

XYLT1 Mutations in Desbuquois Dysplasia Type 2

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Desbuquois dysplasia (DBQD) is a severe condition characterized by short stature, joint laxity, and advanced carpal ossification. Based on the presence of additional hand anomalies, we have previously distinguished DBQD type 1 and identified *CANT1* (calcium activated nucleotidase 1) mutations as responsible for DBQD type 1. We report here the identification of five distinct homozygous xylosyltransferase 1 (*XYLT1*) mutations in seven DBQD type 2 subjects from six consanguineous families. Among the five mutations, four were expected to result in loss of function and a drastic reduction of *XYLT1* cDNA level was demonstrated in two cultured individual fibroblasts. Because xylosyltransferase 1 (XT-I) catalyzes the very first step in proteoglycan (PG) biosynthesis, we further demonstrated in the two individual fibroblasts a significant reduction of cellular PG content. Our findings of *XYLT1* mutations in DBQD type 2 further support a common physiological basis involving PG synthesis in the multiple dislocation group of disorders. This observation sheds light on the key role of the XT-I during the ossification process.

Introduction

Desbuquois dysplasia (DBQD [MIM 251450]) belongs to the multiple dislocation group of disorders (group 20, international classification of skeletal disorders).¹ It is characterized by dislocations of large joints, severe pre- and postnatal growth retardation (−5 SD), joint laxity, and flat face with prominent eyes. Radiological features include short long bones with a monkey wrench appearance of the proximal femora (exaggerated trochanter) and advanced carpal and tarsal ossification. Based on the presence or absence of additional hand anomalies (ranging from extra ossification center distal to the second metacarpal, delta phalanx, or bifid distal phalanx of the thumb), we have distinguished DBQD type 1 from DBQD type 2.^{2,3}

In 2009, we identified *CANT1* (calcium activated nucleotidase 1 [MIM 613165]) mutations as responsible for DBQD type 1.⁴ *CANT1* mutations have then been reported in the “Kim variant” of DBQD, characterized by short metacarpals and elongated middle and proximal phalanges but very short distal phalanges.^{5,6} All subjects with the Kim variant were of Japanese or Korean origin and had a p.Val226Met substitution on at least one allele, with a shared common haplotype, supporting a founder effect in this population.

After our initial study, we screened *CANT1* in a series of 38 DBQD-affected individuals and found *CANT1* mutations not only in all subjects with DBQD type 1 but also in a Kim variant subject and in one atypical DBQD-affected individual with thumb anomaly and major joint dislocations.⁷ However, none of our 30 DBQD type 2 individuals was found to carry a *CANT1* mutation.⁷

DBQD types 1 and 2 have overlapping features with the autosomal-dominant form of Larsen syndrome (MIM

150250) resulting from *FLNB* mutations (MIM 603381),⁸ the spondylo-epiphyseal dysplasia form with dislocations (MIM 143095) resulting from chondroitin 6-O-sulfotransferase 3 (C6ST-1) (*CHST3* [MIM 603799]) mutations,⁹ the diastrophic dysplasia (DTD [MIM 222600]) resulting from solute carrier family 26 (sulfate transporter) member 2 (*SLC26A2* [MIM 606718]) mutations,¹⁰ and a chondrodysplasia with joint dislocations resulting from *IMPAD1* (inositol monophosphatase domain-containing protein 1 [MIM 614010]) mutations.¹¹ Apart from Larsen syndrome, all these disorders are characterized by a reduced amount of proteoglycans (PGs), which are known to be essential macromolecules with a large panel of functions, including extracellular matrix organizers and cell signaling mediators.¹² PGs are composed of a core protein with at least one glycosaminoglycan (GAG) chain attached via a Ser-Gly consensus motif and can be either located at the cell surface or secreted in the extracellular matrix.¹²

Because of the clinical overlap of this group of disorders, we screened *CHST3*, *SLC26A2*, and *IMPAD1* in the remaining DBQD type 2 individuals and found a homozygous *CHST3* mutation in 1/30¹³ and *IMPAD1* mutations in 2/30 subjects.

Among the remaining 27 DBQD type 2 subjects with no known molecular basis, only 20 with detailed clinical and radiological data and regular follow-up were included in our study. To identify the DBQD type 2 gene, we selected two siblings from consanguineous parents for whole-exome sequencing analysis and eventually identified homozygous mutations in seven individuals from six families in the xylosyltransferase 1 gene (*XYLT1* [MIM 608124]). *XYLT1* encodes xylosyltransferase 1 (XT-I, EC 2.4.2.26), which is involved in PG synthesis.¹² Indeed, the very first step of this process consists of the initiation of the

