Mutations in the Fukutin-Related Protein Gene (FKRP) Cause a Form of Congenital Muscular Dystrophy with Secondary Laminin α2 Deficiency and Abnormal Glycosylation of α-Dystroglycan

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The congenital muscular dystrophies (CMD) are a heterogeneous group of autosomal recessive disorders presenting in infancy with muscle weakness, contractures, and dystrophic changes on skeletal-muscle biopsy. Structural brain defects, with or without mental retardation, are additional features of several CMD syndromes. Approximately 40% of patients with CMD have a primary deficiency (MDC1A) of the laminin α2 chain of merosin (laminin-2) due to mutations in the LAMA2 gene. In addition, a secondary deficiency of laminin α2 is apparent in some CMD syndromes, including MDC1B, which is mapped to chromosome 1q42, and both muscle-eye-brain disease (MEB) and Fukuyama CMD (FCMD), two forms with severe brain involvement. The FCMD gene encodes a protein of unknown function, fukutin, though sequence analysis predicts it to be a phosphoryl-ligand transferase. Here we identify the gene for a new member of the fukutin protein family (fukutin related protein [FKRP]), mapping to human chromosome 19q13.3. We report the genomic organization of the FKRP gene and its pattern of tissue expression. Mutations in the FKRP gene have been identified in seven families with CMD characterized by disease onset in the first weeks of life and a severe phenotype with inability to walk, muscle hypertrophy, marked elevation of serum creatine kinase, and normal brain structure and function. Affected individuals had a secondary deficiency of laminin α2 expression. In addition, they had both a marked decrease in immunostaining of muscle α-dystroglycan and a reduction in its molecular weight on western blot analysis. We suggest these abnormalities of α-dystroglycan are caused by its defective glycosylation and are integral to the pathology seen in MDC1C.

Introduction

The congenital muscular dystrophies (CMD) are a heterogeneous group of autosomal recessively inherited diseases, presenting at birth or within the first 6 mo of life with hypotonia, muscle weakness, and the variable appearance of contractures and characterized by dystrophic changes on skeletal-muscle biopsy (Dubowitz 1995). The differing degrees of motor developmental delay, physical disability, muscle pathology, and elevation of serum creatine kinase (CK), along with the variable presence of mental retardation and structural brain defects, underscores the heterogeneous nature of this condition. Recent epidemiological data suggest that the incidence of CMD is 4.65 × 10⁻⁵ and that its prevalence is 8 × 10⁻⁶ (Mostacciuolo et al. 1996), indicating that it is among the most frequent autosomal recessively inherited neuromuscular diseases.

One of the main objectives of the International Consortium on CMD has been to delineate and subdivide the various forms of CMD (Dubowitz 1994, 1995, 1997, 1999; Muntoni et al., in press). Seven genetically distinct entities have been identified so far. The most common form of CMD (~40% of cases) is due to a primary deficiency in the laminin α2 chain of merosin (laminin-2), resulting from mutations in the LAMA2 gene (MDC1A [MIM 156225]; Tome et al. 1994; Helbing-Leclerc et al. 1995). Other CMD disease genes identified so far are ITGA7, leading to integrin α7 deficiency...
(MIM 600536); Hayashi et al. 1998), COL6A2, coding for the α2 chain of collagen VI, in the Ullrich variant (UCMD [MIM 254090]; Camacho Vanegas et al. 2001); and fukutin, responsible for Fukuyama congenital muscular dystrophy (FCMD [MIM 253800], Fukuyama et al. 1981; Kobayashi et al. 1998). Other forms of CMD that have been mapped but for which the disease genes remain elusive are muscle-eye-brain disease (MEB [MIM 236670]), mapped to chromosome 1q23-24 (Brockington et al., 2000), MDC1B (Brockington et al., 2000), MEB (Haltia et al. 1992), and disruption of this axis is associated with several forms of muscular dystrophy (Campbell 1993).

