Aberrant Mitochondria in a Bethlem Myopathy Patient with a Homozygous Amino Acid Substitution That Destabilizes the Collagen VI α2(VI) Chain*

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Background: Collagen VI amino acid substitutions are common, and it is difficult to determine if they are pathogenic.

Results: COL6A2 p.D871N chains are abnormal and cannot assemble. Alternatively spliced chains lacking the mutation cannot functionally substitute.

Conclusion: COL6A2 p.D871N is a recessive mutation.

Significance: Protein studies reveal the consequences of amino acid substitutions in the globular domains of collagen VI.

Bethlem myopathy and Ullrich congenital muscular dystrophy (UCMD) sit at opposite ends of a clinical spectrum caused by mutations in the extracellular matrix protein collagen VI. Bethlem myopathy is relatively mild, and patients remain ambulant in adulthood while many UCMD patients lose ambulation by their teenage years and require respiratory interventions. Dominant and recessive mutations are found across the entire clinical spectrum; however, recessive Bethlem myopathy is rare, and our understanding of the molecular pathology is limited. We studied a patient with Bethlem myopathy. Electron microscopy of his muscle biopsy revealed abnormal mitochondria. We identified a homozygous amino acid substitution in the C-terminal C2 A-domain. Mutant α2(VI) chains are unable to associate with α1(VI) and α3(VI) and are degraded by the proteasomal pathway. Some collagen VI is assembled, albeit more slowly than normal, and is secreted. These molecules contain the minor α2(VI) C2a splice form that has an alternative C terminus that does include the mutation. Collagen VI tetramers containing the α2(VI) C2a chain do not assemble efficiently into microfibrils and there is a severe collagen VI deficiency in the extracellular matrix. We expressed wild-type and mutant α2(VI) C2 domains in mammalian cells and showed that while wild-type C2 domains are efficiently secreted, the mutant p.D871N domain is retained in the cell. These studies shed new light on the protein domains important for intracellular and extracellular collagen VI assembly and emphasize the importance of molecular investigations for families with collagen VI disorders to ensure accurate diagnosis and genetic counseling.

Bethlem myopathy (MIM 158810) and Ullrich congenital muscular dystrophy (UCMD; MIM 254090) sit at opposite ends of a clinical spectrum of muscular dystrophies caused by mutations in the extracellular matrix protein collagen VI (1, 2). Bethlem myopathy is a relatively mild disorder that usually becomes apparent in early childhood. Its characteristic features include proximal muscle weakness and wasting, and contractures commonly involving the finger, elbow, and ankle joints. Patients usually remain ambulant in adulthood and only rarely require mechanical ventilation at night. By contrast, UCMD is a serious congenital disorder. Severe muscle weakness, joint contractures and joint hypermobility mean that many patients lose independent ambulation by their teenage years and can succumb to early respiratory failure without effective interventions (2). A major challenge for diagnosing and counseling UCMD and Bethlem myopathy patients is that both recessive and dominant mutations are found in patients across the entire clinical spectrum (2–6) and so molecular diagnosis and an understanding of the biochemical consequences of mutations is crucial for each patient and their family.

Collagen VI is an abundant and widely expressed extracellular matrix protein that forms a microfibrillar network closely associated with basement membranes (7). The most common isoform is a heterotrimer containing one α1(VI), one α2(VI) and one α3(VI) subunit, encoded by the COL6A1, COL6A2, and COL6A3 genes, respectively. Humans have two recently iden-

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‡ The abbreviations used are: UCMD, Ullrich congenital muscular dystrophy; DAPI, 4',6-diamidino-2-phenylindole; COX, cytochrome c oxidase; SDH, succinate dehydrogenase.
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Ethical Standards—This study was approved by the Royal Children's Hospital Human Research Ethics Committee in accordance with the National Health and Medical Research Council's National Statement on Ethical Conduct in Human Research (2007).

Muscle Biopsy and Staining—Frozen sections (7 μm) were cut from sartorius muscle biopsies and stained for cytochrome c oxidase (COX) or succinate dehydrogenase (SDH) according to standard protocols (21). Some sections were immunostained with a collagen VI antibody (clone VI-26, Millipore, 1:1000 dilution) following antigen retrieval at pH 9. Specific staining was visualized using an Envision FLEX detection and amplification system (Dako). For electron microscopy muscle samples were fixed in 2.5% cacodylate-buffered glutaraldehyde containing 1% osmium tetroxide, dehydrated in an alcohol series, and embedded in Epon 812. Ultrathin sections were observed in a Zeiss Libra 120 PLUS EF transmission electron microscope.

RNA and DNA Extraction, PCR, and Sequencing—Total RNA was extracted from control and patient fibroblasts using RNeasy (Qiagen). RNA was reverse transcribed using murine leukemia virus reverse transcriptase and an oligo(dT) primer according to the manufacturer's protocol (GeneAmp RNA PCR kit; Applied Biosystems). The resulting cDNA was used as a template for PCR amplification and sequencing of the entire coding regions of the α1(VI) and α2(VI) chains and the region coding for α3(VI) domains N2-C5, as described previously (17). Genomic DNA was extracted from patient fibroblasts and from blood samples obtained from both parents. COL6A2 exon 28 containing the mutation was PCR amplified and sequenced using the forward primer 5'-AATGGAAGGGCACAGGT-GCG-3' and the reverse primer 5'-CTTAGCACCATGG-ACGGGGT-3'.

