

Table 2. Functional analysis of NMJs in *myd* mice

	MFPP amplitude (mV) ^a	Quantal content (<i>m</i>) ^b	<i>d</i> -Tubocurarine concentration (μg/ml) ^b
Control 1	0.52 ± 0.07 (<i>n</i> = 16)	49.1 ± 5.5 (<i>n</i> = 10)	0.38
Control 2	0.64 ± 0.10 (<i>n</i> = 18)	52.8 ± 3.8 (<i>n</i> = 14)	0.45
<i>myd</i> 1	0.32 ± 0.04 (<i>n</i> = 11)	65.6 ± 6.5 (<i>n</i> = 5)	0.30
<i>myd</i> 2	0.36 ± 0.06 (<i>n</i> = 7)	75.4 ± 38.7 (<i>n</i> = 5)	0.30

^aValues given are mean ± SEM and number of endplates (in parentheses).

^bThe amount of *d*-Tubocurarine that can inhibit the muscle contraction induced by the EPP.

columns). The utrophin signal was exclusively seen on NMJs, but weakly seen on muscle sarcoplasmic membrane in FCMD (Fig. 6C, right column), which is usually seen in regenerating muscle fibers (26). NMJs from fetal wild-type or *myd* showed similar staining patterns as expected (data not shown), suggesting that this staining pattern is indicative of immature muscle fibers. This observation is consistent with our hypothesis that NMJ formation in secondary α-DGopathies is developmentally arrested during myotube maturation in the fetal phase.

DISCUSSION

FCMD has long been classified as 'muscular dystrophy', although the clinical characteristics differ from those of DMD. Muscular dystrophy is defined generally by necrotic change and active regeneration of muscle fibers. However, FCMD muscle in infantile stage seems more likely to have additional features, implying a more complex pathogenesis for FCMD. Indeed, our microarray analysis showed that general expression profiling clusters FCMD and DMD distinctively. Expression of mature muscle components was surprisingly low in FCMD, indicating less active regeneration of muscle fibers. We also saw similar expression profiles among all FCMD patients in our samples, indicating that FCMD is a chronic rather than progressive disorder, at least at the infantile period.

We confirmed maturational delay and aberrant NMJs in FCMD skeletal muscle fibers by expression profiling, morphological and histochemical analysis, and electrophysiological examination. These findings are common to secondary α-DGopathies but are not seen in DMD. In this study, we demonstrate that the etiology of secondary α-DGopathy skeletal muscle abnormalities may stem from maturational arrest caused by aberrant NMJs in addition to dystrophy. MDC1A, clinical characteristics of which are similar to FCMD, is described with an initial phase of necrosis and regeneration in the early steps of the disease (6). Although aberrant NMJ is also seen in MDC1A (27), the muscle fragility due to defect in the component of basement membrane mainly affects the phenotype and causes necrosis and regeneration.

A considerable body of evidence indicates that muscle differentiation ceases at NMJ formation in FCMD. First, immature type 2C fibers are predominant in FCMD (Table 1). At the initiation of muscle differentiation, satellite

cells proliferate to become myoblasts, fuse to organize myotubes. These early myotubes contain type 2C fibers. Following NMJ formation, immature fibers are induced to differentiate further into mature muscle fibers such as type 1, 2A and 2B. Second, down-regulated expression of matured MYHs in FCMD also suggests arrest at this stage. Following completion of NMJ formation, embryonic, neonatal-type MYHs in immature myofibers are replaced by adult-type, slow MYHs, which are induced by extracellular matrix (ECM) components, growth factors or programmed cell differentiation (28).

Third, MRF4, which is postulated to be induced by AChR clustering in NMJ formation, is down-regulated in FCMD. Normally, during the early phase of muscle development, a series of myogenic regulatory genes such as *myf5*, *MyoD* and *myogenin* are sequentially expressed, followed by MRF4 up-regulation just after NMJ formation. AChRs bind to myotubes, associate with specific factors and trigger a signal to induce expression of myogenic factors including MRF4, which is suggested to play a crucial role in muscle terminal maturation and maintenance (20,21,29). Therefore, MRF4 is distinct from *MyoD*, *myogenin* and *myf5* in that it is expressed mainly in matured myotubes and myofibers. MRF4 has been reported to co-localize with AChR in NMJs and to function in terminal muscle maturation and fiber maintenance (30). It is reasonable to assume that MRF4 function is required at the time and place of NMJ development during skeletal muscle differentiation (23). These results prompt the hypothesis that the differentiation process of muscle fibers arrests at this point in FCMD and *myd*.

Fourth, the fact that high *CHRNA* and low *CHRNA* expression in FCMD muscle relative to that in age-matched normal control or DMD muscle is striking, although the age of the DMD patient was slightly higher than that of the FCMD patients. It clearly indicates that most of the AChR in FCMD muscle is fetal type. It also supports our hypothesis that skeletal muscle in secondary α-DGopathy is immature and that delay of differentiation might be involved with maturation defect of NMJs.

What causes aberrant NMJs in secondary α-DGopathies? Normally, NMJs are built by a dense tangle of sarcoplasmic membrane and neural endings through a layer of basement membrane. It is possible that connections between the neural terminal and glycosylated α-DG, made through a layer of basement membrane and mediated by molecules such as laminin or agrin, are important to normal NMJ formation and subsequent muscle differentiation (31,32). MDC1A, clinical characteristics of which are similar to FCMD (6), also shows aberrant NMJs with fewer synaptic folds (27). In contrast, Musk or rapsyn deficiency does not resemble severe muscular dystrophy in spite of the abnormal NMJ formation. The interaction of laminin and α-DG is thought to play an essential role in the transition of AChR microaggregates into macroaggregates at the developing NMJ, followed by the concentration of α-DG on NMJs (32,33). These facts lend support to our hypothesis that the laminin/α-DG interaction fulfills a pivotal function in normal NMJ formation.

Alternatively, it is also possible that attachment of neural terminals to NMJs is affected pre-synaptically in secondary α-DGopathies. Abnormalities in the pre-synaptic peripheral nerve would affect the neural endings of NMJs, leading to

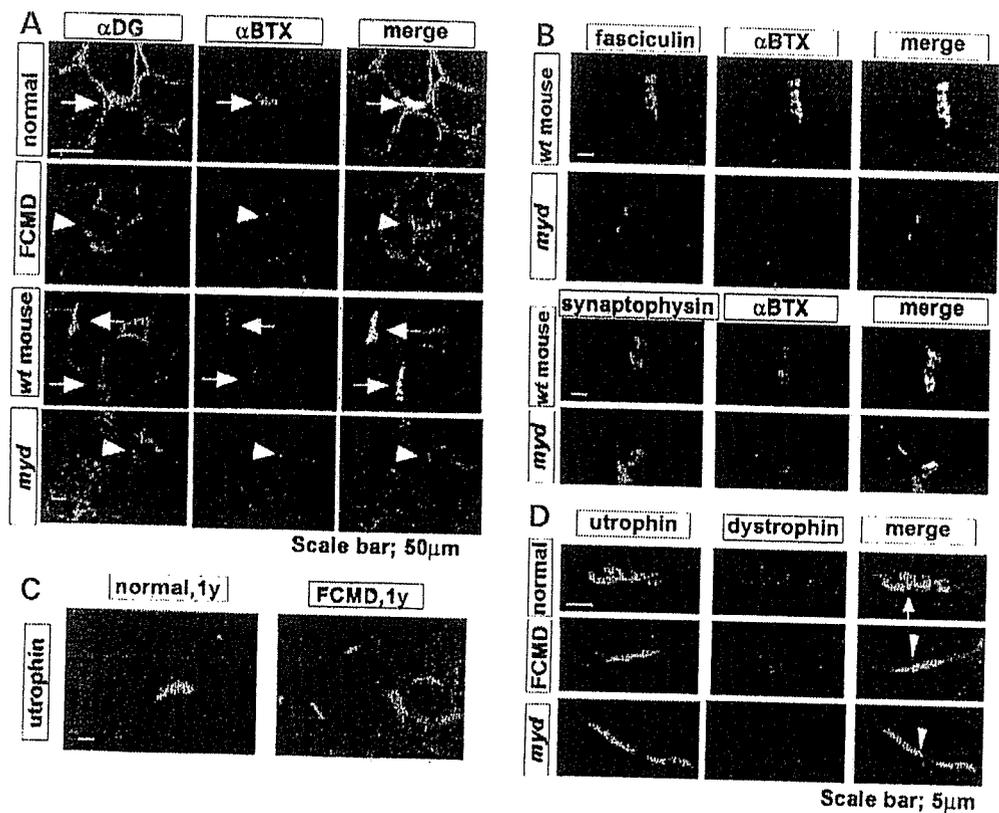


Figure 6. Aberrant NMJs and lack of α -dystroglycan aggregation in NMJs affect pre- and post-synaptic formation of NMJs in secondary α -DGpathy. (A) α -DG (green) strongly co-localizes with α -BTX in NMJs of normal skeletal muscle, merged yellow (arrows). However in secondary α -DGpathies, the lack of α -dystroglycan aggregation on aberrant formed NMJs is seen despite positive staining on muscle peripheral membrane, as evidenced by the lack of co-localization signal in merged column (arrowheads). (B) Green-stained pre-synaptic markers (fasciculin, AChE marker; synaptophysin, pre-synaptic membrane on secondary α -DGpathies. Although green-stained utrophin aggregates exclusively on NMJs of normal muscle fibers, utrophin remains on sarco-plasmic membrane of FCMD muscle fibers. (C) Confocal imaging of utrophin and dystrophin on NMJs of secondary α -DGpathy. Merged column in normal NMJs (arrow) shows that fine primary and secondary folds lined with utrophin are tangled with dystrophin staining on NMJs, which also line the peripheral membrane of muscle fibers. Note that in secondary α -DGpathies, secondary folds are severely lacking (arrowheads). Thin, scattered utrophin staining and markedly decreased staining of dystrophin on NMJs are seen in both FCMD and *myd*. Scale bars = 5 μ m.

deficient differentiation signal transduction to FCMD muscle fibers and arrested post-synaptic muscle differentiation. It has been suggested that α -DG in the central or peripheral nervous system is hypoglycosylated in secondary α -DGpathies (34). O-Mannose-type glycoprotein is suggested to contribute to the stability and maintenance of muscle cell membrane, synaptic formation and myelination of peripheral nerves, although the precise mechanism of α -DG activity in peripheral nerve tissue is unclear. The dystrophin-glycoprotein complex on Schwann cells is also thought to be important in peripheral myelinogenesis, regeneration, differentiation, apoptosis and polarity of skeletal muscle cells (35). Although Ishii *et al.* (36) reported that electron microscopy of Schwann cells on FCMD muscle revealed no pathologic findings, neural transmission may be developmentally impaired and collapsed as a result of hypoglycosylated α -DG.

