

Myotonia congenita with strabismus in a large family with a mutation in the *SCN4A* gene

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Abstract

Background/Aims To determine the genetic basis of myotonia congenita (MC) and strabismus in a large Caucasian family. **Methods** Seven patients making up four generations of a family with MC and strabismus were recruited. All patients had at least one standard ophthalmic examination, including best-corrected visual acuity, refraction, and ocular motility measurements. *CLCN1* and *SCN4A* genes were sequenced and analysed for mutations.

Results Five out of the seven family members were diagnosed with MC by clinical history and electromyography. Ophthalmic history and exam revealed eyelid myotonia and strabismus. All patients with MC were diagnosed with strabismus between the ages of 3 and 6 and required surgical restoration of ocular alignment. Sequencing results revealed a c. 1333G > A; p. Val445Met mutation in the *SCN4A* gene. **Conclusion** There are few reports describing eyelid myotonia and strabismus in patients diagnosed with MC. We found significant ocular involvement in a family with a mutation in *SCN4A*. Future studies may confirm that MC with significant ocular involvement can be used to direct genetic analysis.

Eye (2012) 26, 1039–1043; doi:10.1038/eye.2012.80; published online 1 June 2012

Keywords: myotonia congenita; strabismus; esotropia; genetics

Introduction

Non-dystrophic myotonic (NDM) syndromes are a heterogeneous group of skeletal muscle disorders that can be caused by mutations in

genes encoding chloride (*CLCN1*) or sodium (*SCN4A*) channels. *CLCN1* and *SCN4A* encode ion channels in the t-tubules of muscle cells and are important for skeletal muscle fibre contraction.

Mutations in *SCN4A* can cause a range of sodium channel myotonias, including paramyotonia congenita, hyperkalemic periodic paralysis, and potassium-aggravated myotonias. Mutations in *CLCN1* can cause myotonia congenita (MC), a rare disorder with an estimated worldwide prevalence of 1 in 100 000. However, the prevalence varies between geographical regions, with an estimated prevalence in Scandinavia of 1 in 10 000.¹ The two major types of MC are Becker disease and Thomsen disease. Thomsen disease is autosomal dominant with early onset and mild myotonia.² Becker disease is autosomal recessive with late onset and severe myotonia.³

Earlier, only 40–70% of individuals diagnosed clinically with MC were found to harbour the *CLCN1* mutation. Recently, a study was conducted on patients with either Thomsen or Becker disease, where patients were screened for mutations in *SCN4A* in addition to *CLCN1*. Of all the patients without a mutation in *CLCN1*, ~20% had a mutation in *SCN4A*.⁴ It has also become apparent that mutations in the two different genes can lead to clinically indistinguishable myotonias, while identical mutations within either gene can also result in a spectrum of clinically heterogeneous phenotypes.^{5,6}

In previous studies, ocular manifestations of MC have been described, including eyelid myotonia^{7–9} and saccade alterations.^{10,11} There have been few reports, however, of the disease's association with strabismus. Wakeman *et al*¹¹ reported two cases, a father and a son, with Thomsen MC that were evaluated for ocular motility abnormalities. Magnetic resonance

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Received: 13 January 2012
Accepted in revised form: 13 March 2012
Published online: 1 June 2012

imaging revealed ocular muscle hypertrophy and genetic analysis revealed a mutation in *SCN4A*. Another study described four women from a family with Thomsen disease and associated strabismus, but the disease-causing mutation was not reported.¹⁰ In this study, we discuss the largest reported family with both strabismus and MC with a mutation in *SCN4A* gene.

Materials and methods

Participants and clinical examinations

This study was approved by the Institutional Review Board of the University of California San Diego (UCSD), USA. The subjects were recruited at the Shiley Eye Center and the Abraham Ratner Children’s Eye Center at UCSD. All subjects signed informed consent before participation in this study. Five out of the seven members in the family were diagnosed with MC by clinical history and electromyography. All participants had at least one standard complete ophthalmic examination. Examinations included best-corrected visual acuity measurements, refraction, and ocular motility measurements.

The family being studied consisted of seven Caucasian individuals, including five affected and two unaffected with MC (Figure 1). The proband (IV-1) and her mother (III-3) were seen for their ophthalmic treatments at the UCSD Abraham Ratner Children’s Eye Center. However, the other family members were seen for treatment of their strabismus by different ophthalmologists. Therefore, clinical records from their original diagnosis were not available to us. A thorough history was taken and a current ophthalmic exam was performed.