Collagen VI Biosynthetic Labeling and Immunoprecipitation—Patient and control fibroblasts were grown to confluence in 10 cm² dishes and incubated overnight in DMEM with 10% FBS containing 0.25 mM sodium ascorbate. Fibroblasts were radiolabeled for 18 h with 100 mCi/ml [35S]methionine (Trans-3S-label 1032 Ci/mmol; ICN Pharmaceuticals, Inc.) in 750 μl of serum-free and methionine-free DMEM containing 0.25 mM sodium ascorbate. The cell and medium fractions were harvested, and collagen VI was immunoprecipitated with an α3(VI) N1 domain antibody and analyzed under reducing and nonreducing conditions as described previously (3). For pulse-chase analysis, the fibroblasts were radiolabeled, as above, for 30 min, and chased for 0, 0.5, 1, 2, 3, 4, 6, or 8 h in serum-free DMEM containing 0.25 mM sodium ascorbate and 10 mM L-methionine. Collagen VI was immunoprecipitated from cell and medium fractions with an α3(VI) N1 domain antibody as described above. In some experiments, fibroblasts were pre-
incubated for 2 h with the protease inhibitors 5 \( \mu \text{M} \) Z-Leu-Leu-Leu-al (MG132, Sigma), 5 \( \mu \text{M} \) clasto-Lactacystin \( \beta \)-lactone (Sigma), 10 \( \mu \text{M} \) trans-epoxysuccinyl-1-leucylamido(4-guanidino)butane (E64, Sigma), or 1 \( \mu \text{g/ml} \) pepstatin A (Sigma), then labeled and chased in the presence of the same protease inhibitors. Cell layer samples were immunoprecipitated with an \( \alpha_2(\text{VI}) \) N1 domain antibody and analyzed as above.

**Fibroblast Immunostaining**—Fibroblasts were grown to confluence in four-well chamber slides and supplemented daily with 0.25 M sodium ascorbate for 2 days. The extracellular matrix was stained for collagen VI, and the cell nuclei were stained with 4',6-diamidino-2-phenylindole (DAPI) using methods described previously (3).

**Negative Staining Electron Microscopy**—Confluent fibroblasts in 10-cm² dishes were incubated for 18 h in serum-free DMEM containing 0.25 M sodium ascorbate. The medium was collected and clarified by centrifugation. Microfibrils were deposited on a carbon-coated electron microscopy grid, stained for 30 s with 1% aqueous uranyl acetate, blotted, and quickly air dried. Grids were observed at 300 kV using a Tecnai F30 (FEI, Eindhoven) and micrographs were acquired with an Ultrascan 1000 (Gatan, Pleasanton, CA).

**Constructing Expression Vectors Containing the \( \alpha_2(\text{VI}) \) Wild Type and p.D871N Mutant C2 Domains**—Wild type and mutant \( \alpha_2(\text{VI}) \) C2 domains (nucleotides 2464–3057; amino acid residues 822–1019) were amplified from control and UCMD65 cDNA using a forward primer containing a 5' PvuII restriction enzyme site followed by the StrepII tag sequence (22), and a reverse primer containing a 3' HindIII restriction enzyme site. PCR products were purified and sequentially digested with PvuII and HindIII (New England Biolabs). The fragments were ligated into the episomal expression vector pCEP4-BM40-HisEK that had also been digested with PvuII and HindIII. This vector has previously been modified from the pCEP4 vector (Invitrogen) to contain a cleavable BM40 secretory signal peptide at the N terminus of the multiple cloning site (23). Selected clones were sequenced with a pCEP4 specific primer (Invitrogen) to contain a Precision Plus Protein Standard (Bio-Rad) then transferred to nitrocellulose membranes (GE Healthcare). Membranes were blocked in 5% BSA (w/v) in PBS for 1 h at room temperature, washed four times in PBS and then incubated at room temperature for 1 h in PBS containing a 1:10,000 dilution of horseradish peroxidase-conjugated anti-mouse antibody (Dako). Proteins were detected by incubating membranes with Amersham Biosciences Plus ECL Western blotting detection reagents (GE Healthcare Life Sciences) for 5 min and visualized by exposure to Amersham Biosciences high performance chemiluminescent film (GE Healthcare Life Sciences).

**Protein Homology Modeling**—A three-dimensional protein homology model of the human \( \alpha_2(\text{VI}) \) C2 domain (amino acids 831–1014) was generated using the EasyModeller 4.0 graphical user interface (24) to the MODELLER 9.12 program. The best template for the model, \( \alpha_X \beta_2 \) integrin (1N3Y.pdb), was identified by BLAST searching the PDB sequence database. The final model was visualized using Discovery Studio Visualizer (Accelrys.com).

**RESULTS**

**Case Report**—UCMD65 (Fig. 1), is a 61-year-old male with slowly progressive proximal limb muscle weakness. He has no family history of muscle disease but his parents are second-degree relatives. From the first to third decade our patient complained of clumsiness while playing sport or running but could walk and work normally. In his thirties, he developed gait dis-
turbance with bilateral genu valgus deformities and underwent a tibial osteotomy. When he was 40 years old, he noticed decreased proximal muscle strength and muscle atrophy in his arms that made combing his hair difficult. At age 60, muscle weakness had progressed to his legs, he had difficulty walking, and had begun to use an orthopaedic walker. Simultaneously, he complained about joint contractures in his upper and lower limbs, which restricted his movements, and occasional difficulty swallowing liquids. A neurological examination at age 61 showed proximal weakness with symmetrical muscle atrophy involving all limbs, absent osteotendinous reflexes, and joint contractures in elbows, wrists, fingers, and knees. No skin alterations or sensory disturbances were noted. His gait was unsteady. Electromyography revealed reduced amplitude and frequency of the motor unit potentials in vastus medialis, iliopsoas, deltoids, and right tibialis anterior muscles. Blood tests were normal except for mildly elevated creatine phosphokinase (CK, 149 – 161 units/liter). Muscle magnetic resonance imaging (MRI) showed generalized muscle atrophy with diffuse muscular fatty infiltration, predominantly in proximal limb muscles. Our patient’s clinical presentation thus suggested a mitochondrial myopathy, Emery-Dreifuss muscular dystrophy, or Bethlem myopathy.