Naturally, our data do not rule out the other possibilities for immaturity of the skeletal muscles of α -DGpathies. Although the expression profile and histochemical data indicate that skeletal muscles are in a persistent undifferentiated state, it is possible that the regeneration process fulfills a crucial function in the phenomenon observed in skeletal muscles of α -DGpathies. It is also possible that most of the muscle

fibers are under denervation status, because motor neurons are unable to maintain strong attachments to myofibers, leading to a constant stimulation of denervation signal pathways. Although we demonstrated substantial evidences for the aberrant NMJ, the pathogenesis of skeletal muscles in α -DGpathies is likely to be a combination of these problems and of multifactorial origin.

Taken together, these findings show that muscle fibers in secondary α -DGpathies are developmentally arrested more to 'dystrophic', perhaps because hypoglycosylated α -DG precludes proper aggregation on NMJs, preventing AChR clustering. These defects may disrupt terminal muscle maturation, which is induced after innervation of neurons on the muscle peripheral membrane via a basement membrane layer. Dystrophic changes traditionally thought to underlie 'muscular dystrophy' are caused by attenuated physical connections between α -DG and the muscle basement membrane. We propose that the muscle lesion in secondary α -DGpathies is caused by complex pathogenesis, not only by dystrophic change but more importantly, maturational arrest resulting from chronically delayed terminal muscle fiber maturation and NMJ deficiency. To date, no clinical approaches to secondary α -DGpathies exist. These findings open a possible

avenue for treating muscle tissues in these congenital muscular dystrophies, via targeted induction of NMJ maturation in muscle fibers.

MATERIALS AND METHODS

Samples

All clinical materials were collected for diagnostic purposes. Four muscle specimens (biceps branchii) from FCMD patients (ages: 20 days, 7 months, 1-year 4 months, and 1-year 6 months) were used in the analysis. Genetic screening identified a homozygous retrotransposal insertion into the 3' untranslated region of *fukutin* in all FCMD patients (3). For the non-dystrophic muscle controls, muscle RNAs from two children (ages: 1 year) was used. These patients were selected based on normal laboratory findings, normal plasma CK levels and no histopathological evidence for muscular dystrophy.

We also obtained myodystrophy (*Large^{myd}*) mice and control littermates (*Large^{myd/+}* or *Large^{+/+}*), aged 3 and 6 months, by mating heterozygous pairs provided by Jackson Laboratories.

RNA isolation and expression profiling

Generation of cDNA microarrays containing skeletal muscle transcripts has been reported previously (19). Using similar methods, we constructed a new cDNA chip containing 5600 genes expressed in skeletal muscle. RNA isolation, hybridization and detection methods also have been reported previously. Microarray experiments were carried out using a competitive hybridization method with two labeled targets: one for muscle RNAs from FCMD patients or normal children, and another for pooled muscle RNAs (Origene), which served as a template control for per-chip normalization. Each analysis was conducted at least twice. The hybridization intensities of each spot and the background intensities were calculated using a ScanArray 5000 microarray scanner with Quant Array software (Perkin-Elmer Life Science).

Microarray data analysis

Analysis of microarray data was performed using Genespring version 6.1 (Silicon Genetics) software. Data used for further analysis were calculated using a previously reported method (19). To avoid 'false-positive' signals, we excluded genes from the analysis for which average normal expression level constraints are under 500. We sorted 1790 genes from a total of 5600 for further analysis.

Quantitative real-time PCR

Two patients with FCMD (ages: 10 months and 1 year), two patients with DMD (ages: 1 and 7 years) and two normal children (ages: 1 and 2 years) were used for quantitative RT-PCR using skeletal muscle RNAs. Single-strand cDNA was produced with random primers, and quantitative real-time RT-PCR using SYBR-green was performed using the ABI Prism 7900 sequence detection system (Applied Biosynthesis). Data analysis was performed in duplicate experiments.

Statistical significance was evaluated using Student's *t*-test, and $P < 0.05$ was considered significant. The primers used for the experiments are shown in Supplementary Material, Table S1. *Gapdh* was used as an internal control.

Imaging analysis

HE staining and ATPase staining were performed on cryosections. ATPase staining was performed at pH 9.4–10.6 and 4.2–4.6. For ATPase staining, average data from 10 DMD patients (ages: 3–9) and 10 normal control cases (ages: around 1 year) were selected. Scion Image Beta 3b (Scion Corporation) was used for estimating the content of type 2C fibers, muscle fibers, adipose tissues and interstitial tissues (Table 1). Statistical analysis was performed using Student's *t*-test.

Immunohistochemistry

All muscle specimens were processed for cryosectioning (8- μ m thick) and fixed in 50% ethanol and 50% acetic acid for 1 min. Anti-core α -dystroglycan [GT20ADG, gift from Dr Kevin Campbell (5)], rhodamine-conjugated α -bungarotoxin (BTX, Molecular Probes), anti-synaptophysin (NCL-SYNAp, Novocastra), Alexa-fluor488 labeled anti-fasciculin2 (F4293, Sigma), anti-MRF4 (C-19, Cruz Biotechnology), anti-utrophin (UT2, gift from Dr Michihiro Imamura), anti-human dystrophin (NCL-DYS2, Novocastra), anti-human dystrophin (MANDRA-1, Sigma) and anti-myosin heavy chain for developmental, neonatal, fast and slow types (NCL-MHCd, MHCn, MHCf and MHCs, respectively, Novocastra) were used to stain NMJs.

Electron microscopy

Muscle specimens were obtained from four patients diagnosed as FCMD (ages: 6 months to 4 years old) and from two normal children (ages: 1 year). Five NMJs were found in three FCMD patients' skeletal muscle and four normal NMJs were examined in two normal children. Intercostal muscles and soleus muscles were dissected from myodystrophy and control mice, and cut into 1-mm thick cubes. Four NMJs were examined in *myd* and normal controls, respectively. Samples were fixed in 2% glutaraldehyde and embedded in epoxy resin as described previously (36). Ultrathin (50–90-nm thick) sections were cut on an Ultracut S ultramicrotome (Reichert).

Electrophysiologic examination

Diaphragms with its motor nerve were dissected and used for the conventional intracellular microelectrode study (37). MEPPs, EPPs and resting membrane potentials (RMPs) were recorded. For EPP recording, the phrenic nerve was stimulated using a suction electrode at 0.5 Hz. *d*-Tubocurarine chloride (Curaren, Sigma) was used at a concentration sufficient to inhibit muscle contraction. The potentials were corrected for non-linear summation and the last 64 responses in a train of 114 were saved for later analysis. The quantal content *m* was calculated by the variance method. MEPP and EPP amplitudes were corrected to a standard RMP of -80 mV. To correct MEPP amplitude by the fiber diameter, the geometric

mean of the shortest and longest diameters of muscle fibers was determined in 30 randomly selected muscle fibers in cryostat sections.

SUPPLEMENTARY MATERIAL

Supplementary Material is available at HMG Online.

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Conflict of Interest statement. None declared.

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Central core disease is due to *RYR1* mutations in more than 90% of patients

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Ryanodine receptor 1 (*RYR1*) gene mutations are associated with central core disease (CCD), multiminicore disease (MmD) and malignant hyperthermia (MH), and have been reported to be responsible for 47–67% of patients with CCD and rare cases with MmD. However, to date, the true frequency and distribution of the mutations along the *RYR1* gene have not been determined yet, since mutation screening has been limited to three ‘hot spots’, with particular attention to the C-terminal region. In this study, 27 unrelated Japanese CCD patients were included. Clinical histories and muscle biopsies were carefully reviewed. We sequenced all the 106 exons encoding *RYR1* with their flanking exon–intron boundaries, and identified 20 novel and 3 previously reported heterozygous missense mutations in 25 of the 27 CCD patients (93%), which is a much higher mutation detection rate than that perceived previously. Among them, six were located outside the known ‘hot spots’. Sixteen of 27 (59%) CCD patients had mutations in the C-terminal ‘hot spot’. Three CCD patients had a probable autosomal recessive disease with two heterozygous mutations. Patients with C-terminal mutations had earlier onset and rather consistent muscle pathology characterized by the presence of distinct cores in almost all type 1 fibres, interstitial fibrosis and type 2 fibre deficiency. In contrast, patients with mutations outside the C-terminal region had milder clinical phenotype and harbour more atypical cores in their muscle fibres. We also sequenced two genes encoding *RYR1*-associated proteins as candidate causative genes for CCD: the 12 kD FK506-binding protein (*FKBP12*) and the $\alpha 1$ subunit of L-type voltage-dependent calcium channel or dihydropyridine receptor (*CACNA1S*). However, no mutation was found, suggesting that these genes may not, or only rarely, be responsible for CCD. Our results indicate that CCD may be caused by *RYR1* mutations in the majority of patients.

Keywords: central core disease; genotype–phenotype correlation; muscular dystrophy; myopathy; ryanodine receptor 1 mutations

Abbreviations: *CACNA1S* = $\alpha 1$ subunit of L-type voltage-dependent calcium channel; CCD = central core disease; CICR = calcium-induced calcium release; EC = excitation–contraction; *FKBP12* = FK506-binding protein, 12 kD; mGT = modified Gomori–Trichrome; MmD = multiminicore disease; MH = malignant hyperthermia; *RYR1* = ryanodine receptor 1; SERCA = sarco-endoplasmic reticulum Ca^{2+} ATPase; SR = sarcoplasmic reticulum

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Introduction

Central core disease (CCD) was the first described congenital myopathy in humans (Shy and Magee, 1956), usually inherited in an autosomal dominant pattern, except for few reports on autosomal recessive cases (Manzur *et al.*, 1998;

Jungbluth *et al.*, 2002). The clinical features are quite variable (Quinlivan *et al.*, 2003), ranging from lack of visible weakness or abnormality to lack of independent ambulation. Most of the patients, in classical descriptions, have a slowly or

non-progressive proximal muscle weakness and hypotonia during infancy that can persist throughout adolescence/adulthood, and have associated delayed motor development and reduced muscle bulk. In addition, musculoskeletal alterations including congenital hip dislocation, kyphoscoliosis and joint contractures are also common findings in CCD patients (Lorenzon *et al.*, 2000; Quinlivan *et al.*, 2003). The diagnosis is commonly made from muscle biopsy by the presence of cores in type 1 fibres, which are typically well demarcated and located centrally in the fibres. A few cores, however, can be located in the subsarcolemmal regions of individual type 1 muscle fibres. Longitudinal sections show that the core runs the whole length of the fibre.

Recent reports have documented that the phenotypic presentation varies considerably from no visible disability to lack of independent ambulation. While the overall incidence of CCD is rare, the absence of symptoms in a significant number of patients may suggest that the actual incidence of CCD may be considerably higher than that perceived currently.