FCMD is one of a number of CMD forms that have been shown to have a secondary reduction in the expression of laminin α2 (Hayashi et al. 1993). Recently, a profound depletion of skeletal and cardiac muscle α-dystroglycan has also been reported in FCMD (Hayashi et al. 2001). FCMD is common in Japan, because of a founder haplotype containing a 3.0-kb retrotransposoral insertion into the 3′ UTR of the FCMD gene (Kobayashi et al. 1998). Fukutin, a protein of unknown function, is predicted to be a secreted protein that has sequence similarities to a family of proteins involved in modifying cell-surface molecules such as glycoproteins and glycolipids (Aravind et al. 1999). Fukutin also contains a conserved DxD motif that is found in many glycosyltransferases (Aravind et al. 1999; Breton and Imberty 1999).

Although FCMD is rare outside Japan, other CMD syndromes with secondary laminin α2 deficiency with or without brain involvement are not, including MDC1B (Brockington et al., 2000), MEB (Haltia et al. 1997), and other, as-yet-unmapped variants (Talim et al. 2000; Villanova et al. 2000). We have used sequence analysis to identify fukutin homologues in the human genome as candidate genes for these variants of CMD.

In the present study, we report the identification of a gene encoding a new member of the fukutin-related-protein family (FKRP [EMBL accession number AJ314847]). This gene is mutated in a severe form of CMD, which we have called “MDC1C.” Patients harboring mutations in this gene had a secondary deficiency of laminin α2 and a profound reduction in α-dystroglycan immunostaining. In addition, the molecular weight of α-dystroglycan was reduced in muscle. Together, these findings suggest that this protein is abnormally glycosylated in MDC1C and is central to the pathology seen in this disorder.

Families, Material, and Methods

Family 1

We recently reported the clinical features of the two children in this Scottish family and suggested that they are affected by a novel form of CMD (Mercuri et al. 2000). Both children presented soon after birth with hypotonia and feeding difficulties. They never acquired the ability to walk, because of severe weakness, which also affected their facial muscles. Weakness affected the arms more than the legs, with prominent wasting of the deltoids and pectoralis muscles, whereas both calf and quadriceps muscles were hypertrophied. Cognitive development, intelligence, and vision were normal, as was brain magnetic resonance imaging. Serum CK was markedly elevated (9,000–14,550 IU/L, normal values <190 IU/liter). The older sibling died suddenly at age 7 years, following an upper respiratory tract infection. The younger sibling is now 4 years old and recently had a gastrostomy because of feeding difficulties.

Family 2

This is a consanguineous family from Libya with one affected child. In addition, two children on the paternal side of the family had died in the first decade of life with a similar illness. She presented in the first few weeks of life with hypotonia and feeding difficulties, followed by motor delay. On examination at age 16 mo, she could not walk and was weaker in her arms than in her legs. She had calf hypertrophy and facial weakness. Her serum CK was 2,478 IU/liter and she had a myopathic EMG. Her intelligence was normal.

Family 3

The affected child in this family is a 6.5-year-old girl from southern England who is currently unable to stand or walk. She presented with hypotonia in the first few
weeks of life, followed by delayed motor milestones. She could stand with support between the ages of 4 and 6 years. Her intelligence was normal, her serum CK was markedly elevated, and she had a muscle biopsy at age 9 mo that was dystrophic. At age 6 years, she had marked hypertrophy of the leg muscles with wasting and weakness of shoulder-girdle muscles, particularly her deltoid, pectoralis, and sternocleidomastoid muscles. Serum CK was elevated (2,695 IU/liter). A brain MRI showed normal results; an echocardiogram showed mildly impaired left-ventricular function.

Family 4

The third child of this Sudanese consanguineous family is affected. A detailed clinical description of this patient and those from families 5, 6, and 7 has recently been submitted elsewhere (S. Quijano-Roy, L. Galan, A. Ferreiro, F. Cheliout-Heraut, F. Gray, M. Fardeau, A. Barois, P. Guicheney, N. B. Romero, and B. Estournet, unpublished data). He presented at birth with hypotonia, and his motor development was severely delayed; he never acquired the ability to walk. Calf and tongue hypertrophy were noticed at the ages of 4 and 10 years, respectively. At age 14 years, he went into respiratory failure and has been ventilated via a tracheostomy ever since. His cognitive development and brain MRI were normal; he developed a transient cardiac insufficiency after a respiratory infection at age 11 years, but current cardiac function is normal. Serum CK was elevated (2,277 IU/liter) at age 10 years. He underwent three muscle biopsies (at the ages of 2, 3, and 9 years) that showed a dystrophic picture with normal expression of dystrophin and sarcoglycans. Laminin α2 expression was reduced with an antibody against the 80-kDa fragment and was absent with an antibody against the 300-kDa fragment. No muscle was available to study the expression of α-dystroglycan.

