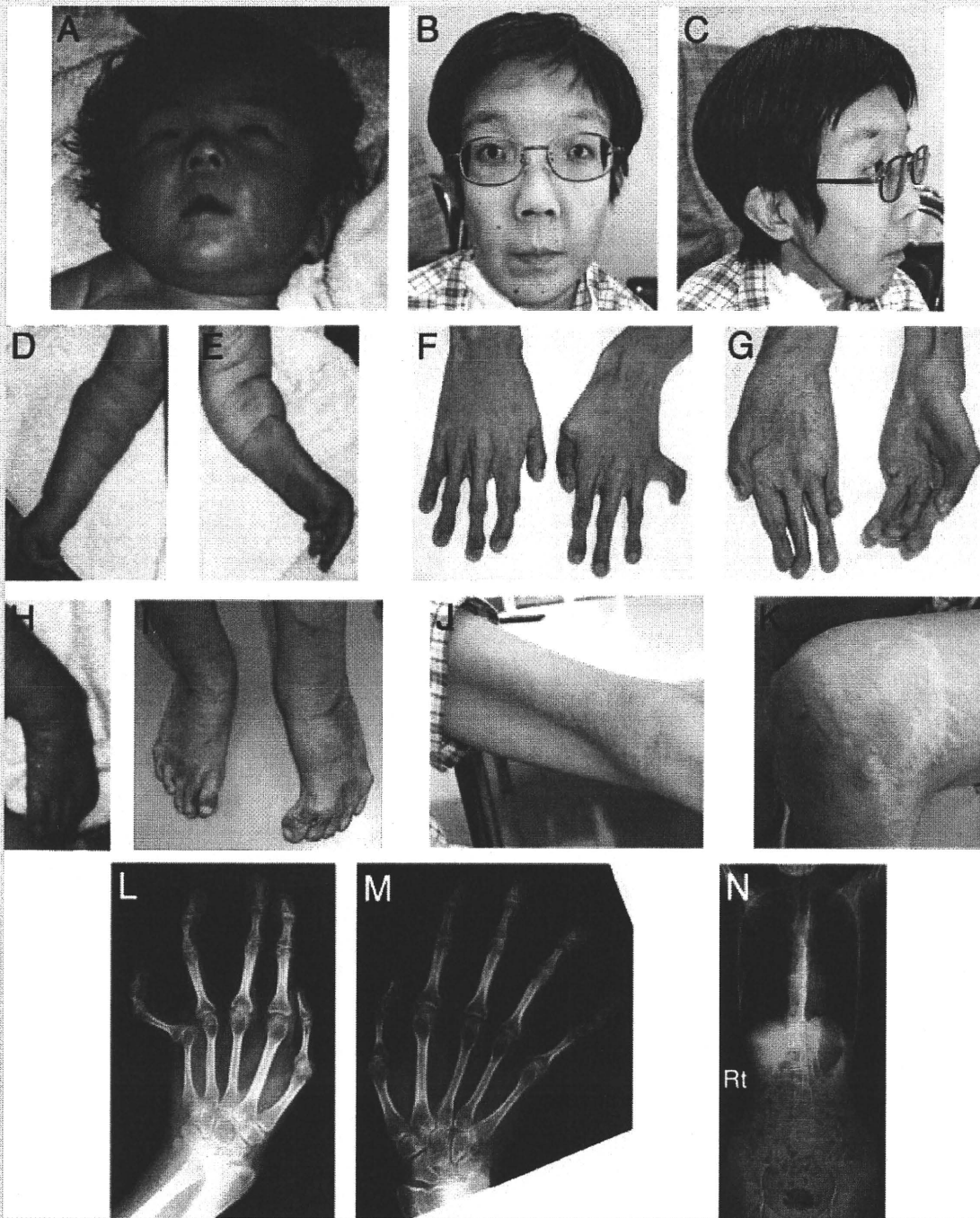


FIG. 4. Patient 3. Clinical photographs of the face in the neonatal period (A) and at age 30 years (B,C); the right (D) and the left (E) arms in the neonatal period; the hands at age 30 years (F,G); the left foot in the neonatal period (H); the feet at age 30 years (I); the skin on the left upper arm at age 30 years (J); and the left knee at age 30 years (K). Radiographs of the left (L) and the right (M) hands, and the spine (N) at age 31 years. [Color figure can be viewed in the online issue, which is available at www.interscience.wiley.com.]