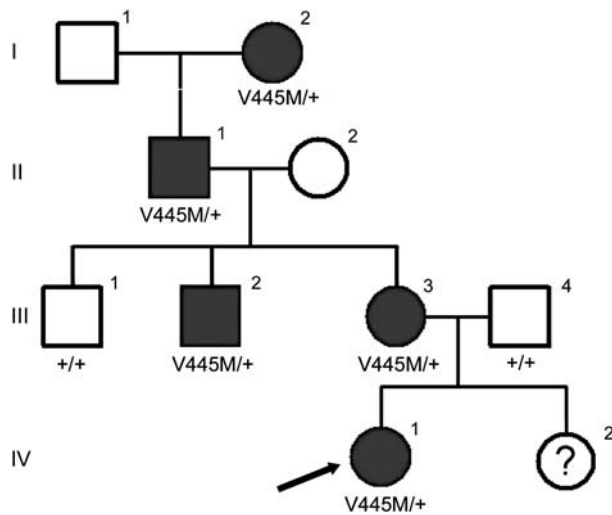


Figure 1 Pedigree of a family affected by MC. The arrow points to the proband. Results from the genetic analysis of *SCN4A* are depicted beneath each investigated individual. Only affected individuals (I-2, II-1, III-2, III-3, and IV-1) harbour the V445M mutation.

DNA sequencing

Genomic DNA samples from all seven family members were extracted from peripheral blood leucocytes with the Qiagen kit (Qiagen Inc., Chatsworth, CA, USA) according to the manufacturer’s instructions. Direct nucleotide sequence analysis was completed for both *CLCN1* and *SCN4A*. The 23 exons of *CLCN1* and 24 exons of *SCN4A* were amplified by PCR and sequenced on Genetic Analyzer 3130 (Applied Biosystems, Carlsbad, CA, USA). The primers used to amplify the exons in *CLCN1* and *SCN4A* are presented in Supplementary Table S1.

Results

Clinical examinations

The family consists of seven members, five of which were diagnosed with MC (Figure 1). Clinical history and exams of the five affected family members are presented in Tables 1 and 2. All affected individuals were diagnosed with esotropia between the ages of 3 and 6, and had at least one surgery in both eyes for strabismus. All affected family members reported myotonic symptoms, including muscle stiffness, upon standing and after exercising and improvement of stiffness with repeated muscle contractions. Several individuals described experiencing eyelid myotonia, delayed opening of the eyes after forceful closure. Of note, III-3 also had increased myotonic symptoms during pregnancy and a history of malignant hyperthermia, an associated risk of MC.¹²

The proband (IV-1) was presented at 5 months of age with esotropia and eyelid myotonia. Other than ocular involvement, she was only mildly affected by myotonia. Despite patching and full cycloplegic spectacles, she developed amblyopia of the left eye by age 3. She subsequently underwent bilateral medial rectus recession and had a positive outcome.

Genetic results

The family history is consistent with an autosomal dominant inheritance pattern (Figure 1). We screened all exons of *CLCN1* and *SCN4A*. All five affected individuals were found to have a G > A transition (c. 1333G > A) in exon 9 of *SCN4A* gene, resulting in a V445M substitution. This variation was absent in the two unaffected family members (Figure 2). The change (c. 1333G > A) was also absent in 1000 normal control chromosomes. No novel or previously reported mutations were found in *CLCN1* gene in all the affected family members.

Discussion

We report here clinical features and genetic analysis of the largest family with MC and strabismus. We initially screened

Table 1 Clinical features of patients with the SCN4A mutation

Patient	Pedigree ID	Age	Sex	Age of disease onset	Number of surgeries OU	Eyelid myotonia	Current VA OD	Current VA OS	WRx OD	WRx OS
Maternal great-grandmother	I-2	87	F	5	2	No	20/30	20/25	-0.75 + 1.25 × 025	-1.00 + 1.50 × 175
Maternal grandfather	II-1	65	M	3	2	Yes	20/20	20/20	N/A	N/A
Maternal uncle	III-2	29	M	4	2	Yes	20/20	20/20	N/A	N/A
Mother	III-3	32	F	5	5	Yes	20/20	20/20	+1.00	-0.75
Proband	IV-1	3	F	3	1	Yes	20/30	20/40	+6.00 + 0.50 × 085	+6.00 + 0.50 × 123

Abbreviations: F, female; M, male; N/A, not applicable; VA, visual acuity; WRx, wearing prescription.

Table 2 Ocular motility exam findings of patients with the SCN4A mutation

Patient	Pedigree ID	Primary	Up gaze	Down gaze	Right gaze	Right tilt	Left gaze	Left tilt	Near
Maternal great-grandmother	I-2	ET10 RHT5	ET8 RHT6	ET6 RHT3	ET8 RHT4	ET8 RHT2	ET8 RHT4	ET8 RHT2	RH(T)'3
Maternal grandfather	II-1	E(T)6	E(T)6	E(T)6	E(T)8	E(T)8	E(T)2	E(T)6	Flick
Uncle	III-2	E(T)12 RH(T)2	E(T)12 RH(T)2	E(T)14 RH(T)2	E(T)10	E(T)14	N/A	E(T)14	E'4
Mother	III-3	ET12	N/A	N/A	ET12	ET12	ET10	ET12	ET'6
Proband	IV-1	ET16	ET16	ET16	ET14	N/A	ET14	N/A	N/A

Abbreviations: E', esophoria at near; ET, esotropia; ET', esotropia at near; E(T), intermittent esotropia; N/A, not applicable; RHT, right hypertropia; RH(T), intermittent right hypertropia.