CCD has been linked to the gene encoding the skeletal muscle ryanodine receptor, *RYR1*, and is considered to be an allelic disease of malignant hyperthermia (MH) susceptibility. This is a pharmacogenetic disorder with autosomal dominant inheritance in which susceptible individuals develop generalized muscle contracture followed by a hypermetabolic state due to massive calcium release from the sarcoplasmic reticulum (SR), when they are exposed to inhaled general anaesthetics or to the depolarizing muscle relaxant succinylcholine. CCD patients have higher probability to be susceptible to MH. The *RYR1* mutations linked to MH and CCD are clustered in three relatively restricted regions of the protein or 'hot spots': N-terminal (residues p.M1-R614), central (p.R2163–p.R2458) and C-terminal (p.R4136–p.P4973). They are also called domains 1–3, respectively (Treves *et al.*, 2005). The first two domains are located in the soluble cytoplasmic regions of the protein. Most of the CCD mutations are clustered in domain 3, which is located in the C-terminus and comprises the transmembrane/luminal and pore-forming region of the channel (McCarthy *et al.*, 2000; Monnier *et al.*, 2001; Davis *et al.*, 2003). Hence, mutations in this region of the protein may directly alter the permeation/selectivity/gating properties of the channel (Balshaw *et al.*, 1999).

The *RYR1* is one of the largest described genes in humans, spanning >159 kb in size on chromosome 19.q13.1. A 15 117-nucleotide-long open reading frame encoded in 106 exons (two of which are alternatively spliced) produces a 563 kD protein. In addition, it forms a homotetrameric structure that functions as an SR calcium-release channel regulating Ca²⁺ content in skeletal muscle during excitation–contraction (EC) coupling. Because of the size of the *RYR1* gene, efficient routine screening for mutations has been difficult. Most of the *RYR1* mutation screenings in CCD patients have been limited to the above-mentioned three 'hot spots' or even to the C-terminal region alone, and 47–67% patients were found carrying *RYR1* mutations (Monnier *et al.*, 2001; Davis *et al.*,

2003; Shepherd *et al.*, 2004), which suggested that CCD is a genetically heterogeneous disease. Here, we screened all the exons and flanking exon–intron boundaries of *RYR1* in order to determine the frequency and distribution of mutations, and describe the genotype–phenotype correlation in Japanese patients with CCD.

Although no mutation in other genes has been associated with CCD, many studies have shown that mutants of the *RYR1*-associated proteins FKBP12 and CACNA1S cause EC uncoupling *in vitro* just as some *RYR1* mutants do (Avila *et al.*, 2003b; Lyfenko *et al.*, 2004; Weiss *et al.*, 2004), raising a possibility that FKBP12 and/or CACNA1S mutations may also be responsible for CCD. In addition, the mutations in the *RYR1*-binding domain of CACNA1S are thought to account for 1% of MH patients (Stewart *et al.*, 2001). We therefore sequenced the entire open reading frame of FKBP12 and the part of CACNA1S that encodes the *RYR1*-binding region.

Methods

Subjects

Unrelated Japanese CCD patients were selected for the study from the National Center of Neurology and Psychiatry (NCNP) database from 1982 to 2004. CCD diagnosis was established on the basis of characteristic muscle pathology findings of cores almost exclusively in type 1 fibres. We excluded multiminicore disease (MmD) cases, which had multiple cores in >70% of type 1 fibres in our series. Available blood samples from the patients' relatives were also included in the analysis. In addition, DNA samples from 150 subjects without any known muscle disease were studied. Informed consent was obtained from the patients or their parents, as well as from control subjects.

The patients' clinical features were assessed by careful review of their medical records. Pathological features of all patients were independently evaluated by four authors (S.W., M.C.V.M., I.N. and I.N.). All patients have undergone a battery of histochemical stains, including haematoxylin and eosin, modified Gomori–trichrome (mGT), NADH-tetrazolium reductase (NADH-TR) and myosin ATPase.

In order to evaluate the genotype–phenotype correlation in CCD patients, the patients were divided into four groups, respectively, provided that they had one heterozygous C-terminal mutation, one heterozygous non-C-terminal mutation, two heterozygous mutations or no mutation.

Data were entered in Statistics Software for Social Sciences (SPSS version 11.0). Demographic characteristics were analysed by computation of the frequency, the mean \pm standard deviation (SD), or the mean \pm standard error of means (SEM), whichever was appropriate. The data then were subjected to a univariate analysis (Fisher's exact test). For comparing age and type 2 fibre deficiency, Mann–Whitney test was employed.

Mutational analysis

Genomic DNA was extracted from either muscle biopsy samples or peripheral blood lymphocytes according to standard protocols (Sambrook *et al.*, 2001). PCR primers were designed to amplify all 106 exons of *RYR1*, all five exons of FKBP12 and the seven

exons of *CACNA1S* (exon 14–17 and 25–27) that encode the RYR1-interacting region (Peng *et al.*, 1998). Amplified fragments were directly sequenced using BigDye Terminator[®] v3.1 Cycle Sequencing kits on ABI3100 automated Genetic Analyzer (Applied Biosystems[™], USA). DNA sequences were analysed with the SeqScape program and compared with the reference genomic sequence of: *RYR1* (Genbank J05200), *FKBP12* (Genbank M92423) and *CACNA1S* (Genbank L33798).

Results

Patients

A total of 27 unrelated Japanese CCD patients were selected for the study, consisting of 7 males and 20 females (Table 1) ranging in age from 2 to 63 years [29 ± 19 years (mean \pm SD)] at the time of muscle biopsy. DNA from the parents of two patients (Patients 23 and 25) were also studied.

Patients 1–8 underwent muscle biopsy because they had pertinent family history of MH and were considered to be MH susceptible. In Japan, the diagnosis of MH susceptibility is achieved by detecting enhancement in the rate of calcium-induced calcium release (CICR) from the sarcoplasmic reticulum in chemically skinned muscle fibres (Ibarra *et al.*, 2005), reflecting the underlying pathomechanism of this disorder, that is, a lower activation threshold of the SR calcium-release channel. Cores were incidentally found in the muscle biopsies of Patients 1–8; thus a working diagnosis of CCD was made despite the absence of any muscle symptoms in most of these patients. CICR rate was enhanced in all eight patients. There was no description about the MH susceptibility status on the medical records of other subjects (Patients 9–27).

Nemaline bodies were identified in Patients 11, 12 and 23 on mGT stain; hence they were labelled as having core/rod disease (CRD).

Mutations

A total of 23 different missense mutations affecting 19 residues in *RYR1* were identified, consisting of 3 previously reported and 20 newly identified missense mutations in 93% of CCD patients (Tables 1 and 2). Two patients did not have any mutation in *RYR1*. All mutations were heterozygous single nucleotide changes, except for the substitution of two consecutive nucleotides (c.14761_14762TT>AC) in Patient 21, which was predicted to result in a single amino acid change (p.F4921T). There were two common heterozygous *RYR1* mutations in this cohort: c.14581C>T (p.R4861C) and c.7522C>G (p.R2508C), identified in 4 out of 25 (16%) and in 3 out of 25 (12%) patients, respectively. No mutations were found in either *FKBP12* or *CACNA1S*.

Twenty-two of 25 (88%) had only one heterozygous mutation: 14 (56%) with a heterozygous C-terminal mutation and 8 (32%) with a heterozygous non-C-terminal mutation. The remaining three (12%) patients had two heterozygous mutations (Patients 23, 24 and 25). Only

the parents of Patients 23 and 25 were screened for the respective pair of mutations, as samples were not available from the parents of Patient 24. In Patient 23, the mutation p.D60N was identified in exon 3 of her maternal allele while p.L3606P was found in exon 73 of her paternal allele. In Patient 25, p.E512K in exon 14 was inherited from her mother while p.R4893P in exon 102 was acquired from her father. No muscle samples were available for histological studies from the parents of these two patients. *RYR1* mutations were found in patients with CRD (Patients 11, 12 and 23) (Table 1, Fig. 1). None of the newly identified missense mutations were found in 300 control chromosomes. Eighteen of 19 amino acids predicted to be changed were highly conserved through *RYR1* evolution and most of them were also conserved across the *RYR* species, *RYR1*, *RYR2* and *RYR3* (Table 2).

The mutations found in this cohort of CCD patients are shown in Fig. 2 at their respective location in the *RYR1*, along with those reported previously. In this study, 6 mutations were located outside the mutational 'hot spots', 4 in N-terminal 'hot spot' and 13 in the C-terminal region. These mutations were carried by 8 out of 25 (32%), 4 out of 25 (16%) and 16 out of 25 (59%) patients, respectively. We did not find any mutation in the central 'hot spot' or domain 2 in our cohort.

Genotype–phenotype correlation

Among eight patients with a heterozygous non-C-terminal mutation, two showed mild limb muscle weakness while the others were asymptomatic. The exact age of onset was difficult to ascertain because of the paucity of symptoms. Significant clinical findings during pregnancy and birth were not reported in this group. On muscle histochemistry, type 2 fibre deficiency was seen in only one patient, as he had <1% of type 2 fibres, while these were >12% in others [$37 \pm 4\%$ (mean \pm SD)]. Minimal endomysial fibrosis was found in three patients (37%), while the others had no increase in the interstitial fibrous tissue.

In contrast, limb muscle weakness was present in all patients with C-terminal mutations, and significant symptoms were manifested during the perinatal period (Table 1). There was statistically significant association between the presence of mutation in the C-terminal domain and clinical phenotypic characteristic in the following categories: limb muscle weakness, poor foetal movement during pregnancy, presence of joint dislocation and delayed motor milestone (Fig. 3A). Endomysial fibrosis and type 2 fibre deficiency were observed in most, if not all, patients with C-terminal mutations (Fig. 3B).