Family 5

The only child from this nonconsanguineous family of Caribbean origin is affected. She was symptomatic at birth with weakness and hypotonia, and her motor development was severely delayed. She could walk only with assistance between the ages of 3 and 6 years. Hypertrophy of the calves had been noticed in infancy, and macroglossia appeared around puberty. She developed progressive contractures in the elbows, knees, and fin-

### Database Screening

We performed BLAST analysis for new fukutin homologues, using the BLAST server at NCBI. The identification of transcribed sequences (ESTs) and their genomic location was also performed using BLAST analysis. Sequence alignments were obtained using the ClustalW Service at the European Bioinformatics Institute.

### Northern Blot Analysis

We determined the expression of FKRP and an estimate of transcript size by northern blot analysis using a blot containing poly(A)⁺ RNA isolated from eight tissues (Hu-
Figure 2 Schematic representation of the FKRP gene. The gene consists of 4 exons and spans no more than 12 kb. The entire coding sequence is located within exon 4. The 5' UTR is blackened, the FKRP open reading frame is unblackened, and the 3' UTR is hatched.

Figure 1 Sequence of the human FKRP cDNA and analysis of the primary sequence. A, Partial nucleotide sequence and open reading frame are shown. The 4-kb cDNA encodes a protein of 495 amino acids, with a predicted molecular weight of 54.6 kDa. B, Multiple sequence alignment of fukutin homologues. Missense and nonsense mutations that were found to occur within the region aligned are indicated by red asterisks (*) and slashes (/), respectively. Amino acid residues are colored according to an 80% consensus: plus signs (+) indicate positively charged residues (H, K, and R; blue), minus signs (−) indicate negatively charged residues (D and E; red), black asterisks indicate serine or threonine residues (light blue on grey background), “a” indicates aromatic residues (F, H, W, and Y; blue on yellow background), “b” indicates big residues (E, F, H, I, L, M, Q, R, W, and Y; green on yellow), “c” indicates charged residues (DEHKR; pink), “h” indicates hydrophobic residues (A, C, D, G, H, I, L, M, T, V, W, and Y; black on yellow background), “l” indicates aliphatic residues (I, L, and V; grey on yellow), “p” indicates polar residues (C, D, E, H, K, N, Q, R, S, and T; blue), and “s” indicates small residues (A, C, D, G, N, P, S, T, and V; green). Residues that are conserved in >80% of sequences are shown as yellow on black. Residues excised from the alignment are indicated by numbers in parentheses. Predicted secondary structures are indicated below the alignment (e/E, extended or β-strand structure; h/H, helix). Residue numbers and GeneInfo identifiers (or else cosmid names) are shown following the alignment. Ca = Candida albicans; Ce = Caenorhabditis elegans; Dm = Drosophila melanogaster; HELP1 = haemolysin erythrocyte lysis protein 1; Hi = Haemophilus influenzae; Hs = Homo sapiens; Lj = Lactobacillus lactis; Pi = Prevotella intermedia; Rp = Rickettsia prowazekii; Sc = Saccharomyces cerevisiae; Sp = Streptococcus pyogenes; Tb = Trypanosoma brucei. Fukutin homologues partially encoded by ESTs in T. brucei (EST AQ635742), T. cruzi (AI077222), and Leishmania major (AQ852136), as well as a Bacillus subtilis sequence similar to L. lactis TrpF C-terminal domain (GenInfo number 1750106; bases 12108–12419), are not shown. Two C. elegans sequences, marked with a dagger (†), are alternative gene predictions that differ from published versions by the prediction of additional exons.

Genotyping
Genomic DNA was extracted from whole blood by standard methods. Linkage to the FKRP locus was assessed by genotyping subjects for the dinucleotide repeat markers D19S219 and D19S606. Genotyping was performed using an ABI 377 Automated sequencer and was analyzed using Genescan 3.1 and Genotyper 2.0 (Applied Biosystems).