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FIGURE 2. UCMD65 muscle pathology. a, UCMD65 had COX negative fibers (*) indicative of a mitochondrial myopathy. b, SDH staining was normal suggesting there was no mitochondrial proliferation. c, electron microscopy revealed frequent mitochondrial paracrystalline inclusions (second panel from left). The panel on the right shows an enlargement of the boxed area to visualize the paracrystalline inclusions. Normal mitochondria in a control sample are shown in the panel on the left. d, immunostaining with a collagen VI specific antibody shows abundant collagen VI surrounding each muscle fiber in the control (nonspecific myopathy); however, collagen VI was not detected in UCMD65 muscle. UCMD65 also had many central nuclei (black arrows).

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Muscle Biopsy—To explore UCMD65 muscle pathology we stained frozen muscle sections for mitochondrial markers cytochrome c oxidase (COX) (Fig. 2a) and succinate dehydrogenase (SDH) (Fig. 2b). In normal muscle virtually all fibers have COX staining with type I fibers staining more intensely than type II reflecting their increased number of mitochondria. UCMD65 muscle had COX negative fibers indicating mitochondrial dysfunction; however, we did not see fibers with increased SDH staining suggesting the disorder was not a primary mitochondrial myopathy (Fig. 2b). Consistent with this, we screened for the common mitochondrial DNA mutations found in MELAS syndrome (mitochondrial encephalopathy, lactic acidosis, and stroke-like episodes), m.3243A>G, m.11084A>G, m.3271T>G, and m.3291T>G; MERRF syndrome (myoclonic epilepsy with ragged red fibers), m.8344A>G and m.8356T>C; and Leigh syndrome, m.8993T>C, and found no alterations (data not shown).

Transmission electron microscopy confirmed the mitochondrial abnormality detected by COX staining. We saw frequent mitochondria with paracrystalline inclusions (Fig. 2c) which are typically found in muscle from patients with mitochondrial myopathies. Having ruled out the most common mitochondrial myopathies, we reasoned that since mitochondria with abnor-
mal cristae are present in skeletal muscle from collagen VI knock-out mice (25), UCMD65 may have a collagen VI disorder, and so we stained skeletal muscle with a collagen VI antibody. There was abundant collagen VI staining around each muscle fiber in the control biopsy (nonspecific myopathy) but collagen VI was not detected in UCMD65 muscle (Fig. 2d), confirming that this patient has a collagen VI disorder. UCMD65’s clinical presentation, at the mild end of the collagen VI disorder spectrum, indicates that he has Bethlem myopathy; however, absent collagen VI staining is usually associated with severe Ullrich congenital muscular dystrophy (26–28) and has not been reported in Bethlem myopathy.

UCMD65 Has a Homozygous COL6A2 Mutation—To characterize the mutation responsible for Bethlem myopathy in this patient we RT-PCR amplified and sequenced the entire coding regions of the α1(VI) and α2(VI) chains and the region coding for domains N3-C5 of the α3(VI) chain. UCMD65 had a homozygous c.2611G>A mutation in α2(VI) causing a p.D871N amino acid substitution. We confirmed the mutation by sequencing genomic DNA PCR products from UCMD65 and his unaffected mother and father who were both heterozygous carriers of the mutation (Fig. 3). The mutation is in the α2(VI) C2 domain and will be present in the most common α2(VI) splice variant, but not the minor α2(VI) C2a form which has an alternative smaller C2 domain (Fig. 3b).

The α2(VI) p.D871N Mutation Interferes with Collagen VI Assembly, Secretion, and Microfibril Formation—We then examined the effects of the mutation on collagen VI production by UCMD65 fibroblasts. In overnight biosynthetic labeling and immunoprecipitation experiments we consistently found that UCMD65 cells secreted less collagen VI than control fibroblasts (Fig. 4, a and b); however, we noted that under reducing conditions, which allow us to visualize the individual collagen VI chains, the α2(VI) C2a band was a similar intensity in control and UCMD65 medium (Fig. 4a). This suggested that the alternatively spliced α2(VI) C2a chains were able to assemble normally with α1(VI) and α3(VI) to form collagen VI monomers, but assembly and secretion of α2(VI) chains with the p.D871N mutation were impaired. Consistent with this, using quantitative RT-PCR we confirmed that the α1(VI), α2(VI) C2, and α2(VI) C2a mRNAs were all expressed at normal levels (data not shown). As in control cells, collagen VI tetrayers were the major secreted form of collagen VI (Fig. 4b) and there was no intracellular accumulation of collagen VI triple helical monomers or dimers (Fig. 4b) indicating that the mutation most likely prevented incorporation of the chains into monomers. Outside the cell collagen VI tetrayers connect end-to-end to form microfibrils which can be deposited into the extracellular matrix surrounding cells. Control fibroblasts accumulate an extensive collagen VI extracellular matrix but, by contrast, the UCMD65 extracellular matrix contained only a very small amount of collagen VI (Fig. 4c).