In terms of core structure, most patients had single cores (95% of fibres), but multiple cores were also seen especially in patients with heterozygous non-C-terminal mutations. In mutations outside domain 3, most cores were located in the periphery or subsarcolemmal areas (71%). In patients with C-terminal mutations, cores were noted to be

Table 1 Summary of clinical and pathological features of the patients and RYR1 mutations

Patient	Sex	Pregnancy and birth symptoms	Family history	Onset Clinical symptoms			Age at biopsy (years)	CICR					Nemaline bodies	Endomyssial RYR1 mutations (Exon)				
				Delayed motor milestone	Muscle weakness	Skeletal and/or articular malformation		Others	Type 2 fibres [†]	Type 1 fibres with cores [†]	Features of cores [‡]	Central cores			Rimmed cores	Atypical cores	Single core	Multiple core cores
1	F	N	Yes	Ad	No	JC	25	E	32	93	15	39	20	83	17	No	Minimal	c.1422G>T.
2	F	N	Yes	Ad	No	Sc	37	E	27	93	22	9	12	96	4	No	No	p.Q474H (13)* c.7522C>T.
3	F	N	No	Ad	No		63	E	13	89	22	17	4	92	8	No	No	p.R2508C (47)* c.7522C>T.
4	F	N	No	Ad	No	Sc	35	E	13	84	14	11	14	84	16	No	No	p.R2508C (47)* c.7522C>T.
5	F	N	Yes	Ch	No	Sc	9	E	40	98	74	12	5	96	4	No	No	p.R2508C (47)* c.7522C>G.
6	M	N	Yes	Ad	No		34	E	65	78	14	39	10	86	14	No	No	p.R2508G (47)* c.7523G>A.
7	F	N	No	Ch	No		8	E	<1	39	28	9	12	89	11	No	Minimal	p.R2508H (47)* c.7635G>C.
8	M	N	Yes	Ad	No		48	E	75	24	43	8	27	99	1	No	Minimal	p.E2545D (48)* c.10100A>G.
9	F	N	No	Ch	Yes		4	ND	<1	99	34	75	1	>99	1	No	Moderate	p.K3367R (67)* c.13703T>C.
10	F	PFM, FI, RI, PS	No	Bi	Yes	JD, JC	3	ND	1	99	78	62	1	100	0	No	Marked	p.L4568P (94)* c.13891T>A.
11	M	FI	No	In	Yes	JD, JC	3	ND	2	68	41	71	28	100	0	Yes	Moderate	p.Y4631N (95)* c.13900G>A.
12	F	PFM, FI	Yes	Bi	Yes	JD, JC	2	ND	1	98	12	88	1	100	0	Yes	Minimal	p.E4634K (95)* c.13912G>A.
13	F	PFM	No	Ch	Yes	JC, Sc	3	ND	<1	86	80	70	13	98	2	No	Marked	p.G4638S (95)* c.14572A>G.
14	F	FI	No	In	No	JD, JC, Sc	10	ND	1	99	81	74	15	99	1	No	Marked	p.N4858D (101)* c.14581C>T.
15	M	N	No	Ch	Yes		10	ND	<1	99	86	80	1	>98	2	No	Minimal	p.R4861C (101) c.14581C>T. p.R4861C (101) p.R4861C (101)

16	F	RI, PS	No	Bi	Yes	P	JD, JC, Sc, Lo	2	ND	3	60	69	78	3	98	2	No	Marked	c.14581C>T, p.R4861C (101)
17	F	PFM	No	Ch	Yes	G	JD, Sc	2	ND	<1	98	58	76	5	99	1	No	Marked	c.14581C>T, p.R4861C (101)
18	M	N	No	Ch	Yes	P	JC, Lo	3	ND	<1	99	67	80	2	99	1	No	Minimal	p.R4861C (101) c.14582G>A, p.R4861H (101)
19	F	PFM, FI, RI	No	Bi	Yes	G	JD, Sc	2	ND	<1	83	21	55	36	100	0	No	Minimal	p.R4861H (101) c.14693T>C, p.14898T (102)*
20	F	PFM, FI	No	Ch	Yes	P		9	ND	<1	96	97	72	4	99	1	No	Minimal	c.14759C>A, p.T4920N (102)*
21	M	N	Yes	Ch	No	P	Lo	6	ND	<1	>99	74	86	6	99	1	No	Minimal	c.14761T 14762TT>AC, p.F4921T (102)*
22	M	PFM, FI, RI	No	Bi	Yes	P	JD, Sc	3	ND	<1	99	34	82	23	82	18	No	Marked	c.14762T>C, p.F4921S (102)*
23	F	FI	Yes	In	Yes	P	Lo	5	ND	9	66	12	12	30	98	2	Yes	Moderate	c.178G>A, p.D60N (3); c.1081T>C, p.L3606P (73)* c.1280C>T, p.S427L (13); c.14696G>A, p.G4899E (102)*
24	F	FI, RI, PS	No	Bi	Yes	G	JD, Lo	10	ND	<1	14	58	62	9	99	1	No	No	c.1534G>A, p.E512K (14); c.14678G>C, p.R4893P (102)*
25	F	FI, RI, PS	No	Bi	Yes	P	JD, Lo	2	ND	21	>99	89	66	29	98	2	No	Moderate	Not detected Not detected
26	F	N	Yes	Ch	Yes	P	Sc	36	ND	13	99	94	0	74 [†]	98	2	No	Minimal	
27	F	N	No	Ad	No	D		35	ND	72	52	60	18	31	100	0	No	None	

C-terminal mutations are written in bold letters. F = female; M = male; N = normal; Ad = adolescence; Ch = childhood; Bi = birth; In = infancy; PFM = poor foetal movement; FI = floppy infant; RI = respiratory insufficiency; PS = poor sucking; JC = joint dislocation; JD = joint dislocation; Sc = scoliosis; Lo = lordosis; FMI = facial muscle involvement; HAP = high-arched palate; D = distal; P = proximal; G = generalized; CICR = calcium-induced calcium release; E = enhanced; ND = not done. * Novel mutations; † Values expressed as percentage of fibres; ‡ Refer to text and/or figure for description of cores.

Table 2 Comparison of amino acid in the affected residues with that from mouse, pig, rabbit and other human RYR proteins, RYR2 and RYR3

Residue	60	427	474	512	2508	2545	3367	3606	4568	4631	4634	4638	4858	4861	4893	4898	4899	4920	4921
Predicted change	N*	L*	H*	K*	G/H/C*	D*	R*	P*	P*	N*	K*	S*	D*	H/C	P*	T	E*	N*	S/T*
RyR1																			
Human	D	S	Q	E	R	E	K	L	L	Y	E	G	N	R	R	I	G	T	F
Mouse	D	S	Q	E	R	E	K	L	L	Y	E	G	N	R	R	I	G	T	F
Pig	D	S	Q	E	R	E	K	L	L	Y	E	G	N	R	R	I	G	T	F
Rabbit	D	S	Q	E	R	E	K	-	L	Y	E	G	N	R	R	I	G	T	F
RyR2																			
Human	D	K	Q	E	R	D	K	L	L	Y	E	G	N	R	R	I	G	T	F
RyR3																			
Human	D	-	Q	M	R	E	K	V	L	Y	Q	G	N	R	R	I	G	T	F

*Novel mutations.

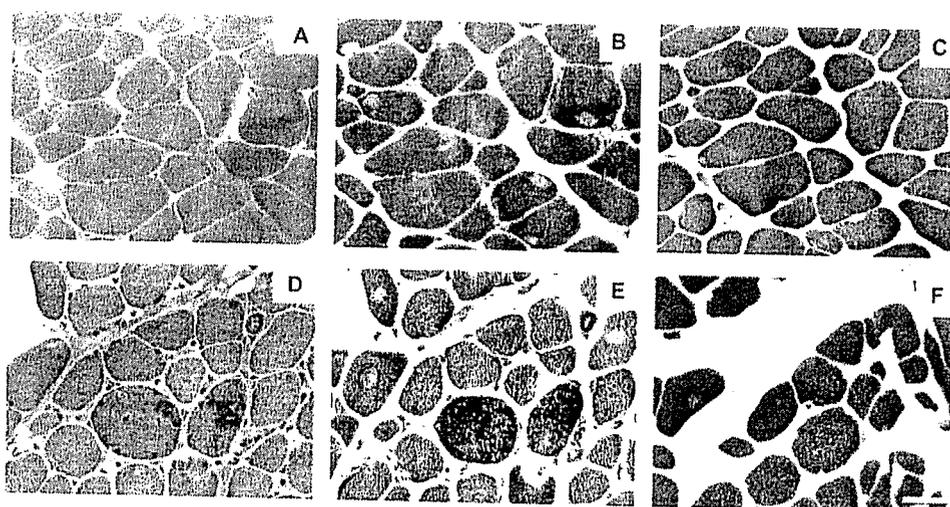


Fig. 1 Muscle biopsy. (A–C) A 10-year-old patient with a double mutation. (A) Minimal fibrosis is observed in mGT section. (B) NADH shows cores in type I fibres. (C) ATPase staining at pH 4.6 shows type 2 fibre deficiency. (D–F) A 3-year-old with nemaline bodies in muscle fibres. (D) Nemaline bodies and moderate fibrosis are depicted in mGT section. (E) Fibres with nemaline bodies have high enzymatic activity; typical central cores are noted. (F) ATPase staining with pH 4.3 pre-incubation demonstrates type 2 fibre deficiency. Bar denotes 50 microns.

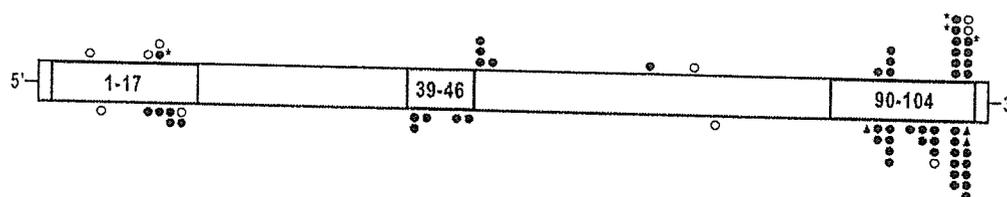


Fig. 2 *RYR1* mutation map for CCD. The three mutational hot spot areas are shaded: CCD domain 1: 1–17; CCD domain 2: 39–46; CCD domain 3: 90–104. Missense mutations (closed circle) found in this study are shown at the top; previously reported mutations are indicated at the bottom. Open circle: recessive mutation; triangle: deletions; asterisk: mutations identified in this study but were also included in previous reports.

characteristic: these were ovoid in shape and with clearly demarcated borders, and were predominantly located in the centre of the fibres [$60 \pm 7\%$ (mean \pm SEM)]; they also occurred singly ($98 \pm 4\%$). In addition, most of the cores ($72 \pm 5\%$) in this group appeared to be ‘rimmed’ (Fig. 4B), which may connote high enzymatic activities around the cores on NADH-TR stain. More cores in type I fibres were also noted ($90 \pm 4\%$), but in general the percentage of the fibres with cores varied from 3 to 100%.

No correlation was seen between clinical severity and the percentage of cores in fibres.

Some fibres show ‘atypical’ cores, characterized by indistinct borders and whose shapes were inexplicitly ovoid (Fig. 4E). These were seen more in non-C-terminal mutations but no statistical significance was observed as compared with C-terminal mutations (Fig. 3B). Like in typical core structures, these were also noted to be either in the centre of the fibre or in the subsarcolemmal areas.