Mutational Analysis
A 1.7-kb fragment containing the FKRP coding sequence was amplified using Advantage-GC Genomic program on the UK-MRC HGMP Resource Centre Web site.
Polymerase Mix (Clontech) and primers FKRP-1F (AAA-GGGATTTGAGAAAGAGC) and FKRP-5 (GCTCAGACAGCCTCTCC). PCR products were separated by agarose gel electrophoresis, were purified (Qiagen), and were used for direct sequencing. Sequencing reactions were performed using an ABI Prism BigDye Terminator Cycle Sequencing kit (Applied Biosystems) and primers FKRP-1R (GCAGGAAGAGTCTACAG), FKRP-2R (CCGAGAGTTGAAGAGGT), FKRP-3R (CTCGCTGCAGAGGTAGCC), FKRP-4R, and FKRP-5R. Sequencing products were separated on an ABI377 automated sequencer (Applied Biosystems) and analyzed using SeqEd (Applied Biosystems).

**Immunohistochemistry**

Unfixed frozen 8-mm sections were incubated with monoclonal antibodies to β-spectrin; laminins α2, β1, and γ1 (Chemicon); α- and δ-sarcoglycan (Novocastra); perlecain (Chemicon); and α-dystroglycan (VIA4-1:1,000), β-dystroglycan (Novocastra 1:25), and α- and γ-sarcoglycan (Novocastra 1:50, 1:100); and were washed and incubated with HRP-antimouse (1:50,000; Jackson). Membranes were visualized using chemiluminescence (ECL+Plus, Amersham).

**Results**

**Database Screening**

Comparison of the mouse fukutin sequence with sequence databases using TBLASTN (Altschul et al. 1997) revealed a murine expressed sequence tag (EST; AA016785) that showed significant similarity \((E = 8 \times 10^{-5})\) across the proposed fukutin active-site region. Reciprocal BLAST searches using a corrected version of this sequence as a query demonstrated its significance to fukutin homologues (Aravind and Koonin 1999), in particular Rickettsia prowazekii RP689 \((E = 2 \times 10^{-5})\) and putative orthologous sequences in Homo sapiens and Drosophila melanogaster (fig. 1b). The human version of this sequence was determined by a combination of EST assembly, reverse transcriptase PCR, and RACE (fig. 1a). The cDNA has a 1,488-bp open-reading frame that encodes a 495-amino-acid protein, which we have called fukutin-related protein (fig. 1a). Sequence analysis of FKRP predicts the presence of a hydrophobic transmembrane-spanning region (amino acids 4–28) followed by a “stem region” and the putative catalytic domain. A similar molecular organization is found in several Golgi-resident glycosyltransferases (Munro 1998).

**Radiation-Hybrid Mapping**

Radiation-hybrid mapping using the Genebridge 4 Hybridization DNA panel gave a maximum two-point LOD score of 17.25 with marker D19S219. The most likely order of markers is centromere-D19S219-FKRP-D19S606-telomere on chromosome 19q13.3. D19S219 and D19S606 are placed 3 cM apart on the Genethon map.

**Genomic Organization and Northern Blot Analysis**

The genomic organization of the FKRP gene was determined by comparing the cDNA sequence with a contiguous segment of chromosome 19 in contig NT-011166.2. The 12-kb gene is composed of three noncoding exons and a single large exon of 3.8 kb that contains part of the 5′ untranslated region and the entire open frame and 3′ untranslated region (fig. 2). Northern blot analysis demonstrated a transcript of ∼4.0 kb in all tissues studied. The FKRP transcript is expressed predominantly in skeletal muscle, placenta, and heart and relatively weakly in the remaining tissues (fig. 3). An
additional transcript of ∼3.6 kb was visible in skeletal muscle, kidney, and lung.

Mutation Analysis of the FKRP Gene

We initially studied the FKRP gene in 25 families with CMD with brain involvement. No family’s results were consistent with linkage to the FKRP locus. In addition, direct sequencing of the FKRP coding region in noninformative families showed that it was normal. We concluded that FKRP was not a major locus for these disorders, at least in our patient population.