Having established that the homozygous α2(VI) p.D871N mutation ultimately leads to a severe collagen VI matrix deficiency, we then examined all the stages of collagen VI assembly in detail to establish the molecular mechanisms responsible for this deficiency. We first used pulse-chase experiments where cells were biosynthetically radiolabeled for 30 min, chased for up to 8 h and collagen VI immunoprecipitated with an α3(VI) chain antibody, to establish the time-course of collagen VI intracellular assembly and secretion. In control cells, α1(VI) and α2(VI) chains were co-immunoprecipitated with α3(VI) from 0.5 h (Fig. 5a). On non-reduced gels collagen VI triple helical monomers, dimers and tetrayers were all present at 0.5 h of chase (Fig. 5b) indicating that once the three chains come together further assembly is rapid. Collagen VI was present in the medium at 1 h of chase and most of the collagen VI synthesized during the 30 min pulse was secreted by two 2 hours of chase (Fig. 5, a and b). By contrast, collagen VI assembly and secretion was reduced in UCMD65. Some α1(VI) and α2(VI) C2a chains co-immunoprecipitated with α3(VI) at 0.5 h of chase (Fig. 5c) but these bands were very feint when compared with the α3(VI) chains and the relative intensities of these bands in the control cells (compare intracellular α3(VI) with α1(VI) and α2(VI) in Fig. 5, a and c). These data confirm that...
assembly of the mutant α2(VI) chains into disulfide bonded monomers with α1(VI) and α2(VI) was indeed disrupted in UCMD65 and suggest that assembly of the minor α2(VI)-C2a chain without the mutation proceeded normally.

Following secretion, collagen VI tetramers associate end-to-end to form microfibrils. We visualized and quantitated microfibrils in the medium of control and UCMD65 fibroblasts using negative staining electron microscopy (Fig. 6a–d). Around 60% of the collagen VI microfibrils in the medium of control cells contained more than 5 tetramers (Fig. 6e); however, there were very few of these large microfibrils in UCMD65 medium and around 40% of the microfibrils contained only a single tetramer (Fig. 6, b–e), indicating that end-to-end assembly of the tetramers secreted by UCMD65 fibroblasts was severely compromised.

α2(VI) p.D871N Chains Are Degraded by the Proteasome—We then looked at the fate of the mutant α2(VI) p.D871N chains in pulse-chase experiments. UCMD65 fibroblasts were preincubated for 2 h with and without inhibitors of lysosomal or proteasomal protein degradation, biosynthetically radiolabeled for 30 min, then chased for up to 6 h. The mutant α2(VI) p.D871N and alternatively spliced α2(VI)-C2a chains were precipitated with an antibody raised to the α2(VI) N1 domain and analyzed under reducing conditions. In the absence of protease inhibitors the mutant α2(VI) chains were almost completely eliminated by 6 h of chase (Fig. 7); however, mutant α2(VI) chains were protected by MG132 and clasto-Lactocystin β-lactone (Fig. 7) indicating that they are normally degraded by proteasomal proteases. The lysosomal protease inhibitors E64 and pepstatin A did not protect the mutant α2(VI) p.D871N chains from degradation. This is consistent with the conclusion that mutant α2(VI) p.D871N chains are selectively degraded by the proteasome.

Mutant α2(VI) C2 Domains Are Not Secreted—To explore the effect of the p.D871N mutation on protein folding we expressed recombinant Strep II tagged wild-type and mutant α2(VI) C2 domains in HEK293-EBNA cells. We separated secreted proteins, soluble intracellular proteins and intracellular proteins that had formed aggregates by SDS-PAGE and detected recombinant C2 domains by immunoblotting with a Strep II antibody. Recombinant wild-type domains were efficiently secreted as expected for a correctly folded protein (Fig. 8a). By contrast, C2 domains containing the p.D871N mutation were predominantly found in the soluble intracellular fraction and were not detected in the medium. This indicates that the mutant domains are recognized as abnormal by endoplasmic reticulum quality control mechanisms and are retained in the cell.

The C2 domain contains 2 potential N-linked glycosylation sites, N897 in the sequence NLT and N954 in the sequence NDS, both are located in loop regions on the outside of the folded protein structure (Fig. 8b) and our data indicate that both can be substituted with an N-linked carbohydrate. When we expressed C2 domains with a deletion of Asn-897 we saw an unglycosylated form and a larger form that is glycosylated at N954 (Fig. 8a). We also detected two forms of the wild-type C2 domain (Fig. 8a). The smaller faster migrating band co-migrated with the glycosylated Asn-897 band indicating that it is folded protein structure (Fig. 8b) and our data indicate that both can be substituted with an N-linked carbohydrate. When we expressed C2 domains with a deletion of Asn-897 we saw an unglycosylated form and a larger form that is glycosylated at N954 (Fig. 8a). We also detected two forms of the wild-type C2 domain (Fig. 8a). The smaller faster migrating band co-migrated with the glycosylated Asn-897 band indicating that it is

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FIGURE 4. Collagen VI production in UCMD65 fibroblasts. Control and UCMD65 fibroblasts were labeled overnight with [35S]methionine and collagen VI was immunoprecipitated from the cell (C) and medium (M) with an α3(VI) N1 domain antibody. a, immunoprecipitated collagen VI was analyzed under reducing conditions on 3–8% Tris acetate gels to resolve the α1(VI), α2(VI), and α3(VI) chains. Collagen VI tetramers and fibronectin dimers (Fn2) are indicated. There was no accumulation of collagen VI monomers or dimers in UCMD65 cells indicating that dimer and tetramer assembly were not affected. c, immunostaining of the collagen VI matrix deposited by fibroblasts. Control and UCMD65 fibroblasts were grown for 2 days post-confluence in the presence of sodium ascorbate and stained with a collagen VI antibody. The cell nuclei were stained with DAPI. Control fibroblasts deposited an extensive collagen VI network; however, the collagen VI matrix was fine and sparse in UCMD65 cultures.