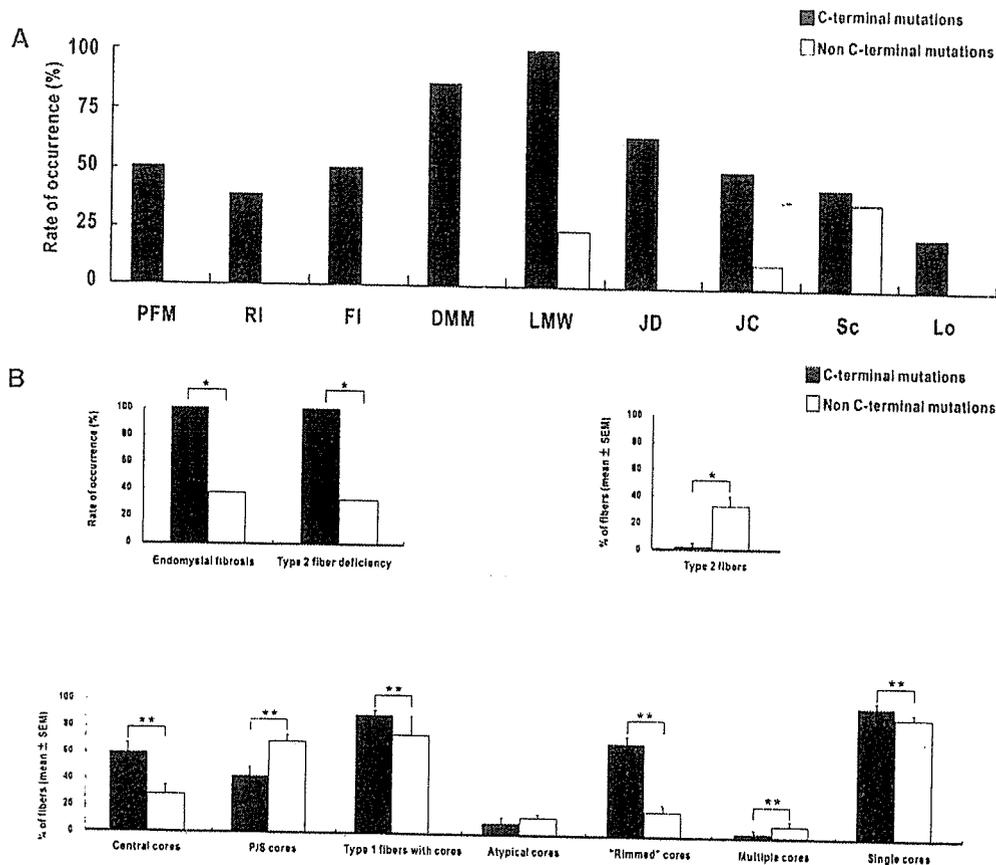


Fig. 3 (A) Genotype–phenotype correlation (clinical). Comparison of the clinical features between the patients with a heterozygous C-terminal mutation and those with a heterozygous non-C-terminal mutation. PFM = poor foetal movement; RI = respiratory insufficiency; FI = floppy infant; DMM = delayed motor milestone; JD = joint dislocation; JC = joint contracture; S = scoliosis; L = lordosis; LMW = limb muscle weakness. * $P < 0.05$, Fisher's exact test. (B) Genotype–phenotype correlation (pathological). Comparison of the pathological features between the patients with a heterozygous C-terminal mutation and those with a heterozygous non-C-terminal mutation. Note the characteristic endomysial fibrosis and fibre type 2 deficiency in patients with C-terminal mutations (upper panel). P/S = peripheral or subsarcolemmal. * $P < 0.05$, Fisher's exact test. ** $P < 0.05$, Mann–Whitney test; SEM, standard error of means.

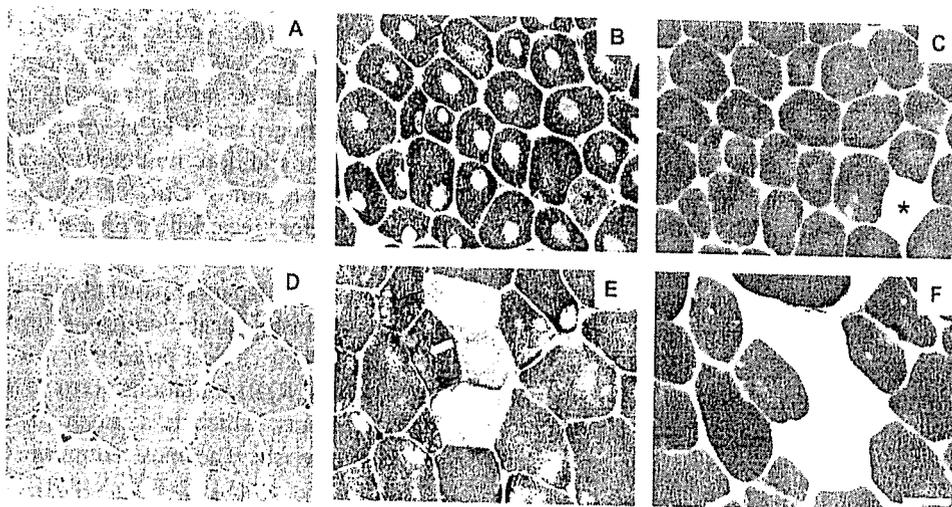


Fig. 4 Muscle biopsy. (A–C) A 9-year-old patient with a mutation in C-terminal region. (A) Minimal fibrosis is seen in mGT. (B) NADH-TR stain reveals the 'central core' in almost all fibres; note the 'rimming' of some cores (arrow). (C) Myosin ATPase staining, pH 4.4, clearly demonstrates type 2 fibre deficiency; type 2 fibre is marked with asterisk. (D–F) A 63-year-old patient with non-C-terminal mutation. (D) Minimal fibrosis is observed in mGT. (E) NADH shows cores but not on all type 1 fibres; note atypical cores (red arrow) and multiple cores in fibres (yellow arrow). (F) ATPase at pH 4.5 shows type 2 fibres in higher frequency compared with patient with C-terminal mutation. Bar denotes 50 microns.

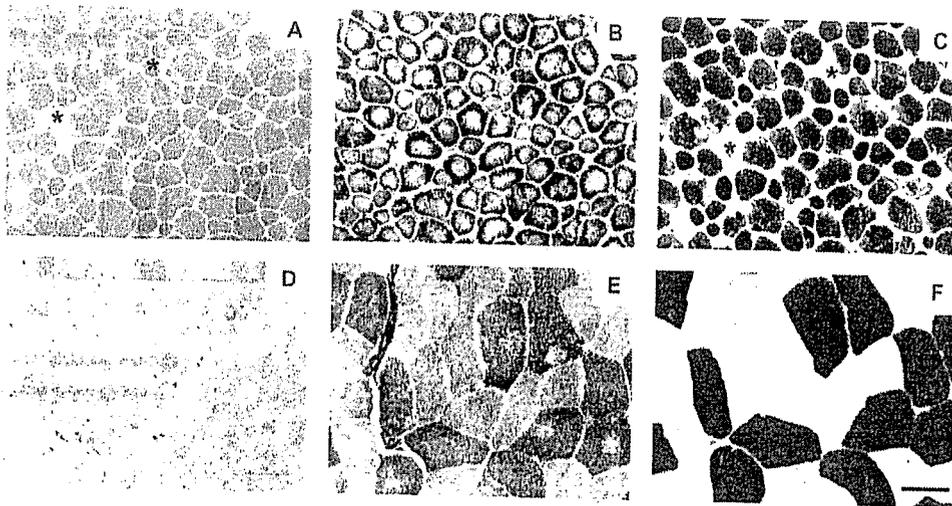


Fig. 5 Sections from patients in whom mutations were not found. (A–C) Sections from a 36-year-old female. (A) Moderate endomysial fibrosis is observed in mGT. (B) Cores are seen in almost all fibres; note that cores occupy the whole diameter of fibres, and the subsarcolemmal area surrounding the cores has high oxidative enzyme activity. (C) Type 2 fibre (asterisk) deficiency is seen in ATPase with pH 4.4 pre-incubation. (A–C) Biopsy findings from a 35-year-old female. (D) No endomysial fibrosis is noted in mGT. (E) Cores were seen but only in few type 1 fibres. (F) No fibre type 2 deficiency is observed (type 2 fibres comprised 72%). (C) Scale bar denotes 50 microns.

In Patients 1–8, from whom muscle biopsy was performed for CICR and cores were incidentally identified, all had non-C-terminal heterozygous mutations.

The clinical features of patients with two heterozygous mutations were more similar to those of the group with one heterozygous C-terminal mutation. In terms of muscle pathology, type 2 fibre deficiency was likewise seen (Table 2), similar to mutations involving the C-terminal region. This idea may also be buttressed by the fact that among a total of three patients with CRD, two carried a single C-terminal mutation, while the other had a compound heterozygous mutation. In terms of core pathology, however, they resemble that of non-C-terminal mutations: the cores were seen in 14 and 49% of type 1 fibres in Patients 24 and 25, respectively; and the characteristic ‘rimming’ of cores is less appreciated (Fig. 1).

In two patients in whom no mutation was found, the cores had peculiar features distinguishable from that seen in patients with identifiable mutations. Particularly, in Patient 26, the cores, albeit located centrally, occupied almost the whole diameter of the fibre, and, notably, the subsarcolemmal area around the cores had increased oxidative activity (Fig. 5). The cores in Patient 27 were initially considered to be regular cores but a closer look has shown that almost all the cores had indistinct borders; moreover, only <20% of type 1 fibres have cores (Fig. 5). In such cases, the term ‘core-like’ may be more appropriate to describe these structures.

Discussion

RYR1 mutations in CCD

At least 44 reported *RYR1* mutations have been associated with CCD, including 39 missense mutations and 5 deletions (Quane *et al.*, 1993; Zhang *et al.*, 1993; Fletcher *et al.*, 1995;

Manning *et al.*, 1998; Lynch *et al.*, 1999; Monnier *et al.*, 2000, 2001; Avila and Dirksen, 2001; Tilgen *et al.*, 2001; Ferreiro *et al.*, 2002; Jungbluth *et al.*, 2002; Robinson *et al.*, 2002; Sewry *et al.*, 2002; Avila *et al.*, 2003a; Quinlivan *et al.*, 2003; Zorzato *et al.*, 2003; Shepherd *et al.*, 2004). All these mutations are clustered in the three ‘hot spots’, except for two (Fig. 2). *RYR1* mutations have been reported to be responsible for 47–67% of patients suffering from CCD, implying that the disease is genetically heterogeneous (Vainzof *et al.*, 2000; Monnier *et al.*, 2001). However, neither the true frequency nor the distribution of CCD-causative mutations has been accurately determined to date, since mutation screening has been limited to the three ‘hot spots’ of the *RYR1*, or even only to the C-terminal region. According to *in vitro* studies, two *RYR1*-binding proteins, *FKBP12* and *CACNA1S*, directly participate in or modulate EC coupling, while EC uncoupling is thought to be the linchpin in the pathogenesis of CCD (Avila *et al.*, 2001; Lyfenko *et al.*, 2004). In addition, mutations have been identified in *RYR1*-binding region of *CACNA1S* in 1% of patients with MH. We therefore regarded *FKBP12* and *CACNA1S* genes as rational candidates for CCD.