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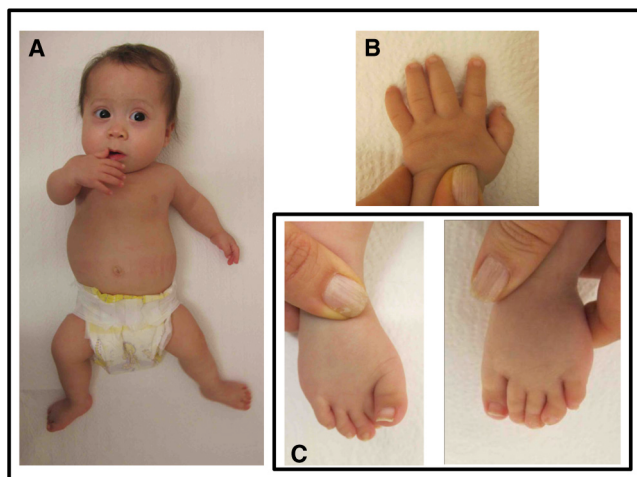


Figure 1. Clinical Features of One DBQD Type 2 Subject with *XYLT1* Mutation

Individual 7 (family 6) at 12 months. Note the flat face, narrow thorax, short limbs (A) with hip dislocation, hyperlaxity of fingers (B), and deviation of toes (C).

glycosaminoglycan (GAG) chains and can be catalyzed by the two paralogs XT-I and XT-II.¹⁴

Material and Methods

Individuals

The 20 DBQD-affected individuals selected for this study fulfilled the diagnostic criteria for DBQD type 2, namely severe pre- and postnatal short stature, short extremities, dislocations with monkey wrench appearance of the femora, short long bones with metaphyseal widening, epiphyseal dysplasia, and advanced carpal and tarsal ossification (Figure 1). Among them, the two siblings selected for whole-exome sequencing analysis were born to healthy consanguineous Tunisian parents ($f = 1/32$). At birth,

they both had a severe short stature, hypotonia, flat face with prominent eyes, hyperlaxity, hip dislocation, and narrow thorax leading to a respiratory distress that spontaneously resolved in the first year of life. Skeleton X-rays showed a monkey wrench appearance of the femoral neck, epiphyseal dysplasia, knee dislocation, and advanced carpal bone age (Figure 2, Table 1).

Samples

Informed consent for participation and sample collection were obtained via protocols approved by the Necker Hospital ethics board committee. Venous blood was obtained for DNA extraction from DBQD-affected individuals (QIAamp DNA blood Maxi kit, QIAGEN). Fibroblast cultures were established from skin biopsies obtained from scalp incision.

Exome Sequencing

Exome capture was performed at the genomic platform of Fondation IMAGINE (Paris, France) with the SureSelect Human All Exon kit (Agilent Technologies).¹⁵ Single-end sequencing was performed on an Illumina Genome Analyzer IIx (Illumina) generating 72-base reads. Sequence data were analyzed by the Bioinformatic platform and visualized via the interface created by the Bioinformatic platform (Université Paris Descartes, Paris). For sequence alignment, variant calling, and annotation, the sequences were aligned to the human genome reference sequence (UCSC Genome Browser, hg18 build) by BWA aligner.¹⁶ Downstream processing was carried out with the Genome Analysis Tool-kit (GATK),¹⁷ SAMtools¹⁸ and Picard Tools. Substitution calls were made with GATK Unified Genotyper, whereas indel calls were made with a GATK IndelGenotyperV2. All calls with a read coverage $\leq 2\times$ and a Phred-scaled SNP quality of ≤ 20 were filtered out. All the variants were annotated with an in-house-developed annotation software system.

Microsatellite Analysis of *XYLT1* Locus

Microsatellite analysis was performed in consanguineous families at the *XYLT1* locus on chromosome 16p12.3, by means of four

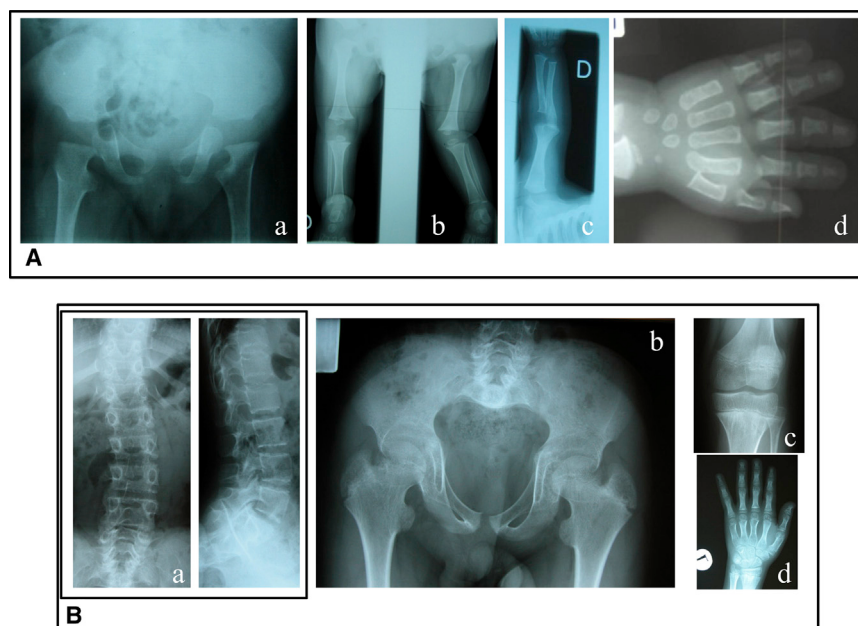


Figure 2. Radiological Features of Two DBQD Type 2 Subjects with *XYLT1* Mutations

(A) Subject 1 (family 1) at 8 months. Note the monkey wrench appearance of the proximal femora, absence of upper femoral epiphyseal centers (a), knee dislocation (b), short long bones (c), and advanced carpal bone age (d).