Next, we looked at families with CMD with a secondary deficiency in laminin α2 and no brain involvement. In seven families that shared a common, severe phenotype, we identified mutations in the coding region of FKRP that are expected to affect protein function (fig. 4 and table 1). In brief, we identified 10 different mutations—8 missense and 2 nonsense. Both affected children from consanguineous families 2 and 4 had homozygous missense changes, as did the affected child from nonconsanguineous family 5. The affected children in the remaining four families were compound heterozygotes. Families 1 and 3 had one missense and one frameshift mutation, whereas families 6 and 7 had two missense mutations (table 1). In family 1, the C1154A change was present in the mother, but the A926G change was unexpectedly not found in the father’s lymphocyte DNA, suggesting a germline mutation (fig. 4). The changes were considered to be pathogenic, because they segregated with the disease in a recessive fashion and were not present in 100 control chromosomes. A small number of silent changes were identified and are listed in table 1.

Muscle Biopsy: Immunohistochemistry and Immunoblot Analysis

The skeletal muscle biopsies showed a severe dystrophic picture in all cases (a haematoxylin- and eosin-stained section from families 1 and 2 is shown in fig. 5a–b). Antibodies to α- and β-dystroglycan in families 1 and 2 showed preserved β-dystroglycan (fig. 5e–f) but a severe depletion of α-dystroglycan on most fibers in family 1 (fig. 5c), whereas >50% of fibers showed a reduction in family 2 (fig. 5d). Only the muscle from family 1 showed reduced labeling for laminin α2 (fig. 5g), β1, and γ1, and this tended to be on the larger rather than smaller fibers. In this family, only a minority of fibers with a severe depletion of α-dystroglycan were also associated with a reduction in laminin α2, β1, and γ1 labeling. However, the muscle from family 2 did not

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**Figure 4**  An example of mutation analysis in FKRP. **a**, The pedigree of family 1, showing the segregation of two heterozygous mutations, A926G and C1154A. Both affected children are compound heterozygotes, whereas the unaffected sib has inherited only one of the at-risk alleles. The father unexpectedly did not have the C1154A mutation in his lymphocyte DNA and most likely is a germline mosaic. Haplotype analysis at this and other loci confirmed paternity (data not shown). **b**, DNA sequence analysis of FKRP, identifying the mutations in family 1.
show any reduction in laminin α2 (fig. 5h), β1, and γ1 labeling, suggesting that the absence of α-dystroglycan in a large proportion of fibers is not directly associated with a reduction in these laminin chains. Immunolabeling of the membrane-associated proteins β spectrin and dystrophin, along with the glycosylated proteins α- and δ-sarcoglycan, were within normal limits. Perlecan immunolabeling in families 1 and 2 was indistinguishable from controls. A marked reduction in α-dystroglycan expression was reported in two of our patients from France, the only other families from which muscle was available.

Immunoblot analysis of α-dystroglycan in families 1 and 2, adult control muscle, and Duchenne muscular dystrophy (DMD) muscle was performed. A broad band of ~156 kDa was seen in normal muscle that was reactive with antibody V1A4-1, which recognizes a glycosylated epitope (fig. 6a). Families 1 and 2 showed a narrowing of the broad reactive band with a marked shift towards a lower molecular weight. Only minor alterations in the molecular weight of α-dystroglycan were apparent in the DMD muscle, despite the presence of a similar number of immature fibers to MDC1C, as judged by the expression of neonatal myosin (not shown). Both MDC1C muscles showed the same abnormality in α-dystroglycan on western blot. This suggests that the reduction in the mean α-dystroglycan molecular weight observed in MDC1C was not simply a reflection of an immature muscle but likely reflects differences in the pattern of glycosylation in these MDC1C patients. The reduction in β-dystroglycan in DMD was expected, given the absence of dystrophin in these patients. The molecular weight of the heavily glycosylated membrane proteins α- and γ-sarcoglycan, along with β-dystroglycan, were normal (figs. 6a–b).