We screened 27 Japanese patients with CCD, diagnosed on the basis of histological findings, without putting much emphasis on their clinical presentation as inclusion criteria, and found that *RYR1* mutations occur in 93% (25 out of 27) of CCD patients, which is a much higher rate than that thought previously. The most common CCD mutation in our cohort was c.14581C>T (p.R4861C). Interestingly, the c.14582G>A (p.R4861H) mutation that affects the same amino acid is the most common CCD mutation in European countries, while it was identified in only one patient from our series. Almost all reported patients bearing c.14582G>A had a positive family history, but all patients carrying mutation c.14581C>T were sporadic in this study and also in the

literature (Monnier *et al.*, 2001; Tilgen *et al.*, 2001; Davis *et al.*, 2003), suggesting that the 14 581 nucleotide might be susceptible to change.

To our knowledge, this is the first endeavour to screen the entire coding region of *RYR1* in a cohort of CCD patients. By screening the C-terminal region alone, we would have found *RYR1* mutations only in 59% (16 out of 27) of CCD patients, which is consistent with detection rates in previous reports, missing the two compound heterozygous mutations. Even extending the mutation screening to the three 'hot spots', we would have found *RYR1* mutations in only 67% (18 out of 27), also missing one compound heterozygous mutation. It thus becomes necessary to screen the entire *RYR1* coding region in CCD patients, however impractical it may be owing to the size of the gene. Interestingly, most of mutations outside the 'hot spots' were localized in exons 47 and 48, which neighbour the central region 'hot spot' (exons 39–46). Hence, mutation detection rate in CCD patients could be increased up to 89% by including exons 47 and 48, which may be a more practical alternative than sequencing the entire *RYR1* gene. In addition, *FKBP12* and *CACNA1S* may not be causative genes for CCD or an extremely rare, if they are, since there was no mutation in *FKBP12* or in the *RYR1*-binding region of *CACNA1S* genes.

We did not find any mutation in two patients in our cohort, suggesting that CCD may still be genetically heterogeneous even though there still remains the possibility that mutations may exist in unexamined regions, such as promoter region and introns, or that we may have overlooked a mutation. Notwithstanding this likelihood of other probabilities, the pathological characteristics of these patients are clearly different from the rest. If these patients were excluded from the analysis solely on the basis of the 'uncharacteristic' core-like structures, that is, if the inclusion criteria used were more stringent, the detection rate of mutations involving the *RYR1* gene will considerably and significantly increase to 100%.

Probable autosomal recessive CCD

CCD was once thought to be inherited solely via an autosomal dominant mechanism, but actually rare instances of recessive inheritance have also been identified (Manzur *et al.*, 1998; Ferreira *et al.*, 2002; Romero *et al.*, 2003).

In this study, sequencing of the entire *RYR1* coding regions led to the identification of three patients (Patients 23, 24 and 25) with two heterozygous mutations. Patients 24 and 25 needed respiratory mechanical assistance after delivery, while their parents were completely healthy and without any skeletal abnormalities. In both patients, each of the two heterozygous mutations was, respectively, found in each of the parents, confirming that these patients had a compound heterozygous mutation. In Patient 25, p.R4893P affected the third residue of a very well conserved GVRAGGGIGD luminal motif (amino acids p.G4891-D4900) that has

been proposed to be a pore-forming fragment responsible for the electrophysiological characteristics of the channel (Zhao *et al.*, 1999).

The parents of Patients 23 and 25 were asymptomatic, while they had heterozygous mutation, suggesting the recessive nature of these mutations. Since CCD patients could be clinically asymptomatic, however, we may not be able to completely exclude the possibility that their parents could have CCD. Nevertheless, judging from the clinical features and the mutation data, autosomal recessive mode of inheritance is most likely in these particular patients. Patient 24 also carried two heterozygous mutations (p.S427L in exon 13, p.G4899E in exon 102), but further analysis was inevitably limited by the unavailability of samples from her parents, hampering full evaluation if this was indeed an autosomal recessive case. The p.G4899E mutation also affected the very well conserved GVRAGGGIGD luminal motif (G4899 underlined). Interestingly, this mutation has been reported in two papers as a causative heterozygous mutation (Monnier *et al.*, 2001; Romero *et al.*, 2003); however, since only the 'hot spots' were screened in these studies, the presence of another mutation is still possible and hence it might be impetuous to conclude that p.G4899E mutation is a causative dominant mutation.

Genotype–phenotype correlation

In general, CCD has a wide spectrum of phenotypic expression, ranging from the apparent absence of symptoms to the presence of perinatal complications and generalized muscle weakness. Mutations in the C-terminal region seem to be associated with certain clinical and pathological features: hypotonia during infancy, delayed motor development and limb muscle weakness; type 2 fibre deficiency and interstitial fibrosis; and characteristic cores with clearly demarcated borders, which are observed in almost all type 1 muscle fibres. In addition, 'rimming' on the borders of these cores is observed in much higher frequency; similarly, this phenomenon was also noted in the biopsy specimens of three families determined to have mutation in domain 3 (Sewry *et al.*, 2002). These unique features therefore delineate C-terminal mutations from other groups, at least in terms of muscle pathology.

In contrast, most of CCD patients harbouring at least one mutation outside the C-terminal region had only mild musculoskeletal abnormalities such as joint contractures and scoliosis. This phenomenon may be explained by the leaky-channel model and EC uncoupling model. Some non-C-terminal mutations in *RYR1* promote the leak of Ca^{2+} ions from the SR that may or may not be compensated by the activity of the sarco-endoplasmic reticulum Ca^{2+} ATPase (SERCA), resulting in an elevation of resting cytosolic Ca^{2+} and a depletion of SR Ca^{2+} stores. On the other hand, C-terminal mutations, especially those in the pore region of *RYR1*, may directly affect the channel gating properties, resulting in an abolition of orthograde activation by the

voltage-gated L-type Ca^{2+} channel or, in other words, EC uncoupling. However, there is no compensatory mechanism to increase Ca^{2+} release as SERCA pumps do in the leaky model (Tong *et al.*, 1999; Avila *et al.*, 2001, 2003*b*; Lyfenko *et al.*, 2004). This may explain why C-terminal mutations are associated with clinically evident muscle weakness.

Interestingly, all patients with enhanced CICR had a non-C-terminal heterozygous mutation. Although we did not measure CICR in patients with C-terminal mutations, our results may suggest that majority of CCD patients with MH susceptibility might have a non-C-terminal mutation, considering the facts that no patient with C-terminal mutation had a family history of MH susceptibility, and that one C-terminal mutation, p.I4898T, has been associated with normal halothane and caffeine sensitivities by an *in vitro* study (Lynch *et al.*, 1999). In a recent study made by Monnier *et al.* (2005), where individuals from IVCT-confirmed MH-susceptible families have been screened for *RYR1* mutations and CCD, some MH-normal patients had cores in type 1 fibres albeit the absence of mutation. It would thus be interesting to screen the whole *RYR1* in these patients, as only selected exons were included for genetic screening. In our study, however, we definitely cannot assume that patients with C-terminal mutations may have normal CICR test, as further evaluations are necessary. Furthermore, in only such patients, multiple cores were seen in addition to central cores. Thus, non-C-terminal heterozygous mutations may be more associated with multiple cores and MH susceptibility, although the number of patients is too small to enable us to draw a conclusion. Previous reports have asserted that exclusive MH susceptible patients have mutations in the C-terminal region, but there is only limited study on the histopathological evaluation of these patients, revealing the presence of cores; it is also important to stress that these cores are rather not characteristic of CCD (Ibarra *et al.*, 2006).

We have shown that the muscle pathology on the patients may differ among those with C-terminal mutations and those with mutations outside this area. Cores, moreover, were not only located in the centre of the muscle fibres but also in subsarcolemmal or peripheral areas, and atypical cores can also be seen in such cases, indicating that the possibility of *RYR1* mutation cannot be excluded in patients with such atypical cores. Here, atypical cores of CCD were still different from the cores of MmD, for most of the cores were clearly demarcated and only some fibres appeared to have more than one core; nevertheless, histological, clinical, genetic features and mode of inheritance overlap between CCD and MmD, making the boundary of these two diseases blurred (Lyfenko *et al.*, 2004; Mathews *et al.*, 2004). Patients 24 and 25, who possessed double mutations, had pathology akin to characteristics of both C- and non-C-terminal mutations: type 2 fibre deficiency is seen, and cores are rather atypical and are only seen in moderate number on type 1 fibres. This may, in part, be explained by the fact that these patients carry mutations in and outside the C-terminal region.

In the past, we may have been biased in interpreting the phenotype–genotype correlation between CCD and *RYR1* mutations, since mutation screening had been limited to the ‘hot spots’. Quane *et al.* (1993) claimed that various phenotypes could be observed even among patients with the same mutation. However, their patients may have had a compound heterozygous mutation and one of the mutations may have been overlooked. In fact, if we had just screened the *RYR1* C-terminal region of patients with two heterozygous mutations (Patients 24 and 25), we would have found only one mutation, giving an impression of remarkable phenotypic difference between the patients and their healthy parents, regardless of the presence of the same ‘heterozygous’ mutation.

In summary, our comprehensive mutation screening revealed a surprisingly higher detection rate of *RYR1* mutations in CCD patients than that revealed in previous reports, which accounts for >90% of the cases. Our results also suggest the possibility that many *RYR1* mutations may have been overlooked by the regularly used screening methods that cover only three ‘hot spots’. At least, exons 47 and 48 should be included into the hot spot screening if a comprehensive mutation analysis of *RYR1* is not feasible. In terms of muscle pathology, patients with mutations involving the C-terminal region have characteristic cores that can be easily distinguished from those with mutations outside this domain, and also from those patients without mutation in the *RYR1*. Needless to say, the pathomechanism of how these cores develop still remains to be elucidated.

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A *Gne* knockout mouse expressing human V572L mutation develops features similar to distal myopathy with rimmed vacuoles or hereditary inclusion body myopathy

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Distal myopathy with rimmed vacuoles (DMRV) or hereditary inclusion myopathy (h-IBM) is an early adult-onset distal myopathy caused by mutations in the UDP-*N*-acetylglucosamine 2-epimerase/*N*-acetylmannosamine kinase (*GNE*) gene which encodes for a bifunctional enzyme involved in sialic acid biosynthesis. It is pathologically characterized by the presence of rimmed vacuoles especially in atrophic fibers, which also occasionally contain congophilic materials that are immunoreactive to β -amyloid, lysosomal proteins, ubiquitin and tau proteins. To elucidate the pathomechanism of this myopathy and to explore the treatment options, we generated a mouse model of DMRV/h-IBM. We knocked out the *Gne* gene in the mouse, but this resulted in embryonic lethality. We therefore generated a transgenic mouse that expressed the human *GNE*V572L mutation, which is the most prevalent among Japanese DMRV patients, and crossed this with *Gne*^(+/-) mouse to obtain *Gne*^(-/-)h*GNE*V572L-Tg. Interestingly, these mice exhibit marked hyposialylation in serum, muscle and other organs. Reduction in motor performance in these mice can only be seen from 30 weeks of age. A compelling finding is the development of β -amyloid deposition in myofibers by 32 weeks, which clearly precedes rimmed vacuole formation at 42 weeks. These results show that the *Gne*^(-/-)h*GNE*V572L-Tg mouse mimics the clinical, histopathological and biochemical features of DMRV/h-IBM, making it useful for understanding the pathomechanism of this myopathy and for employing different strategies for therapy. Our findings underscore the notion that hyposialylation plays an important role in the pathomechanism of DMRV/h-IBM.