(B) Subject 4 (family 3) at 12 years of age. Note the scoliosis with irregular vertebral endplates (a), monkey wrench appearance of the proximal femora with short femoral necks, short and broad iliac wings (b), flat knee epiphyses (c), short metacarpals, and advanced carpal bone age and prominent wrist epiphyses (d).

Table 1. Clinical Features of the Seven DBQD Type 2-Affected Individuals from Six Families

	Ethnic Origin	Csg	Parameters at Birth	Clinical Features at First Exam^a	Radiological Features	Follow-up: Growth and Skeleton	Other
Family 1, sibling 1 (female)	Tunisian	1/32	length 37 cm (term)	hyperlaxity respiratory distress hypotonia flat face	monkey wrench of the femoral neck epiphyseal dysplasia knee dislocation advanced carpal bone age	at 24 years of age: height 111.5 cm (< –6 SD)	mild intellectual disability
Family 1, sibling 2 (male)	Tunisian	1/32	length 41 cm (term)	hypotonia narrow thorax hip dislocation flat face	monkey wrench of the femoral neck epiphyseal dysplasia knee dislocation advanced carpal bone age	at 20 years of age: height 121 cm (< –6 SD)	intellectual disability
Family 2 (female)	Mauritian	1st cousins	weight 2,000 g	lower limb deformity multiple dislocations (hip, knee)	monkey wrench of the femoral neck brachymetacarpus epiphyseal dysplasia	at 13 years of age: flexion of hips and knees valgus deformation of the lower limbs patella instability multiple surgeries weight 35 kg (–1 SD) height 98 cm (< –6 SD) toe deformations	mild intellectual disability
Family 3 (male)	Belgian	1st cousins	weight 2,570 g, length 39 cm, HC 33 cm (term); transient respiratory problems in the neonatal period	flat face low nasal bridge blue sclerae cleft palate short neck narrow thorax short limbs	coronal clefts in the neonatal period, thereafter mild platyspondyly shortening of tubular bones absent ossification of distal femoral epiphyses at birth	at 12 years, 9 months of age: weight 23.7 kg (–3.5 SD) height 109.5 cm (–6 SD) span 111 cm HC 50.8 cm (–2 SD) flat face, prominent eyes, low nasal bridge pectus carinatum, narrow thorax hyperlaxity of fingers and knees (genua valga) broad feet, toe clinodactyly	intellectual disability
Family 4	Turkish	1st cousins	length 44 cm (term)	at 3.5 months: height 48.5 cm head control: 2 months hip dislocation (right) knee dislocation simian creases hypermobile fingers flat face blue sclerae	neonatal period: advanced carpal ossification right hip and bilateral knee dislocation at 8 months: advanced bone age elbow dislocation	at 11 months of age: height 56 cm at 5.5 years of age: height 84 cm (–5.5 SD) coarse and round face full cheek, long philtrum, mild micrognathia hypermobile fingers moderate truncal obesity pectus excavatum	sitting at 9 months walking at 3 years
Family 5	Turkish	1st cousins	length 43 cm (term)	at 52 days: height 46 cm round and flat face epicanthal folds short extremities and hands bilateral simian crease	neonatal period: monkey wrench advanced carpal ossification, short metacarpals and phalanges widened anterior ribs at 9 years: patella and elbow subluxation short iliac wings	at 13 months of age: height 59 cm at 9 years of age: height 99 cm at 13 years of age: height 109 cm (–9 SD) HC 53 cm coarse and round face, blue sclera, proptotic eyes short extremities increased lumbar lordosis hypermobile joints pectus excavatum pes planus truncal obesity	mild intellectual disability sitting at 8 months walking at 24 months

(Continued on next page)

Table 1. Continued							
	Ethnic Origin	Csg	Parameters at Birth	Clinical Features at First Exam ^a	Radiological Features	Follow-up: Growth and Skeleton	Other
Family 6	Turkish	from the same village	length 33 cm, weight 1,200 g (born at 36 WG)	cleft palate subluxation of right knee	monkey wrench advanced carpal ossification and tarsal extra ossification double proximal femoral epiphyses short phalanges with short 1 st metacarpal	at 12 months of age: height 50 cm (<−6 SD) hypermobile joints respiratory problems 2 first months of life	normal motor development at age 2

Abbreviations are as follows: Csg, consanguinity; WG, weeks of gestation; HC, head circumference.
^aFirst exam was performed at birth unless noted otherwise.

repeat-containing microsatellite markers: D16S405, D16S499 located on both sides of *XYLT1*, and two intragenic microsatellites, D16S103 and D16S3017. FAM-labeled PCR products were run on an ABI 3130 sequencer and analyzed with GeneMapper (Applied Biosystems).