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ever, there were very few of these large microfibrils in UCMD65 medium and analyzed under reducing conditions. In the absence of protease inhibitors the mutant α2(VI) chains were almost completely eliminated by 6 h of chase (Fig. 7); however, mutant α2(VI) chains were protected by MG132 and clasto-Lactocystin β-lactone (Fig. 7) indicating that they are normally degraded by proteasomal proteases. The lysosomal protease inhibitors E64 and pepstatin A did not protect the mutant α2(VI) p.D871N chains from degradation. This is consistent with the conclusion that mutant α2(VI) p.D871N chains are selectively degraded by the proteasome.

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FIGURE 5. Pulse-chase analysis of collagen VI assembly and secretion. Control (a and b) and UCMD65 (c and d) fibroblasts were labeled with [35S]methionine for 30 min and chased for up to 8 h. Collagen VI in the cell and medium was immunoprecipitated with an α3(VI) N1 domain antibody and analyzed under reducing conditions on 3–8% Tris acetate gels to resolve the α1(VI), α2(VI), and α3(VI) chains (a and c) and non-reducing conditions to compare chain assembly (b and d). In the control, α1(VI) and α2(VI) chains were co-precipitated with α3(VI) from 0.5 h and collagen VI appeared in the medium from 1 h of chase. By contrast, in UCMD 65 co-precipitating α1(VI) and α2(VI) chains were barely visible in the cell throughout the chase period, and small amounts of secreted collagen VI first appeared in the medium at 2 h of chase. To allow the collagen VI produced by UCMD65 to be clearly seen the gels shown in panels c and d were exposed for three times longer than the control gels in panels a and b. Collagen VI α1(VI), α2(VI), α3(VI), and α2(VI) C2a chains, and contaminating fibronectin (Fn) are indicated in panels a and c. Collagen VI monomers, dimers, tetramers, and contaminating fibronectin dimers (Fn2) are marked in panels b and d.

with paracrystalline inclusions by electron microscopy. Similar mitochondrial changes are seen in collagen VI knock-out mice (25), and have also been reported in a patient with an autosomal dominant benign limb-girdle myopathy with contractures who fitted the clinical criteria for Bethlem myopathy (29). This patient did not have a molecular diagnosis as the report was prior to the discovery that collagen VI mutations were the underlying cause of Bethlem myopathy (30). The patient also had COX negative fibers similar to our patient and had similar mild and very slowly progressing muscle wasting and weakness. Together these data suggest that COX negative fibers and structural mitochondrial abnormalities might be common in Bethlem myopathy muscle.

UCMD65, has a homozygous COL6A2 p.D871N mutation which is in the α2(VI) C2 A-domain. Our data show that the mutant α2(VI) C2 domain is misfolded and the mutant α2(VI) chains are not able to assemble normally with α1(VI) and α3(VI). When compared with control fibroblasts, UCMD65 fibroblasts secrete much less collagen VI and the minor alternatively spliced α2(VI) C2a product, which does not contain the mutation, is a major component of the UCMD65 secreted assemblies. While the α2(VI) C2a chain is clearly able to assemble with α1(VI) and α2(VI), the kinetics are slower than assembly of the major α2(VI) splice form and collagen VI secretion is delayed by approximately one hour. The α2(VI) p.D871N mutation was reported previously in a patient with recessive Bethlem myopathy who had a premature stop codon that resulted in nonsense-mediated mRNA decay on the other COL6A2 allele (4). In that case the authors showed collagen VI secretion was reduced but didn’t detect the α2(VI) C2a chain when they immunoprecipitated collagen VI with an α1(VI) antibody and then immunoblotted with an antibody that recognizes all three collagen VI chains. Two factors could explain these seemingly contradictory findings. Firstly, the patient in the earlier report would have been making half the normal amount of the α2(VI) C2a chain because of mRNA decay of the products from one allele; and secondly, it is likely that radiolabeling is a more sensitive technique for detecting minor protein products than immunoblotting.

Our data fit closely with the demonstrated effects of other α2(VI) C2 amino acid substitutions on collagen VI biosynthesis and intracellular assembly. For example, a homozygous α2(VI) p.R876S mutation destablizes the mutant chain and the small amount of secreted collagen VI contains the α2(VI) C2a chain (31). We reported a patient with recessive α2(VI) C2 domain mutations, p.L873P and p.N897del, and showed that the mutant chains were unstable and only small amounts of collagen VI were secreted (3). At the time we didn’t recognize that the secreted collagen VI contained the α2(VI) C2a chain but more recent analyses clearly show that this is the case (data not shown). Our data showing that the p.N897del C2 domain is misfolded and retained in the cell suggests that the consequences of all these α2(VI) C2 domain amino acid substitutions, p.L873P, p.N897del and p.R876S are very similar to the p.D871N mutation reported here. It is surprising then, that the clinical presentation of these patients is quite different. UCMD65 reported here (p.D871N; p.D871N) and BM2 (p.Q819X; p.D871N; (4)) both have classical mild Bethlem
myopathy and retained independent ambulation well into adulthood, while UCMD2 (p.L873P; p.N897del; (3)) and UC15 (p.R876S; p.R876S; (31)) had a more severe disorder consistent with UCMD and lost ambulation at 10 and 11 years of age, respectively. While we don’t have a definitive explanation for why mutations that have a similar effect on collagen VI biosynthesis and assembly have different clinical outcomes it is possible that uncharacterized modifying genes are expressed differently in these patients or that subtle differences in the ability of the mutant chains to be incorporated into tetramers and microfibrils modulate the phenotype.