**Discussion**

We have identified a new member of the fukutin family of proteins, FKRP, whose gene maps to chromosome 19q13.3. FKRP, like fukutin, contains sequence motifs identifying it as a putative sugar transferase (Aravind and Koonin 1999; Breton and Imberty 1999). Mutations in this gene give rise to a very distinct form of congenital muscular dystrophy, which previously has been recognized as a separate clinical entity (Mercuri et al. 2000; S. Quijano-Roy, L. Galan, A. Ferreiro, F. Cheliout-Heraut, F. Gray, M. Fardeau, A. Barois, P. Guicheney, N. B. Romero, and B. Estournet, unpublished data). The characteristic clinical features of this condition are onset in the first few weeks of life; severe weakness and wasting of the shoulder-girdle muscles; hypertrophy and weakness of the leg muscles, with inability to walk; calf and thigh hypertrophy; and severe restrictive respiratory involvement, leading to respiratory failure in the second decade of life. Several patients had signs of heart involvement, and this suggests that cardiac muscle is also affected in this disorder. Other invariable features were preserved intelligence, normal brain structure on imaging, and marked elevation of serum CK. A secondary reduction in laminin α2 was also observed, although this was less clear in some patients because of the severe generalized muscle damage secondary to the dystrophic muscle process. In keeping with established nomenclature, we have called this disease “MDC1C” (MDC1A represents CMD with a primary laminin α2 deficiency; MDC1B has a secondary laminin α2 deficiency and calf hypertrophy linked to 1q42). We did not observe patients with two null mutations in FKRP, a finding paralleled in FCMD, where it has been suggested that the presence of two such mutations would be embryonic lethal (Kondo-Iida et al. 1999).

Patients with MDC1C had a severe reduction in α-dystroglycan immunolabeling that was much more dramatic than the reduction in laminin α2, the only additional protein in which we could detect an altered pattern of expression. The muscle biopsy from family 1 showed that fibers deficient in laminin α2 were also deficient in α-dystroglycan immunolabeling, but the converse was not true. This suggests that whatever re-

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**Table 1**

**Mutations and Polymorphisms Found in the FKRP Gene**

<table>
<thead>
<tr>
<th>Change</th>
<th>Inheritance</th>
<th>Protein Effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mutations:</td>
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<tr>
<td>Family 1 (British):</td>
<td></td>
<td></td>
</tr>
<tr>
<td>A926G</td>
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</tr>
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<td>C1154A</td>
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</tr>
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</tr>
<tr>
<td>Family 3 (British):</td>
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<tr>
<td>C1016A</td>
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<td></td>
</tr>
<tr>
<td>C649A</td>
<td>Homozygous</td>
<td>Pro217Thr</td>
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<tr>
<td>Family 5 (Caribbean):</td>
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<td></td>
</tr>
<tr>
<td>A1394C</td>
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</tr>
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<td>G1201A</td>
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<td>Asp401Asn</td>
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Figure 5  Muscle sections, from families 1 (a) and 2 (b), stained with haematoxylin and eosin. Immunolabeling for α- and β-dystroglycan is shown in c and e, respectively (family 1), and in d and f, respectively (family 2). α-dystroglycan was markedly reduced on most fibers in family 1, whereas >50% of fibers showed a reduction in family 2. β-dystroglycan was normally expressed in all fibers in both families. Immunolabeling for laminin α2 is shown in g (family 1) and h (family 2). A reduction in laminin α2 expression on the larger fibers was evident only in family 1. Bar = 100 μm.

A relationship exists between the reduction in α-dystroglycan and laminin α2, the deficiency in α-dystroglycan occurs first. Perlecan expression was within normal limits in families 1 and 2, suggesting that other components of the basement membrane are not grossly perturbed. The normal immunolabeling of β spectrin and dystrophin confirmed that the alterations in α-dystroglycan were not due to generalized membrane damage, which is often a feature of dystrophic muscle. These observations, together with the reduction in the average weight of α-dystroglycan, raise the possibility that FKRP is involved in the glycosylation of α-dystroglycan.
Figure 6  

**a**, Immunoblot analysis of muscle proteins extracted from families 1 and 2 (F1 and F2) showed a marked shift in the average molecular weight of α- but not β-dystroglycan. Control adult muscle (CTRL) and muscle from patients with Duchenne muscular dystrophy (DMD), are included for comparison. The significant reduction in α-dystroglycan molecular weight was only observed in patients with MDC1C. 