Sequencing Analysis of *XYLT1*

The exons and exon-intron boundaries of *XYLT1* were amplified with specific primers (available upon request). Amplification products were purified by ExoSapIT (Amersham) and directly sequenced with the Big Dye Terminator Cycle Sequencing Ready Reaction kit v.1.1 on an automatic sequencer (ABI3130xl; PE Applied Biosystems). Sequence analyses were performed with Seqscape software v.2.5 (Applied Biosystems).

RNA Extraction, Reverse Transcription, and Real-Time Quantitative PCR

Skin primary fibroblasts were cultured in DMEM medium supplemented with 10% fetal calf serum and antibiotics at 37°C in a humidified atmosphere containing 5% CO₂. For real-time PCR analyses, fibroblasts were seeded onto 96-well plates after standardization via the cell counter CASY Model TT (Roche). After reverse transcription of mRNA with M-MLV reverse transcriptase kit (Invitrogen), PCR analyses were performed on the 7300 Real Time PCR System (Roche). The cDNA level of *XYLT1*, *XYLT2*, and *B4GALT7* was normalized to *GAPDH*.

Data were generated from three independent experiments performed in quadruplicate and compared to control data with one-way ANOVA followed by Tukey's post hoc correction test with GraphPad Prism 6.0 software (GraphPad).

Analysis of PG Biosynthesis and Profile after Metabolic Radiolabeling in Cultured Fibroblasts

Fibroblasts were seeded onto 6-well plates and incubated the following day in Fischer's medium containing 10 µCi/ml Na₂[³⁵S]SO₄ in the presence of 5 µM of methylumbelliferyl-β-D-xylopyranoside (4-MUX) or vehicle (DMSO) as described previously.¹⁹ After radiolabeling, each cell media or cellular fractions were applied to G-50 columns (GE Healthcare, VWR) and quantified by scintillation counting on Packard 1600TR Tri Carb Liquid Scintillation Analyzer.

To establish the PG profile, the PG fractions were digested for 4 hr at 37°C by either the Chondroitinase ABC (cABC) from *Proteus vulgaris* (Sigma-Aldrich) or by a mixture containing both the Heparinases II and III from *Flavobacterium heparinum* (Sigma-Aldrich). All samples were resolved by SDS-PAGE with Criterion Precast gels (4%–15% Bis-Tris, Bio-Rad) and visualized by autoradiography. In order to compare PG distribution, the same amount of radioactivity was loaded in each well.

Results

Identification of a Homozygous *XYLT1* Mutation by Whole-Exome Sequencing in Two Siblings with DBQD Type 2

For exome-sequencing analysis, we first focused our analyses on nonsynonymous variants, splice acceptor and donor site mutations, and coding indels, anticipating that synonymous variants were far less likely to cause disease (Table 2). We also defined variants as previously

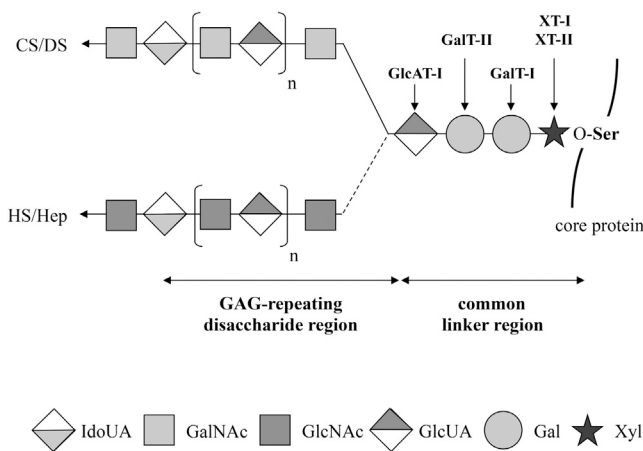
Single-Nucleotide Variations					
Individuals	Total	Without Duplicates ^a	Not in Databases ^b	In Essential Splicing and Coding Regions ^c	Deleterious/Damaging or Unknown ^d
Family 1, sibling 1 (Csg)	11,534	5,560	216	60	11
Family 1, sibling 2 (Csg)	11,324	5,245	159	35	4
Family 1, siblings 1+2	8,881	4,590	70	18	2

Abbreviation is as follows: Csg, consanguinity.
^aWithout duplicates resulting from the homozygous status of subjects.
^bDatabases searched were dbSNP, 1000 Genomes, Exome Variant Server, and in-house exomes.
^cVariants included were essential splicing and nonsynonymous (missense, nonsense, deletion and insertion); eliminated variants were intergenic or intronic non-coding RNA or UTR splicing.
^dAs determined by SIFT or PolyPhen.