FIGURE 6. Collagen VI microfibril formation is compromised in UCMD65. Negative staining electron micrographs of collagen VI microfibrils in the medium from control (a) and UCMD65 fibroblasts (b–d). The white arrowheads in a indicate end-to-end association of the tetramers. e, collagen VI secreted into the medium of control (white bars) and UCMD65 (black bars) fibroblast was visualized by negative staining electron microscopy, and the ability of the tetramers to associate end-to-end was quantitated. The occurrence of microfibrils containing 1–5 or >5 tetramers is shown as a percentage of the total number of microfibrils. Tetramer-tetramer association was severely compromised in UCMD65 compared with the control.

The role of the minor α2(VI) C2a splice variant in normal tissues has not yet been defined. It assembles with the other collagen VI chains into triple helical monomers more slowly than α2(VI) chains and outside the cell tetramers containing the C2a splice variant cannot assemble end-to-end into microfibrils efficiently and only small amounts of this collagen VI isoform are deposited into the extracellular matrix. These data emphasize the critical importance of the α2(VI) C2 domain in monomer and microfibril assembly. Further, electron microscopy of the collagen VI fibrils deposited by the patient with compound heterozygous mutations, α2(VI) p.Q819X and p.D871N, in which only α2(VI) C2a chains are likely to be incorporated into secreted collagen VI molecules, revealed that the microfibrils in the extracellular matrix did not develop the extensive interconnections seen control microfibrils (4). This finding suggests that the α2(VI) C2a domain is also important in promoting interactions between collagen VI microfibrils and raises the possibility that inclusion of this domain in collagen VI tetramers might modulate collagen VI microfibril formation and interactions with other extracellular matrix molecules.

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FIGURE 7. Pulse-chase analysis of α2(VI) chain degradation. UCMD65 (a) and control fibroblasts (b) were labeled with [35S]methionine for 30 min and chased for up to 6 h. Intracellular collagen VI was immunoprecipitated with an α2(VI) N1 domain antibody that recognizes the α2(VI) and the α2-C2a chains, and analyzed under reducing conditions on 3–8% Tris acetate gels. When used, proteasome inhibitors MG132 and clasto-Lactocystin β-lactone (clasto), and lysosomal enzyme inhibitors E64 or pepstatin A (pep A) were added 2 h before the experiment and included in the labeling and chase medium. In the absence of inhibitors (none) UCMD65 α2(VI) chains are almost completely eliminated by 6 h of chase; however, α2(VI) chain degradation is almost completely prevented by the proteasome inhibitors MG132 and clasto-Lactocystin β-lactone. The lysosomal protease inhibitors E64 and pepstatin A do not protect UCMD65 α2(VI) chains from degradation. The no primary antibody control (no Ab) lanes show the fibronectin that binds non-specifically to the protein A-Sepharose beads. Protease inhibitors do not affect clearance of α2(VI) chains from control fibroblasts.
We explored how the p.D871N mutation and N897del mutations affected the structure and trafficking of the α2(VI) C2 domain by expressing wild-type and mutant C2 domains in transfected mammalian cells. Our experiments are the first confirmation that the two consensus sites for N-linked glycosylation in the C2 domain are both used. While wild-type C2 domains were efficiently secreted, both mutant domains were retained in the cell suggesting that they are misfolded and held in the ER by quality control mechanisms. This probable misfolding likely explains why the mutant α2(VI) chains found and were eliminated from patient fibroblasts by proteolysis that could be prevented by two drugs that inhibit proteasomal degradation. Lysosomal protease inhibitors did not slow degradation of the mutant α2(VI) chains. This indicates that the mutant α2(VI) chains likely remain soluble and are transported out of the ER for degradation by the proteasome (ERAD), and that the alternative autophagy/lysosomal pathway is not activated. Collagen type I proα1(I) chains that are unassembled because they have a mutation in the C-terminal propeptide, the domain critical for intracellular chain association, are also retro-translocated to the cytoplasm for degraded by the proteasome (37, 38). This may be a common pathway for degrading collagen subunits with mutations that prevent the initial step of collagen assembly, chain recognition and triple helix formation.

Our detailed study into the functional consequences of this α2(VI) C2 domain amino acid substitution p.D871N sheds new light on the requirements for intracellular and extracellular collagen VI assembly and the cellular degradation pathway responsible for eliminating the mutant chains that are unable to assemble. It also emphasizes the critical importance of molecular diagnosis for each family as patients with very different clinical presentations can have similar mutations that affect collagen VI protein in similar ways. Further detailed studies on molecular mechanisms of the collagen VI disorders are needed to understand the factors contributing to this clinical enigma.