**b**, The molecular weights of α- and γ-sarcoglycan are comparable with those of control and DMD muscle. The normal expression of the glycosylated proteins α- and γ-sarcoglycan suggest that the pathogenesis of this disorder is closely linked to the altered expression of α-dystroglycan, rather than a more generalized effect on glycosylated membrane proteins.

Dystroglycan is widely expressed, but its glycosylation pattern varies in a tissue-specific and developmental pattern (Henry and Campbell 1999; Leschziner et al. 2000), and this may provide a possible mechanism by which defects in individual glycosyltransferases can result in a tissue-specific phenotype. Many of α-dystroglycan's binding partners, including laminin α2, recognize carbohydrate moieties, and their disruption would be expected to have a profound effect on its function (Durbée et al. 1998; Montanaro et al. 1999). This is supported by the recent finding that the gene mutated in the recessive myodystrophy (myd) mouse encodes a glycosyltransferase, LARGE, and results in the altered glycosylation of α-dystroglycan (Grewal et al. 2001). The myd mouse carries an out-of-frame deletion in the LARGE gene that is predicted to give rise to a truncated protein lacking either of its two catalytic domains, in effect creating a functional "knockout." The myd phenotype is characterized by proximal muscle weakness, reduced life span, elevated serum creatine kinase, and a skeletal-muscle pathology with foci of degeneration and regeneration. These features are shared by MDC1C; however, the myd mouse also shows signs of CNS involvement (sensorineural deafness) that is not apparent in MDC1C (Mathews et al. 1995). In addition, patients with MDC1C have a variable cardiac involvement not seen in the myd mouse. Skeletal-muscle α-dystroglycan expression in the myd mouse was found to be virtually absent, using the same antibody as was used in the present study. Because this antibody recognizes a glycosylated epitope, these observations suggest that an abnormality in the glycosylation of dystroglycan may underlie the skeletal-muscle pathology seen in this animal. However, no further comparisons can be made, because data were not presented on the immunohistochemical expression of either α-dystroglycan or laminin α2 (Grewal et al. 2001), and the depletion of α-dystroglycan on immunoblotting was more marked than that observed in MDC1C. The skeletal muscle from patients with FCMD also shows a severe immunohistochemical reduction of α-dystroglycan and almost undetectable levels of α- but not β-dystroglycan, when tested by immunoblotting. The biosynthesis of glycoproteins is a complex pathway involving the controlled activity of many different enzymes in the synthesis of a variety of carbohydrate structures. It is probable that FKRP, like fukutin and LARGE, participates somewhere along one or more of these pathways in the processing of α-dystroglycan. The more severe reduction of α-dystroglycan seen in both FCMD and the myd mouse, compared with that seen in MDC1C, probably represents the differing involvement of the respective glycosyltransferase in the synthesis of the glycosylated epitope recognized by the V1A4-1 antibody.

The altered processing of α-dystroglycan in MDC1C, like FCMD and the myd mouse, is likely to be integral to the pathology in this disease. However, although we could find no evidence of other abnormalities in the processing of other proteins, such as β-dystroglycan and α- and γ-sarcoglycan, the activities of glycosyltransferases are known to be permissive, and the existence of additional substrates and the contribution to the pathology that their altered processing would produce cannot be ignored.

The altered processing of α-dystroglycan due to its abnormal glycosylation in MDC1C, together with the recent corresponding findings in FCMD and the myd mouse, identifies a novel pathogenic mechanism in muscular dystrophy and opens a new avenue of research into these diseases.
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Electronic-Database Information

Accession numbers and URLs for data in this article are as follows:

ClustalW Service at the European Bioinformatics Institute, http://www2.ebi.ac.uk/clustalw/ (for sequence alignments)
EMBL, http://www.ebi.ac.uk/emb/ (for human FKRP [accession number AJ314847])
Online Mendelian Inheritance in Man (OMIM), http://www.ncbi.nlm.nih.gov/Omim/ (for MDC1A [MIM 156225], MEB [MIM 236670], FCMD [MIM 253800], ITGA7 [MIM 600536], UCMD [MIM 254090], and MDC1B [MIM 604801])
UK-MRC HGMP Resource Centre, http://www.hgmp.mrc.ac.uk/Registered/Webapp/rhyme/index.html (for the RHyme program)

References