prominent occiput and a lot of white hair, and a slender and asymmetric face with hypertelorism, strabismus, blue sclerae, a short nose with a hypoplastic columella, low-set rotated ears, a long philtrum, a thin upper lip vermillion, a high palate, crowded teeth, and a protruding jaw (Fig. 4B,C). He had a Marfanoid habitus with a

flat and thin thorax and kyphoscoliosis. His fingers were slender and cylindrical. He had contractures of fingers and toes with limited flexion or extension (Fig. 4F,G,I), chronic dislocations of bilateral distal radio-ulnar joints and radial heads, and progressive talipes valgus and planus with extremely soft subcutaneous tissues at the

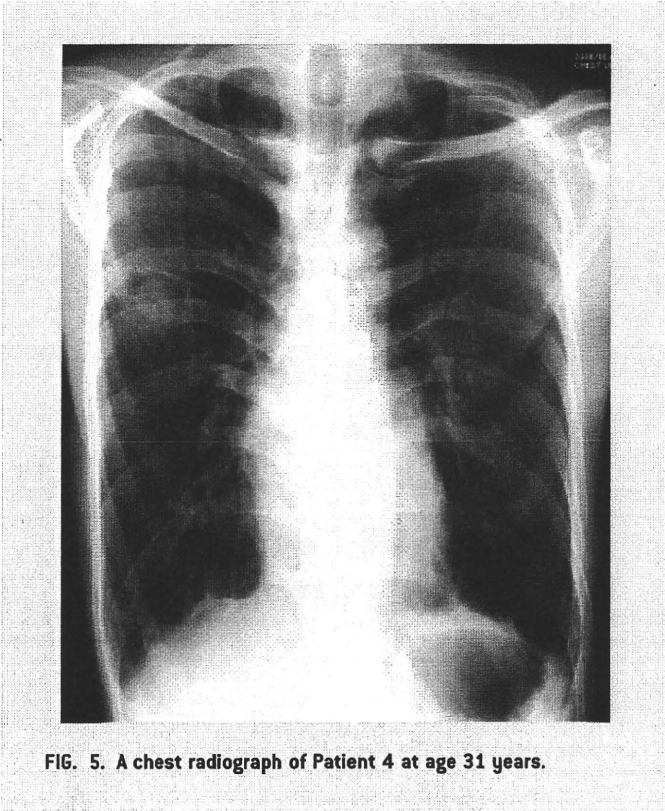


FIG. 5. A chest radiograph of Patient 4 at age 31 years.

heels (Fig. 4I). His skin was redundant (Fig. 4J), bruisable, and fragile with atrophic scars (Fig. 4K). Prominent palmar wrinkles showed acrogeria (Fig. 4G). He showed hyperalgesia to pressure such as measuring blood pressure at the upper arms. He suffered from constipation requiring oral laxatives and *Lactobacillus*, but sometimes had diarrhea. Large bowel sounds resembling a frog croak were frequently heard. Cardiac ultrasonography showed grade 2 aortic valve regurgitation without dilation of the ascending aorta, and a trace of mitral valve regurgitation. Ophthalmological examinations showed microphthalmia, myopia, and astigmatism. He showed hypogonadism without voice changes.

When last seen by us, his physical condition had deteriorated. He was bedridden, because of recurrent subcutaneous large hematomas around the buttocks and thighs possibly caused by spontaneous rupture of vessels or muscles. He also suffered from a dislocation of the right hip joints after falling, followed by another large hematoma throughout the right leg.