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REFERENCES
1. Allamand, V., Bríñas, L., Richard, P., Stojkovic, T., Quijano-Roy, S., and Bonne, G. (2011) CollVI myopathies: where do we stand, where do we go? Skelet Muscle 1, 30
2. Bönnemann, C. G. (2011) The collagen VI-related myopathies: muscle meets its matrix. Nat. Rev. Neurol. 7, 379–390
3. Baker, N. L., Mörgelin, M., Peat, R., Goemans, N., North, K. N., Bateman, J. F., and Lamandé, S. R. (2005) Dominant collagen VI mutations are a common cause of Ullrich congenital muscular dystrophy. Hum. Mol. Genet. 14, 279–293
4. Gualandi, F., Urciuolo, A., Martoni, E., Sabatelli, P., Squarzoni, S., Bovo-lenta, M., Messina, S., Mercuri, E., Franchella, A., Ferlini, A., Bonaldo, P., and Merlini, L. (2009) Autosomal recessive Bethlem myopathy. Neurology 73, 1883–1891
5. Merlini, L., Martoni, E., Grumati, P., Sabatelli, P., Squarzoni, S., Urciuolo, A., Ferlini, A., Gualandi, F., and Bonaldo, P. (2008) Autosomal recessive myosclerosis myopathy is a collagen VI disorder. Neurology 71, 1245–1253
6. Foley, A. R., Hu, Y., Zou, Y., Columbus, A., Shoffner, J., Dunn, D. M., Weiss, R. B., and Bönnemann, C. G. (2009) Autosomal recessive inheritance of classic Bethlem myopathy. Neuromuscul. Disord. 19, 813–817
7. Keene, D. R., Engvall, E., and Glanville, R. W. (1988) Ultrastructure of type VI collagen in human skin and cartilage suggests an anchoring function for this filamentous network. J. Cell Biol. 107, 1995–2006
8. Fitzgerald, J., Rich, C., Zhou, F. H., and Hansen, U. (2008) Three novel collagen VI chains, α4(VI), α5(VI), and α6(VI). J. Biol. Chem. 283,
Abnormal Mitochondria and Protein Instability in Bethlem Myopathy

9. Gara, S. K., Grumati, P., Urciuolo, A., Bonaldo, P., Kobbe, B., Koch, M., Paulsson, M., and Wagener, R. (2008) Three novel collagen VI chains with high homology to the α3 chain. J. Biol. Chem. 283, 10658–10670

10. Whittaker, C. A., and Hynes, R. O. (2002) Distribution and evolution of von Willebrand/integrin A domains: widely dispersed domains with roles in cell adhesion and elsewhere. Mol. Biol. Cell 13, 3369–3387

11. Stokes, D. G., Saitta, B., Timpl, R., and Chu, M. L. (1991) Human α3(VI) collagen gene. Characterization of exons coding for the amino-terminal globular domain and alternative splicing in normal and tumor cells. J. Biol. Chem. 266, 8626–8633

12. Zanussi, S., Doliina, R., Segat, D., Bonaldo, P., and Colombatti, A. (1992) The human type VI collagen gene. mRNA and protein variants of the α3 chain generated by alternative splicing of an additional 5′-end exon. J. Biol. Chem. 267, 24082–24089

13. Timpl, R., and Chu, M. L. (1994) Microfibrillar collagen type VI in Extracellular Matrix Assembly and Structure (Yurchenco, P. D., Birk, D., and Mecham, R. P., eds), pp. 207–242, Academic Press, Orlando

14. Butterfield, R. J., Foley, A. R., Dastgir, J., Asman, S., Dunn, D. M., Zou, Y., Hu, Y., Flanigan, K. M., Svoboda, K. I., Winder, T. L., Weiss, R. B., and Bonnemann, C. G. (2013) Position of glycan substitutions in the triple helix of COL6A1, COL6A2, and COL6A3 is correlated with severity and mode of inheritance in collagen VI myopathies. Hum. Mutat. 34, 1558–1567

15. Pace, R. A., Peat, R. A., Baker, N. L., Zamurs, L., Mörgelin, M., Irving, M., Adams, N. E., Bateman, J. F., Mowat, D., Smith, N. J., Lamont, P. J., Moore, S. A., Mathews, K. D., North, K. K., and Lamandé, S. R. (2008) Collagen VI glycine mutations: perturbed assembly and a spectrum of clinical severity. Ann. Neurol. 64, 294–303

16. Lamandé, S. R., Shields, K. A., Kornberg, A. J., Shield, L. K., and Bateman, J. F. (1999) Bethlem myopathy and engineered collagen VI triple helical deletions prevent intracellular multimer assembly and protein secretion. J. Biol. Chem. 274, 21817–21822

17. Baker, N. L., Mörgelin, M., Pace, R. A., Peat, R. A., Adams, N. E., Gardner, R. J., Rowland, L. P., Miller, G., De Jonghe, P., Ceulemans, B., Hannibal, M. C., Edwards, M., Thompson, E. M., Jacobson, R., Quinlivan, R. C., Afrámos, S., Kornberg, A. J., North, K. K., Bateman, J. F., and Lamandé, S. R. (2007) Molecular consequences of dominant Bethlem myopathy collagen VI mutations. Ann. Neurol. 62, 390–405

18. Jimenez-Mallebrera, C., Mäioli, M. A., Kim, J., Brown, S. C., Feng, L., Lampe, A. K., Bushby, K., Hicks, D., Flanigan, K. M., Bonnemann, C., Sewry, C. A., and Muntoni, F. (2006) A comparative analysis of collagen VI production in muscle, skin and fibroblasts from 14 Ullrich congenital muscular dystrophy patients with dominant and recessive COL6A1 mutations. Neuromuscul. Disord. 16, 571–582