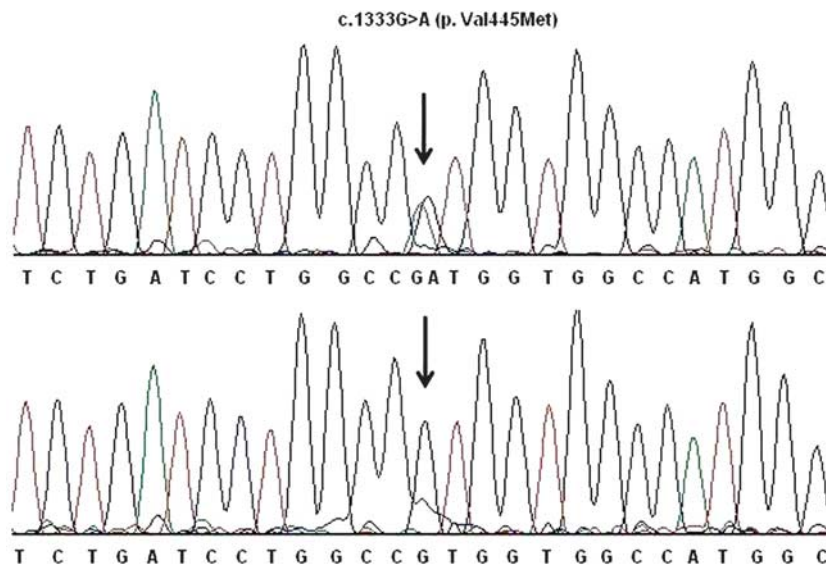


Figure 2 The SCN4A sequencing results. Sequencing data show that a G–A transition in exon 9 (arrow) causes the V445M mutation in affected individuals (top). This alteration was absent in the unaffected individuals (bottom).

for mutations in *CLCN1* based on reports of increased frequency of mutations within this gene in patients with MC.¹⁴ However, as no mutations were found, we screened for mutations in *SCN4A*. *SCN4A* is located on 17q23 and includes 35 kb with 24 exons.¹³ All affected family members were found to have a G>A transition within exon 9 of

SCN4A, resulting in the V445M mutation. This mutation has been associated with debilitating, painful myotonia, and muscle hypertrophy.^{14–16} However, the family we studied did not have these findings, suggesting that other modifying factors may be involved in determining the above-mentioned phenotype.

Results from multiple studies demonstrate that patients with *SCN4A* mutations are more likely to have ocular symptoms, either eyelid myotonia or strabismus.^{8,9,11} One study showed that 96.7% of patients with sodium channel mutations had eyelid myotonia compared with 46.9% of patients with chloride channel mutations. Furthermore, mutations in *SCN4A* cause more severe eyelid myotonia than mutations in *CLCN1*.⁸ An increased number of sodium channels in facial muscles compared with other skeletal muscles may explain the findings associated with *SCN4A* mutations.^{8,17,18} Consistent with our analysis, the only other genetic study performed on patients with strabismus in addition to myotonia also found the V445M mutation.¹⁹ There may be an association between this mutation and extraocular muscles. It has been speculated that esotropia may be caused by a slightly nasally displaced superior and inferior rectus, as well as an inferiorly displaced lateral rectus.¹¹ However, imaging of a larger cohort with more detailed scans are necessary to confirm an association between the mutation and such anatomic anomalies.

In conclusion, even though this is a rare condition, ophthalmologists should be aware of its ocular manifestations. In our study, eyelid myotonia and strabismus in the proband led to the suspicion of MC. As these two ocular findings can present as early as 5–6 months of age, ophthalmologists may be the first to evaluate patients with NDM. Ophthalmologists should be aware of the reported risk of malignant hyperthermia in patients with MC¹² and alert the anaesthesiologist before the use of general anaesthesia. Finally, our results may support the sequencing of *SCN4A* before *CLCN1* in myotonic patients with significant ocular involvement.

Summary

What was known before

- Clinical presentation of myotonia congenita includes a wide range of symptoms and signs, but is often associated with muscle stiffness and hypertrophy.
- *SCN4A* and *CLCN1* genes are associated with myotonia congenita.

What this study adds

- An extended family clinical report of eyelid myotonia and strabismus associated with myotonia congenita.
- Further evidence that mutations in the *SCN4A* gene may be more likely to present with evidence of eyelid myotonia and ocular symptoms in myotonia congenita compared with mutations in other genes associated with the myotonic disease.

Conflict of interest

The authors declare no conflict of interest.

Acknowledgements

We acknowledge Maureen Crocker, Bin Lin, Karen Ysensky and Dr Salma Khayali for their help in this study. KZ is supported by grants from the NEI/NIH (EY014428, EY018660, and EY019270), VA Merit Award, Research to Prevent Blindness, and BWF Clinical Scientist Award in Translational Research. SG is supported by the UCSD Clinical and Translational Research Institute 1TL1RR031979-01.

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Supplementary Information accompanies the paper on Eye website (<http://www.nature.com/eye>)