Patient 4

The patient is a now 32-year-old Japanese man. Part of his clinical course was described previously [Yasui et al., 2003]. He underwent surgery for bilateral talipes equinovarus at age 1 year. He was diagnosed as EDS at age 4 years. He had six operations for bilateral retinal detachment from age 16 years, and had surgery for carpal tunnel syndrome at age 18 years. Talipes valgus and planus was noticed at age 6 years, and progressed. He wore special orthopedic shoes from age 9 years and underwent surgical fixations of bilateral ankle joints at age 22 years.

At age 23 years, he was admitted for surgical fixation of the left toes. His weight was 38.0 kg (-2.4 SD) and height 168.5 cm (-0.4 SD). He had mild excavation of the left thorax and mildly hyperextensible skin with bruisability. During the preoperative evaluation, prolonged bleeding time was noticed (11 min [normal, 1–5]). He developed a large subcutaneous hematoma over the buttocks after a mild injury, with a decline of hemoglobin concentration from 9.2 to 4.9 g/dl. The hematoma was not resorbed with intranasal DDAVP and intravenous carbazochrome sodium sulfate, tranexamic acid, and packed red cells. It took 9 months for resorption of the hematoma after a combined therapy of daily intranasal DDAVP and weekly intramuscular conjugated estrogen. At age 31 years, he suffered from a right hemopneumothorax with a large bulla, which was treated with chest tube drainage and transfusion of packed red cells.

When last seen by us at age 31 years, his hemostatic condition was controlled by weekly oral doses of conjugated estrogen and temporary intranasal DDAVP after injuries. He had a slender and asymmetric face with hypertelorism, blue sclerae, a high palate, and a protruding jaw. His skeletal features included slender fingers with flexion contractures and limited movements, chronic dislocations of bilateral radial heads, flexed and immobile toes, kyphoscoliosis, and talipes valgus and planus with difficulty in walking. Acrogeria-like palmar wrinkles were noted.

Patient 5

The patient, now a 20-year-old Japanese woman, was the first child of a healthy 24 year-old mother and a healthy 25-year-old father. She was delivered by cesarean section for a breech presentation at 39 weeks of gestation. Her birth weight was 2,800 g (-0.4 SD), length 47.0 cm (-1.1 SD), and OFC 35.0 cm ($+1.0$ SD). She exhibited apnea with severe cyanosis on the day of her birth, and was admitted to a neonatal ICU. Her craniofacial features included a large anterior fontanelle, hypertelorism, short and downslanting palpebral fissures, strabismus, a short nose with a hypoplastic columella, a long philtrum, a thin upper lip vermilion, a small mouth, a high palate, and micro-retrognathia. Her skeletal features included pectus excavatum, slender fingers with flexion-adduction contractures of bilateral thumbs and extension contractures in bilateral index to little fingers (Fig. 6A), and bilateral talipes equinovarus (Fig. 6C,D). She hated to be hugged tightly. Bilateral talipes equinovarus were treated with serial plaster casts, followed by surgical corrections at age 1 year. At the operation, anomalous insertions of the flexor muscles were noted: the flexor digitorum longus and the flexor hallucis longus were not inserted to the toes, but were fused in the soles, forming a fascia-like tissue. Bilateral hip dislocations were treated by a brace, followed by a surgical correction in the right hip. Her skin was easily bruised and torn after a mild injury. Recurrent joint dislocations occurred easily or spontaneously at the elbows, shoulders, and knees. Constipation was treated with glycerin enema, and then improved with drinking milk or coffee after age 2 years. She raised her head at age 8 months, sat unassisted at age 1 year, and walked independently at age 2 years and 2 months. She was suspected to have EDS at around age 5 years.

She developed large subcutaneous hematomas over the buttocks after falling at age 6 years and age 7 years. At age 9 years, she suffered