19. Briñas, L., Richard, P., Quijano-Roy, S., Gartioux, C., Ledeuil, C., Lacène, J. F., and Hynes, R. O. (2002) Kinked collagen VI tetramers and reduced microfibril formation in a patient with Bethlem myopathy and introduced triple helical glycine mutations. J. Biol. Chem. 277, 1949–1956

20. Old, S. L., and Johnson, M. A. (1989) Methods of microphotometric assay of succinate dehydrogenase and cytochrome c oxidase activities for use on human skeletal muscle. Histochem. J. 21, 545–555

21. Schmidt, T. G., and Skerra, A. (2007) The Strep-tag system for one-step purification and high-affinity detection or capturing of proteins. Nat. Protoc. 2, 1528–1535

22. Wilson, R., Freddi, S., and Bateman, I. F. (2002) Collagen X chains harboring Schmid metaphyseal chondrodysplasia NCI domain mutations are selectively retained and degraded in stably transfected cells. J. Biol. Chem. 277, 12516–12524

23. Kuntal, B. K., Aparoy, P., and Reddanna, P. (2010) EasyModeller: A graphical interface to MODELLER. BMC Res. Notes 3, 226

24. Irwin, W. A., Bergamin, N., Sabatelli, P., Reggiani, C., Megghian, A., Merlino, L., Braghetta, P., Columbaro, M., Volpin, D., Bressan, G. M., Bernardi, P., and Bonaldo, P. (2003) Mitochondrial dysfunction and apoptosis in myopathic mice with collagen VI deficiency. Nat. Genet. 35, 367–371

25. Higuchi, I., Shiraishi, T., Hashiguchi, T., Suehara, M., Niyama, T., Naka- gawa, M., Arimura, K., Maruyama, I., and Osame, M. (2001) Frasemesh mutation in the collagen VI gene causes Ullrich's disease. Ann. Neurol. 50, 261–265

26. Camacho Vanegas, O., Bertini, E., Zhang, R. Z., Petrinì, S., Minossi, C., Sabatelli, P., Giusti, B., Chu, M. L., and Pepe, G. (2001) Ullrich scleroatonic muscular dystrophy is caused by recessive mutations in collagen type VI. Proc. Natl. Acad. Sci. U.S.A. 98, 7516–7521

27. Brockington, M., Brown, S. C., Lampe, A., Yuva, Y., Feng, L., Jimenez-Mallebrera, C., Sewry, C. A., Flanigan, K. M., Bushby, K., and Muntoni, F. (2004) Prenatal diagnosis of Ullrich congenital muscular dystrophy using haplotyping analysis and collagen VI immunocytochemistry. Prenat. Diagn. 24, 440–444

28. Malandrinì, A., Scarpini, C., Fabrizi, G. M., Parrotta, E., Salvadori, C., and Guazzi, G. C. (1995) Early-onset benign limb-girdle myopathy with contractures and facial involvement affecting a father and daughter. J. Neurol. Sci. 132, 195–200

29. Jobis, G. I., Keizers, H., Vreijling, J. P., de Visser, M., Speer, M. C., Wolter- man, R. A., Baas, F., and Bolhuis, P. A. (1996) Type VI collagen mutations in Bethlem myopathy, an autosomal dominant myopathy with contractures. Nat. Genet. 14, 113–115

30. Zhang, R. Z., Zou, Y., Pan, T. C., Markova, D., Fertala, A., Hu, Y., Squarzoni, S., Reed, U. C., Marie, S. K., Bönnemann, C. G., and Chu, M. L. (2010) Recessive COL6A2 C-globular missense mutations in Ullrich congenital muscular dystrophy: role of the C2α splice variant. J. Biol. Chem. 285, 10005–10015

31. Tooley, L. D., Zamurs, L. K., Beecher, N., Baker, N. L., Peat, R. A., Adams, N. E., Bateman, J. F., North, K. N., Baldock, C., and Lamandé, S. R. (2010) Collagen VI microfibril formation is abolished by an α2(VI) von Willebrand factor type A domain mutation in a patient with Ullrich congenital muscular dystrophy. J. Biol. Chem. 285, 33567–33576

32. Torimitsu, T., and Klionsky, D. J. (2007) Endoplasmic reticulum stress: a new pathway to induce autophagy. Autophagy 3, 160–162

33. Meusser, B., Hirsch, C., Jarosch, E., and Sommer, T. (2005) ERAD: the long road to destruction. Nature Cell Biology 7, 766–772

34. Ding, W. X., and Yin, X. M. (2008) Sorting, recognition and activation of the misfolded protein degradation pathways through macroautophagy and the proteasome. Autophagy 4, 141–150

35. Bateman, J. F., Boot-Handford, R. P., and Lamandé, S. R. (2009) Genetic diseases of connective tissues: cellular and extracellular effects of ECM mutations. Nat. Rev. Genet. 10, 173–183

36. Fitzgerald, J., Lamandé, S. R., and Bateman, J. F. (1999) Proteasomal degradation of unassembled mutant type I collagen pro-α1(I) chains. J. Biol. Chem. 274, 27392–27398

37. Lamandé, S. R., Chesser, S. D., Golub, S. B., Byers, P. H., Chan, D., Cole, W. G., Silence, D. O., and Bateman, J. F. (1995) Endoplasmic reticulum-mediated quality control of type I collagen production by cells from osteogenesis imperfecta patients with mutations in the pro-α1(I) chain carboxyl-terminal propeptide, which impair subunit assembly. J. Biol. Chem. 270, 8642–8649