Based on the recessive mode of inheritance of DBQD, two genes were found to harbor an identical homozygous substitution in the two siblings: the structural maintenance of chromosome 1B (*SMC1B* [MIM 608685], RefSeq accession number NM_148674.3) and the xylosyltransferase 1 gene (*XYLT1*, RefSeq NM_022166.3) (Table 2). These results were confirmed by direct sequencing.²⁰ The *SMC1B* (c.11T>G [p.Leu4Arg]) and *XYLT1* (c.1792C>T [p.Arg598Cys]) substitutions cosegregated with the disease, were present at the heterozygous state in the parents, and were considered pathogenic in the PolyPhen and Sift analytical programs. *SMC1B* belongs to the cohe-

Among the 19 remaining families affected by DBQD type 2, 5/13 inbred individuals were homozygous at the *XYLT1* locus and direct sequencing was finally performed in 11 families (including 6 nonconsanguineous families). A total of five distinct *XYLT1* mutations in seven individuals (six families) with DBQD type 2 were identified (Figures 1 and 2, Tables 1 and 3). Two resulted in a premature stop codon (c.276dupG [p.Pro93Alafs*69], c.439C>T [p.Arg147*]) located N-terminal to the glycosyltransferase family 14 and xylosyltransferase domains; two were splice site mutations (c.1290–2A>C, c.1588–3C>T) located in the donor sites of exons 6 and 8, respectively, involved in the glycosyltransferase family 14 domain; and one was a missense substitution (c.1792C>T [p.Arg598Cys]) located in the xylosyltransferase domain, considered as damaging in PolyPhen and Sift analytical programs (Figure 4, Table 3). All mutations segregated with the disease and were not identified in 200 control chromosomes. *SMC1B* was excluded by linkage analysis in the remaining five consanguineous families with *XYLT1* mutations.

By quantifying the cDNA level of *XYLT1* in the two cultured individual fibroblasts (p.Arg147*, female aged 14; and p.Pro93Alafas*69, male aged 23), we found a dramatic reduction of *XYLT1* cDNA level (Figure 5A). Because the two xylosyltransferases XT-I and XT-II¹⁴ can both initiate the synthesis of GAG chains to PG core proteins, we also measured the cDNA level of *XYLT2* and found a significant decrease of *XYLT2* expression in the two



XT-I and XT-II participate in the initiation of the PG biosynthesis by transferring a Xyl to specific Ser residues of the core protein. Two Gal and one GlcUA will be sequentially added via the action of GalT-I, GalT-II, and GlcAT-I to form the common linker region. The addition of a GalNAc will then initiate the assembly of chondroitin-sulfate/dermatan-sulfate (CS/DS) chains while addition of a GlcNAc will initiate the synthesis toward heparan-sulfate/heparin (HS/Hep) chains. Mutations in genes involved in the assembly of the common linker region have been associated with several disorders, namely DBQD type 2 (*XYLT1*), Ehlers-Danlos progeroid syndrome, type 1 (*B4GALT7*), Ehlers-Danlos progeroid syndrome, type 2/spondyloepimetaphyseal dysplasia with joint laxity type 1 (SEMD-JL1) (*B3GALT6*), and Larsen-like syndrome (*B3GAT3*).

Table 3. *XYLT1* Mutations in the Six DBQD Type 2-Affected Families

Family	Ethnic Origin	Csg	No. of Affected Children	Nucleotide Change	Status	Amino Acid Change	Location
1	Tunisian	yes	2	c.1792C>T	Ho	p.Arg598Cys	Ex9
2	Mauritian	yes	1	c.439C>T	Ho	p.Arg147*	Ex3
3	Belgian	yes	1	c.276dupG	Ho	p.Pro93Alafs*69	Ex1
4	Turkish	yes	1	c.1588-3C>T	Ho	?	In7
5	Turkish	yes	1	c.1290-2A>C	Ho	?	In5
6	Turkish	yes	1	c.1290-2A>C	Ho	?	In5

Abbreviations are as follows: Csg, consanguinity; ho, homozygote; ?, mutation not localized in a protein domain.

individual cell lines when compared to control 2 (male, aged 12). No significant differences were found when individual data were compared to control 1 (female, aged 2.5; Figure S1 available online).

By contrast, the cDNA level of *B4GALT7*, which encodes the glycosyltransferase (GT) that adds the second sugar galactose (Gal) to the xylose (Xyl), was not affected by *XYLT1* mutations (Figures 3 and S1).

The PG Biosynthesis Is Strongly Affected by *XYLT1* Mutations in the Two Cultured Individual Fibroblasts

To evaluate the impact of the loss of function of *XYLT1*, the cells were treated with the 4-MUX, an exogenous acceptor substrate of the GalT-I that can increase GAG synthesis (Figure 3). No major differences in the sulfate incorporation level were found for the extracellular fractions in the absence of 4-MUX. In the presence of 4-MUX, a significant increase of sulfate incorporation was observed, indicating that GAG biosynthesis was initiated and enhanced by the 4-MUX via the GalT-I activity (Figure S2). In contrast, the incorporated sulfate level associated with the cellular fractions was significantly lower in the two cultured individual fibroblasts and no increase was observed after the addition of 4-MUX (Figure 6).