INTRODUCTION

Distal myopathy with rimmed vacuoles (DMRV) is an autosomal recessive myopathy which was originally reported by Nonaka *et al.*, (1) thus it is also known as Nonaka myopathy. It is the same entity with hereditary inclusion body myopathy (h-IBM), which was initially reported among Iranian Jews (2). DMRV/h-IBM usually starts affecting adults from ages 15 to 40 years with an average onset of 26 years and with an initial symptom of altered gait (1). It is gradually progressive, and patients become wheelchair-bound between 26 and 57 years of age, or about an average of 12 years after the onset of symptoms (3).

DMRV/h-IBM is characterized by preferential involvement of the distal muscles of lower extremities especially the tibialis anterior muscles, with relative sparing of the quadriceps, hence the term 'quadriceps-sparing' rimmed vacuolar myopathy (2). Other muscles are involved as well, especially late in the course of the disease (3). Serum creatine kinase (CK) level is normal or mildly elevated. The characteristic finding on muscle biopsy is the presence of rimmed vacuoles (RVs), which are actually empty spaces surrounded by aggregation of autophagic vacuoles. These RVs occasionally contain congophilic materials that are immunoreactive to various proteins, including amyloid β , phosphorylated tau, ubiquitin and

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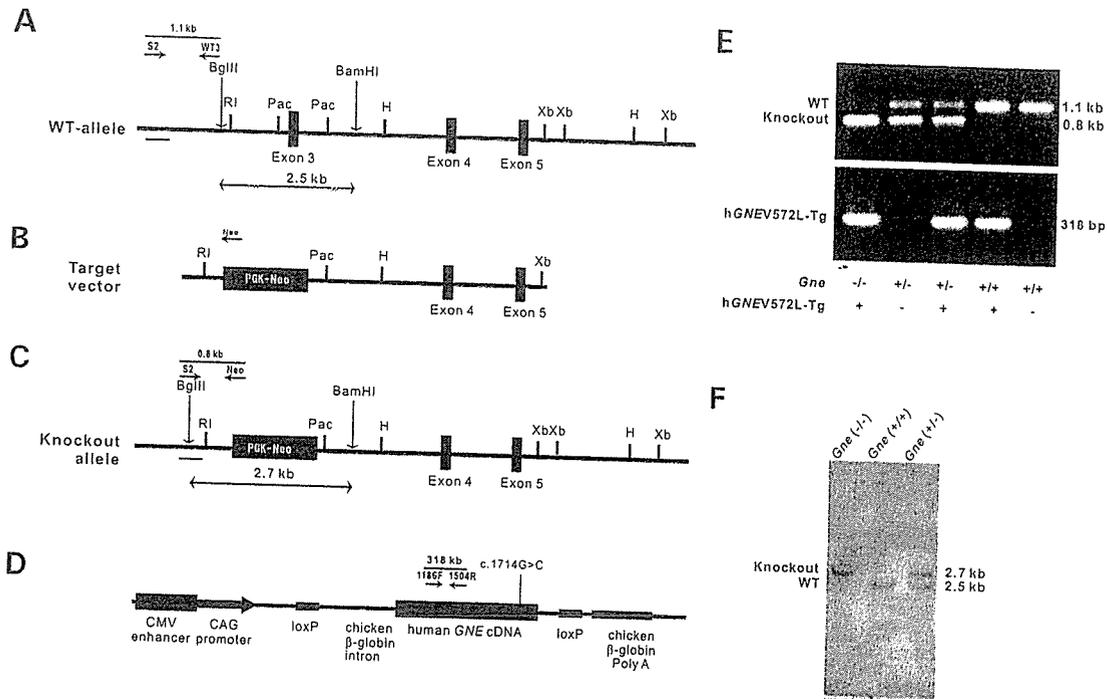


Figure 1. Genomic configuration of the *Gne* gene and targeting vector. Schematic structure of the WT *Gne* gene (A), which contains exon 3–5. Positions of restriction enzyme sites are shown. H, *Hind*III; Pac, *Pac*I; RI, *Eco*RI; Xb, *Xba*I. A 10.5 kb *Eco*RI–*Xba*I fragment was subcloned to make the targeting vector (B). One side of the Neo cassette was inserted 1.4 kb upstream of exon 3, and the other side of which was inserted 124 bp downstream of exon 3. In the knockout allele (C), the Neo cassette replaced the 1.4 kb upstream of exon 3, exon 3 and 124 bp downstream of exon 3. We selected homologous recombinants by neomycin resistance. (D) Structure of mutated human *GNEV572L* construct. CAG promoter was used to achieve expression in various tissues. (E) Genotyping of *Gne* and h*GNEV572L*-Tg mice by PCR. Five genotypes resulting from crossing a $Gne^{+/-}$ h*GNEV572L*-Tg and $Gne^{+/-}$ are shown [from left to right: $Gne^{+/-}$ h*GNEV572L*-Tg (+); $Gne^{+/-}$ h*GNEV572L*-Tg(-); $Gne^{+/-}$ h*GNEV572L*-Tg(-); $Gne^{+/+}$ h*GNEV572L*-Tg(+); and $Gne^{+/+}$ h*GNEV572L*-Tg(-)]. In *Gne* genotyping (upper panel), the 0.8 and 1.1 kb bands represent knockout and WT alleles, respectively [PCR products amplified by primer sets are illustrated in (A) and (C)]. In h*GNEV572L*-Tg genotyping (lower panel), the presence of 318 bp band represents the integration of the h*GNEV572L* transgene. (F) Confirmation of *Gne* genotypes by Southern blot analysis. Tail genomic DNA were digested with *Bgl*II and *Bam*HI and analyzed by Southern blot analyses with 5' probes shown as bars in (A) and (C). The fragments of 2.7 and 2.5 kb represent knockout and WT alleles, respectively.

α -synuclein. Necrotic and regenerating fibers and areas of inflammation are uncommon but can be seen. Ultrastructurally, filamentous inclusions measuring 18–20 nm in diameter are seen in both the cytoplasm and nucleus (3), in addition to the presence of autophagy and various inclusions.

DMRV/h-IBM was mapped to chromosome 9 (4,5), and was shown to be associated with mutations in the *GNE* gene (6,7), which encodes for a bifunctional enzyme that catalyzes the rate-limiting step in sialic acid biosynthesis (8). All patients acquire the disease by autosomal recessive pattern, and have at least one missense mutation in one allele, including the most common mutations V572L and M712T among Japanese and Iranian Jews, respectively. No patient with homozygous null mutation was identified. Genetically confirmed DMRV/h-IBM diseases, initially recognized among Japanese and Iranian Jews (6,9–11), appear to afflict patients with diverse nationalities and cultural backgrounds (12–16).

The mechanism by which mutations in the *GNE* lead to the phenotype in DMRV/h-IBM has remained unclear. We previously demonstrated that mutations in the *GNE* led to the reduction in either the UDP-GlcNAc 2-epimerase or ManNAc kinase activity (17); moreover, we have shown that myotubes from DMRV patients are hyposialylated, and this phenomenon can be corrected by the addition of free sialic acid and/or its precursor. Other groups have shown

similar results regarding *GNE* activity, but in contrast, they suggested that only the cells derived from a patient carrying a homozygous epimerase mutation had a significant reduction in the overall membrane-bound sialic acid (18), and that *GNE* mutations may not contribute to alteration in sialylation in h-IBM myoblasts (19). To address these issues, we developed a mouse model for the disease. In this article, we present the first DMRV/h-IBM mouse model that expressed only the mutated human *GNE* and show that this mouse evidently displays features of DMRV/h-IBM seen in human patients.

RESULTS

Production of $Gne^{-/-}$ h*GNEV572L*-Tg

In Figure 1, the genomic configuration of the *Gne* gene (A) and the targeting construct (B) are shown. The inserted Neo cassette replaced the 1.4 kb upstream of exon 3, exon 3 and 1.4 kb downstream of exon 3. Only WT and $Gne^{+/-}$ mice were generated; no $Gne^{-/-}$ mouse was produced (data not shown), in concurrence with a previous report (20). We then proceeded to generate a transgenic mouse (h*GNEV572L*-Tg) that expressed the human mutated *GNE* with V572L, the most common *GNE* mutation in Japan, the structure of which is shown in Figure 1D. Of the resulting litters, nine

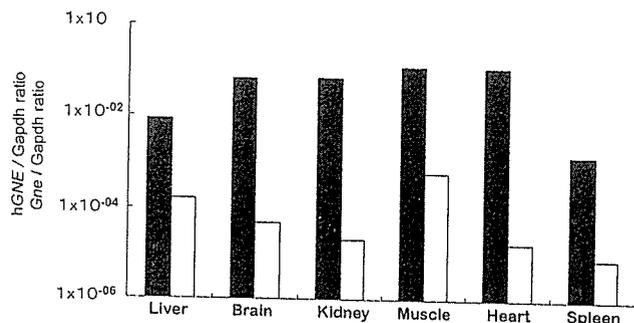


Figure 2. Expression of hGNEV572L (closed bars) and endogenous *Gne* (open bars) relative to Gapdh. Log-values were used for computation of ratio. Highest expression of hGNEV572L is seen in skeletal and cardiac muscles, followed by heart, kidney, brain, liver and spleen.

mice were found to incorporate the hGNEV572L by PCR analysis of genomic DNA isolated from tail snips, but only four lines were able to generate offspring. Using quantitative RT-PCR, we quantified mRNA expression of hGNEV572L and endogenous *Gne* in muscle and other organs of these transgenic mice. Transgene expression was highest in the skeletal muscle, followed by heart, kidney, brain, spleen and liver (Fig. 2), while endogenous *Gne* expression was barely detected. We also determined the copy number using quantitative PCR by comparing the amplification of hGNEV572L with endogenous *Gne*. We calculated the copy numbers for lines 3, 6, 7 and 9 as 2, 4, 3 and 5, respectively. Consequently, we used this transgenic line 9 for producing the model mouse.

We crossed the hGNEV572L transgenic mouse with a *Gne*^(+/-) to obtain a *Gne*^(+/-)hGNEV572L-Tg. Further, we crossed this *Gne*^(+/-)hGNEV572L-Tg with a *Gne*^(+/-) to generate our model mouse, a transgenic mouse on a *Gne* knockout background, *Gne*^(-/-)hGNEV572L-Tg. Analysis of 823 newborn mice from independent heterozygous crosses indicated that the numbers of mice with the five genotypes, *Gne*^(-/-)hGNEV572L-Tg(+), *Gne*^(+/-)hGNEV572L-Tg(+), *Gne*^(+/-)hGNEV572L-Tg(-), *Gne*^(+/+)hGNEV572L-Tg(+) and *Gne*^(+/+)hGNEV572L-Tg(-) were 72 (9%), 225 (28%), 193 (24%), 177 (22%) and 136 (17%), respectively, almost approximating the expected ratio of Mendelian inheritance. Mice of the latter four genotypes did not demonstrate unusual phenotype, and thus were considered as control littermates. Images for routine PCR for checking *Gne* genotype and the incorporation of the human *GNEV572L* are shown in Figure 1E, with the corresponding PCR fragments illustrated in A, C and D.

Hyposialylation is evident in the *Gne*^(-/-)hGNEV572L-Tg

Understandably, mutations in the *GNE* can affect sialylation of glycoconjugates because of the gene's role in sialic acid synthesis. We therefore measured the sialic acid levels in the *Gne*^(-/-)hGNEV572L-Tg mice using HPLC with fluorometric detection. In wild-type (WT) mice, sialic acid levels are highest in the brain, followed by the liver, spleen and kidney (Fig. 3B, open boxes). In both skeletal and cardiac muscles, sialic acid levels are evidently lower than in other

tissues. As we have expected, the total sialic acid in the *Gne*^(-/-)hGNEV572L-Tg mice is remarkably lower than WT. This hyposialylation is most remarkable in the serum (Fig. 3A). A significant reduction in total sialic acid level is seen in various tissues examined (Fig. 3B, closed boxes). We also measured sialic acid level in the hGNEV572L-Tg and noted that sialic acid levels are comparable with WT mice (Fig. 3B, gray boxes), although the transgenic expression was extremely higher than endogenous *GNE*.

Gne^(-/-)hGNEV572L-Tg has lower median of survival than littermate

The *Gne*^(-/-)hGNEV572L-Tg mice were indistinguishable from their littermates at birth and seemed healthy (Fig. 4A and B). After 30 weeks of age, these mice weighed less than their littermates (Fig. 4). Significant difference in weight is more pronounced and earlier in female *Gne*^(-/-)hGNEV572L-Tg (Fig. 4C) mice when compared with male (Fig. 4D). To investigate plausible explanations for this difference in weight, we performed gross inspection of the muscles, and found out that some muscles, especially the gastrocnemius, were atrophic in the *Gne*^(-/-)hGNEV572L-Tg when compared with control (Fig. 4E and F), and this finding was more remarkable among females (Fig. 4E).

Surprisingly, the median survival rate were lower in the *Gne*^(-/-)hGNEV572L-Tg as seen in Figure 4B. The cause of death could not be ascertained, but upon necropsy, no external gross abnormalities were seen and the internal organs appeared normal. On pathological examination, five out of 12 (41%) mice that died had RVs in the skeletal muscle; among these five mice, only one was before 40 weeks of age. Twenty five percent had fibrosis and a few RVs in the diaphragm. Thirty-three percent had fibrosis in the cardiac muscles.

Gne^(-/-)hGNEV572L-Tg shows clinical phenotype

Using a tool for assessing general muscle strength, these mice notably performed worse than their littermates (Fig. 5A). Interestingly, significant change in muscle power is noted after 30 weeks of age. We then proceeded to measure serum CK activity in the mice and found out that CK was significantly elevated in the *Gne*^(-/-)hGNEV572L-Tg mice when compared with their littermates (Fig. 5B), albeit the observation that these values are much lower when compared with muscular dystrophy models like *Large*^{myd} and *Sgcb*^(-/-) mice (data not shown). Because the appearance of phenotype seemed to be related to age, we measured CK activity according to different age groups. From Figure 5C, we note that elevation of CK activity starts at 30 weeks of age. Using gel electrophoresis, we verified that CK-MM isozyme was primarily increased (data not shown).

Gne^(-/-)hGNEV572L-Tg shows characteristic pathological features of DMRV

We checked if the *Gne*^(-/-)hGNEV572L-Tg mice showed muscle phenotype not only by the analysis of muscle power

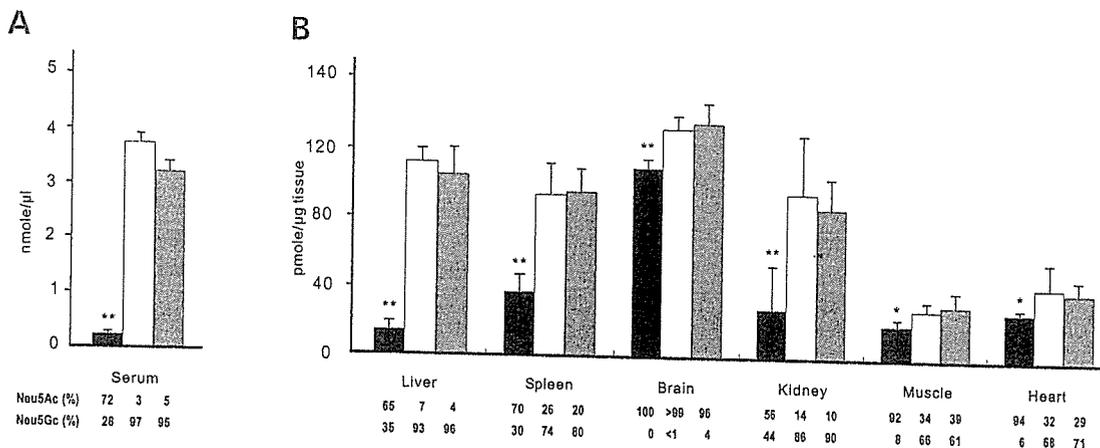


Figure 3. Measurement of total sialic acid in serum (A) and tissues (B) in *Gne*^{-/-}hGNEV572L-Tg (closed bars), WT (open bars) and hGNEV572L-Tg (gray bars); bars represent mean total sialic acid level with SD. Breakdown of sialic acid contents according to standards used (Neu5Ac and Neu5Gc) are shown below. Sialic acid levels of WT and hGNEV572L-Tg are comparable. Note the reduction of sialic acid levels in the serum and tissues of the *Gne*^{-/-}hGNEV572L-Tg mice. Single asterisk, *P* < 0.05; double asterisks, *P* < 0.005.

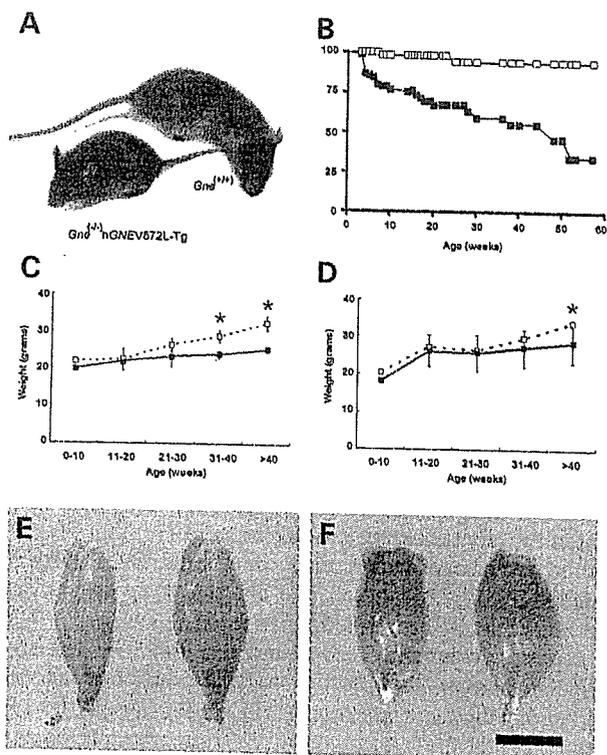


Figure 4. Overall phenotype of the *Gne*^{-/-}hGNEV572L-Tg mice. (A) Picture of WT (up) and *Gne*^{-/-}hGNEV572L-Tg (down) at 16 weeks of age. (B) Survival curve of mice: WT (open squares, *n* = 71 for males and females), *Gne*^{-/-}hGNEV572L-Tg mice (closed squares, *n* = 71 for males and females). *Gne*^{-/-}hGNEV572L-Tg mice have significantly reduced life spans when compared with WT mice (log-rank test: *P* = 0.001). (C) Growth curves of female: WT (open squares, *n* = 10 per age group), *Gne*^{-/-}hGNEV572L-Tg mice (closed squares, *n* = 10 per age group) (D) Growth curves of male mice: WT (open squares, *n* = 10 per age group), *Gne*^{-/-}hGNEV572L-Tg (closed squares, *n* = 10 per age group). *Gne*^{-/-}hGNEV572L-Tg mice have lower body weight after 30 weeks of age, and this is more pronounced in females; asterisk denotes *P* < 0.05, Mann-Whitney *U* test. Gastrocnemius muscles of female (E) and male (F) mice: *Gne*^{-/-}hGNEV572L-Tg (left) and littermate (right). Atrophy is noted on gross inspection of hGNEV572L-Tg gastrocnemius muscles; bar represents 5 mm.

but also by evaluating biopsy samples in five different age groups (10, 20, 30, 40 and 50 weeks). Morphometric analysis of the fibers in different age groups showed that the variation in fiber size becomes more marked with age, preferentially affecting the gastrocnemius and quadriceps muscles (data not shown); for this reason, we used the gastrocnemius muscle in further experiments. We found that at young age, they developed neither clinical nor pathological phenotype, as they were comparable with WT. Figure 6A, D and G shows representative sections from the *Gne*^{-/-}hGNEV572L-Tg mice per age group. Histopathological analysis revealed almost normal findings in the muscle sections before 30 weeks of age (data not shown). In general, necrotic and regenerating processes are not observed in the young mice, although a few necrotic fibers are observed as they grow older. No endomysial or perimysial inflammation is seen.

Scattered small angular fibers are noted by 30 weeks of age (Fig. 6A), which is not seen in the littermates (Fig. 6B and C). The variation in fiber size becomes more noticeable as the mice grow older. In addition, fibers appear atrophic by 40 weeks of age, in support with the observation that the gastrocnemius muscle is relatively atrophic by gross inspection. Remarkably after 40 weeks, RVs are seen in scattered fibers (arrows in Fig. 6D and G). Occasionally, inclusion bodies are found in the fibers with or without RVs (arrowhead in Fig. 7A and B). Like in humans, these RVs are intensely stained with acid phosphatase, giving the impression that autophagic process is activated (Fig. 7C). We confirmed this by checking the expression of lysosomal-associated proteins (LAMPs) 1 and 2, and LC3 in muscle sections, all of which are upregulated. LAMP-1 is predominantly expressed within the vicinity of RVs (Fig. 7E). LAMP-2, on the other hand, noticeably is also expressed in the subsarcolemmal areas aside from its localization in the area of RVs (Fig. 7F). LC3 immunoreactivity is almost similar to LAMP-2, except that the perinuclear region is also highlighted (Fig. 7G).