Modifications in PG Pattern Are a Consequence of *XYLT1* Mutations

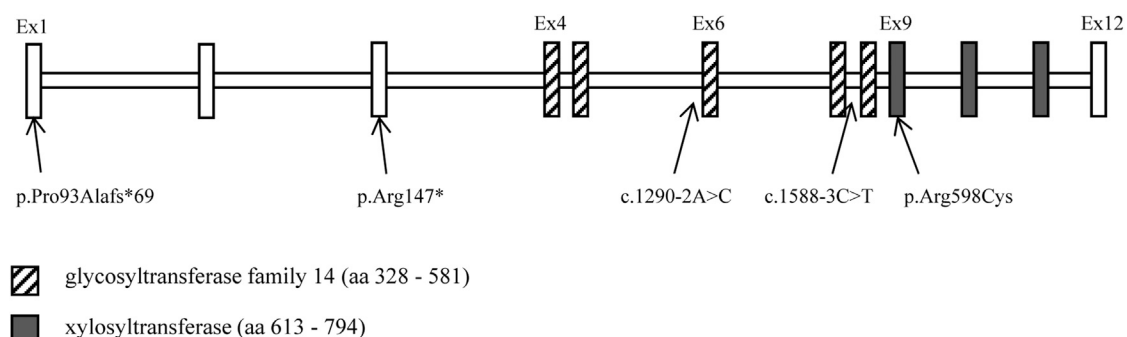
An important reduction of secreted and large chondroitin-sulfate proteoglycan (CSPG) production was observed in

the two cultured individual fibroblasts (lanes 4, 6, 7, and 9, Figure 7). In the p.Arg147* cell line, the presence of a stronger second band at 90 kDa (lanes 7 and 9, Figure 7) was observed, presumably corresponding to the accumulation in the extracellular media of other small CSPGs such as decorin. For individual and control fibroblast cell lines, the addition of 4-MUX led to the production of GAG chains with an increased ratio of lower-molecular-weight GAGs. The sensitivity to Chondroitinase ABC treatment supported that these GAG chains were mainly chondroitin-sulfate (CS) (lanes 11, 14, and 17, Figure 7). In the p.Arg147* cultured fibroblasts, the accumulation of CSPGs at 90 kDa was still observed after the xyloside addition (lanes 16 and 18, Figure 7).

For the cellular fractions, mainly heparan-sulfate proteoglycans (HSPGs) with high molecular weight (>200 kDa) were detected and no major differences in PG profiles and distribution were observed between control 2 and individual fibroblasts in the absence or presence of 4-MUX (data not shown).

Discussion

We report here the identification of *XYLT1* mutations in seven individuals from six families with DBQD type 2. All individuals presented severe pre- and postnatal short stature below -6 SD, flat face with prominent eyes, short extremities, dislocations with monkey wrench appearance of the femora, short long bones with metaphyseal

**Figure 4. Location of the Five *XYLT1* Mutations**

Organization of *XYLT1* gene and protein with the two glycosyltransferase family 14 and xylosyltransferase domains

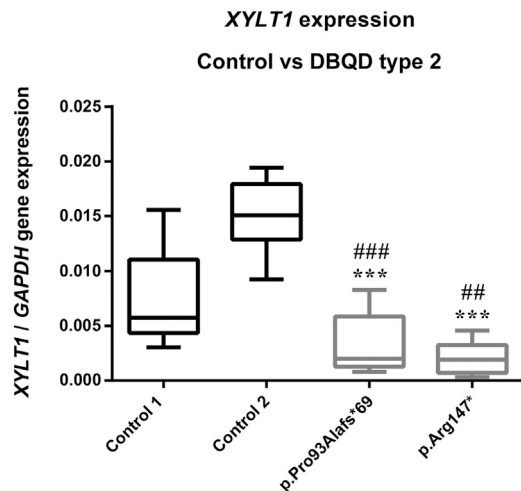


Figure 5. cDNA Level of *XYLT1* in Controls and Two Cultured Individual Fibroblasts

cDNA levels were normalized to *GAPDH*. Individuals p.Arg147* and p.Pro93Alafs*69 are female aged 14 and male aged 23 years, respectively. Control 1 (female) and 2 (male) are aged 2.5 and 12 years, respectively. Control (black boxes) and individual (gray boxes) data are presented as box plots and are the mean of three independent experiments performed in quadruplicate. ## $p < 0.01$, ### $p < 0.001$ compared to control 1, *** $p < 0.001$ compared to control 2 by one-way ANOVA with Tukey's post hoc test. Note that mutations in *XYLT1* affect cDNA level of *XYLT1*.

widening, epiphyseal dysplasia, and advanced carpal and tarsal ossification. No significant clinical or radiological differences could be found with the remaining 14 DBQD type 2 subjects with unknown molecular bases. However, long-term follow-up of *XYLT1*-mutated individuals emphasizes the severity of the short stature (< -6 SD) contrasting with obesity, lower limb and foot deformities requiring often repeated surgeries, and intellectual disability (5/7). Interestingly, respiratory distress was present at birth in 4/7 subjects and spontaneously resolved in the first years of life but thorax narrowness persisted in the eldest children. No major scoliosis was observed.

Among the five mutations identified, four were expected to result in loss of function, confirmed in two by the demonstration of a drastic reduction of gene expression in cultured fibroblasts. The substitution p.Arg598Cys identified is located in a highly conserved region of the protein, between the glycosyltransferase family 14 and the catalytic domains of the enzyme,²¹ pointing out possible changes in the protein folding and/or stability as the positively charged amino acid Arg is replaced by a polar and neutral Cys residue. Very recently, another substitution of XT-I (p.Arg481Trp) has been reported in Turkish siblings presenting short stature and minor skeleton features associated with intellectual disability.²² X-rays of the two siblings (aged 16 and 18 years) do not show monkey wrench appearance of the femoral neck or advanced carpal bone age, but these features disappear in the course of DBQD. However, among features listed in the two siblings, coxa valga, broad thumbs, abnormal feet, and obesity are

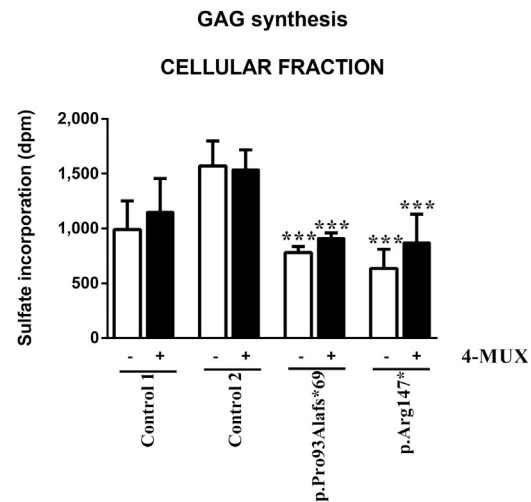


Figure 6. Sulfate Incorporation of Cellular Fractions in Controls and the Two Cultured Individual Fibroblasts in the Presence of 4-MUX or Vehicle

The black (+4-MUX, 5 μ M) and white (DMSO vehicle) bars represent the mean $\pm$ SEM of two independent experiments performed in triplicate. *** $p < 0.001$ when compared to the control 2 by one-way ANOVA with Tukey's post hoc test.

Note that *XYLT1* mutations have an impact on the biosynthesis of cellular PGs in the two cultured individual fibroblasts.

reminiscent of DBQD spectrum. The milder phenotype observed in the two siblings might be due to a partial loss of function of XT-I.²²

PG formation begins with the synthesis of a common linker chain of four sugar residues. The very first step of this process can be catalyzed by the two paralogs XT-I and XT-II¹⁴ leading to the initiation of the two types of GAG chains (Figure 3).²³ Our finding of decreased *XYLT2* cDNA level in two individual fibroblasts suggests that *XYLT1* loss of function may affect *XYLT2* gene expression. However, because of the variability observed in cultured fibroblasts (even in control samples), this observation will need to be confirmed in additional individual samples compared to age- and gender-matched control samples.

The two paralogs XT-I and XT-II exhibit a high degree of homology within their catalytic domain²⁴ and are ubiquitously expressed in human tissues (data not shown).^{25,26} Despite the apparent redundancy in catalytic activity as demonstrated by in vitro enzymatic assays^{25,26} and in gene expression, the identification of *XYLT1* mutations in a severe chondrodysplasia supports a pivotal role of *XYLT1* during the ossification process. The specific implication of XT-I in skeletal development and during cartilage repair has been previously reported.^{27,28} Moreover, a recent study describing the phenotype of the *pug* mouse mutant carrying a homozygous missense mutation in *Xylt1* and characterized by a disproportionate dwarfism, premature chondrocyte maturation, and increased Indian Hedgehog signaling further supports a key role of XT-I in early chondrocyte maturation and skeletal length.²⁹

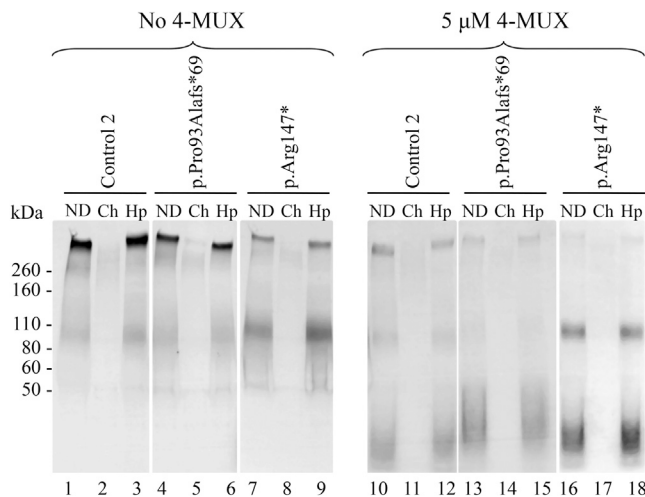


Figure 7. Gel Electrophoresis of Radiolabeled GAG Chains Extracted from Control 2 and the Two Cultured Individual Fibroblasts Media in the Presence 4-MUX or Vehicle

For each sample, three lanes corresponding to nondigested (ND), digested by the Chondroitinase ABC (Ch), and digested by the Heparinases II and III (Hp) were loaded onto the gel and the volumes were adjusted to dpm values to visualize GAG profile and distribution. Control 2 (male) is aged 12.

Note that mutations in *XYLT1* change the distribution and profile of secreted PGs in the two cultured individual fibroblasts.

The presence of intellectual disability was observed in at least five subjects and highlights also a possible role of *XYLT1* in brain development. Interestingly, common CSPGs such as aggrecan, perlecan, and biglycan are both found in cartilage and expressed during brain development.^{30,31} Our findings that mainly CSPGs were secreted into the cell media and that synthesis of large CSPGs was clearly reduced for the p.Arg147* cell line suggest that both PG-specific expression pattern and impaired synthesis of large CSPGs may contribute to the tissue specificity of the disease. Interestingly, a strong decrease of sulfate incorporation in the high-density PG fraction (mainly CSPGs) was also observed in the *pug* mutant.²⁹ Finally, the accumulation of PGs in the range of 90 kDa observed for the p.Arg147* cell line could compensate the important loss of high-molecular-weight CSPGs, whereas for the p.Pro93Alafs*69 cell line, the loss of large CSPGs was less pronounced.

The consequences of *XYLT1* mutations on PG synthesis were also clearly illustrated by a significant reduction of cellular PG synthesis in the two cultured individual fibroblasts. This reduction could not be rescued by the addition of 4-MUX, indicating that most of the 4-MUX-primed GAGs were presumably secreted into the cell media. Because XT-I catalyzes the first step of PG synthesis, an altered expression of this enzyme is likely to impact PG synthesis either by limiting the initiation of the GAG chains or (less probably) by leading to an abnormal decrease of the PG core protein level.

Interestingly, trials based on the use of either heparin-like molecules in animals³² to repair skull defects or HS

supplementation³³ in cultures of human mesenchymal stem cells have already been tested. Although protein xylosylation is known to be the rate-limiting step in PG biosynthesis, this can be bypassed by the addition of exogenous xylosides (such as 4-MUX) into cell media owing to the GalT-I activity. Our findings of identical *B4GALT7* cDNA level in controls and individual fibroblasts but increased GAG production in response to 4-MUX treatment in individual fibroblasts may support the development of new approaches based on the use of GAGs.

To date, mutations in genes involved in the synthesis of the common linker region have been identified in β 1,4-galactosyltransferase 7 (GalT-I = *B4GALT7* [MIM 604327]),^{34–36} β 1,3-galactosyltransferase 6 (GalT-II = *B3GALT6* [MIM 615291]),^{37,38} and β 1,3-glucuronosyltransferase I (GlcAT-I = *B3GAT3* [MIM 606374]),³⁹ and the human phenotypes associated included Ehlers-Danlos progeroid type (type 1 MIM 130070, type 2 MIM 615349), spondylo-epiphyseal-meta-dysplasia with joint laxity type 1 (SEMD-JL1 [MIM 271640]), and a Larsen-like phenotype with heart defect (MIM 245600). Perturbations in GAG modifications (mainly sulfation) have been also reported in (1) spondyloepiphyseal dysplasia (SED) with dislocations resulting from *CHST3* mutations (C6ST-1) responsible for altered sulfation of CSPGs,⁴⁰ (2) a chondrodysplasia with joint dislocations resulting from mutations in *IMPAD1* that encodes for the golgi-resident nucleotide phosphatase gPAPP and resulting in an under-sulfation of PGs,¹¹ and (3) DBQD resulting from *CANT1* mutations accounting for shorter GAG chains.⁷

Our findings of *XYLT1* mutations in DBQD type 2 further confirm a common physiological basis in the group of multiple dislocation disorders involving PG synthesis and also shed light on the pivotal and specific role of XT-I during the ossification process.

Supplemental Data

Supplemental Data include two figures and can be found with this article online at <http://www.cell.com/AJHG/>.

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Web Resources

The URLs for data presented herein are as follows:

Alamut Interpretation Software 2.0 (gateway for PolyPhen-2, SIFT, SpliceSiteFinder-like, MaxEntScan, NNSPLICE, and Human Splicing Finder), <http://www.interactive-biosoftware.com>
Ensembl Genome Browser, <http://www.ensembl.org/index.html>
GeneCards, <http://www.genecards.org>

NCBI, <http://www.ncbi.nlm.nih.gov/>
 Online Mendelian Inheritance in Man (OMIM), <http://www.omim.org/>
 Primer3, <http://bioinfo.ut.ee/primer3-0.4.0/primer3/>
 RefSeq, <http://www.ncbi.nlm.nih.gov/RefSeq>
 UCSC Genome Browser, <http://genome.ucsc.edu>

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