

# Clinical Reasoning: Cardioembolic stroke in a 23-year-old man with elbow contracture

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## Section 1

A 23-year-old right-handed man with a diagnosis of spinal muscular atrophy (SMA) type 3 presented with right facial droop, hemiparesis, and expressive aphasia. CT angiography revealed a thrombus in the distal left middle cerebral artery. IV tissue plasminogen activator was administered and he recovered without any residual deficit. His cardiac monitoring revealed episodes of atrial fibrillation (figure, C) and episodes of narrow junctional escape rhythm in 20–30 s. ECG showed normal left ventricular ejection fraction (>55%). Cardiac MRI showed moderate dilation of right atrium, moderate left atrial elongation, and severely increased left ventricular cavity size with normal systolic function.

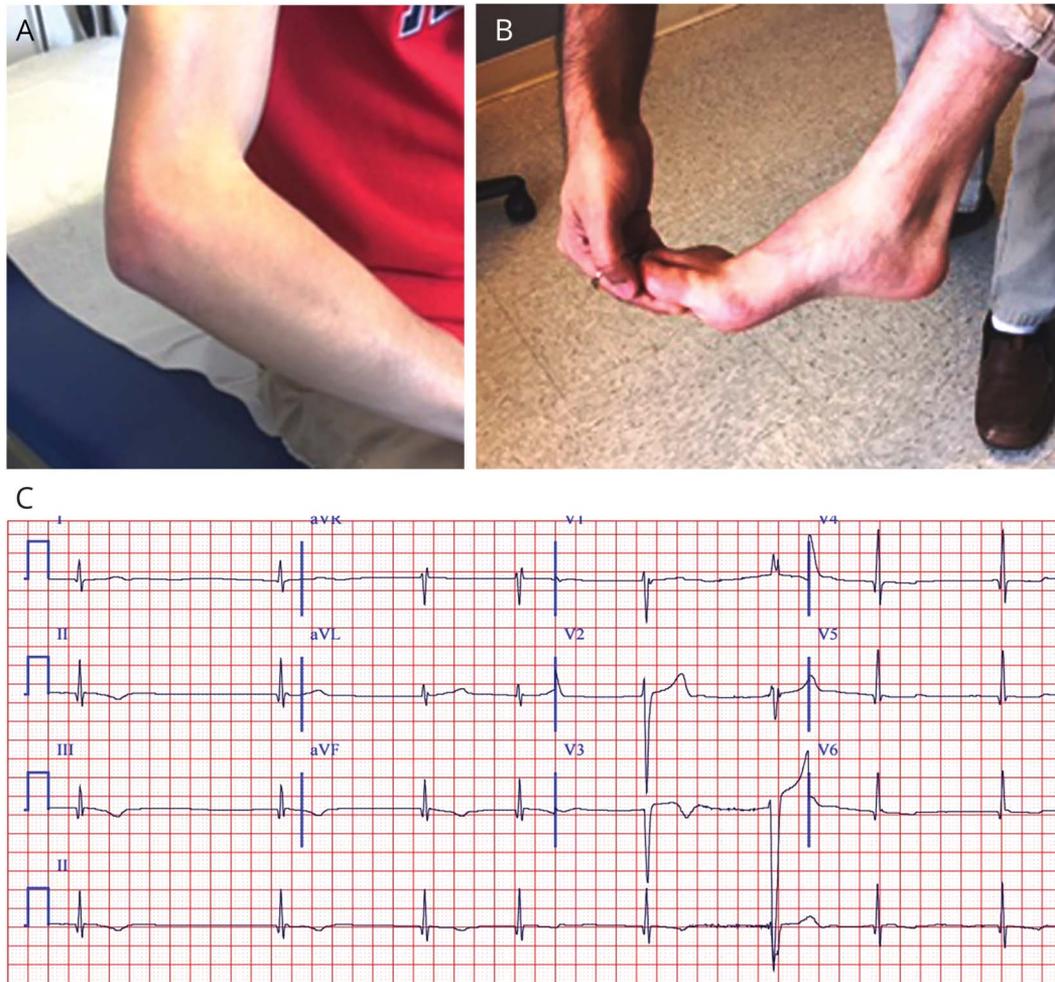
Neurologic examination revealed muscle atrophy in a predominantly humeroperoneal distribution. There was mild weakness in deltoid and triceps (4+/5 Medical Research Council [MRC] scale) and moderate weakness in ankle dorsiflexion (3+/5 MRC scale) bilaterally. There were prominent elbow and heel contractures along with exaggerated lumbar lordosis and cervical spine rigidity (figure, A and B). Facial and bulbar muscles were spared. Mild neck flexion weakness was noted. There was no scapular winging. Sensation was intact to all modalities. Reflexes were globally absent. The patient walked on his toes with normal stride.

The patient had an uncomplicated birth history and attained motor milestones appropriately. He was a slow runner and could not keep up with his peers. He was diagnosed with SMA type 3 at age 3 based on EMG and muscle biopsy. Around age 8, he started to develop elbow and ankle contractures. Eventually, he became a toe walker and had frequent falls. He noticed gradual loss of muscle bulk in upper and lower extremities, especially around the shoulder and hip girdle without severe proximal weakness. He had a normal cardiac evaluation with ECG and echocardiogram at age 14.

## Questions for consideration:

1. Is this phenotype consistent with the diagnosis of SMA type 3?
2. What is your differential diagnosis?

**Figure** Elbow and heel cord contractures and ECG findings



(A) Elbow contracture. (B) Heel cord contracture. (C) ECG shows atrial fibrillation.

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## Section 2

Patients with SMA type 3 present with progressive proximal muscle atrophy and weakness, but early development of joint contractures and cardiac involvement is unusual. Joint contractures can be seen in SMA type 1 and type 2 patients only in the setting of severe weakness in nonambulatory patients. Occasionally, early contractures of the heel cord develop in SMA with lower extremity predominance (SMA-LED) from *BICD2* mutation or in distal SMAs with *TRPV4* mutation. These non-5q SMAs are usually associated with intrauterine involvement or delayed motor milestones, and do not affect the heart. Moreover, elbow contractures would be distinctly unusual in these disorders.<sup>1</sup> This patient did not have typical phenotype of SMA.

In contrast, early flexion contractures of tendons, especially at the elbow, are a hallmark feature of certain inherited muscular dystrophies, including Emery-Dreifuss muscular dystrophy (EDMD), limb-girdle muscular dystrophy 1B (LGMD 1B), and Bethlem myopathy.<sup>2</sup>

Early-onset elbow, heel, and posterior cervical muscle contractures, humeroperoneal distribution of muscle atrophy and weakness, and cardiac conduction defects are cardinal features of EDMD. EDMD can be X-linked, autosomal dominant (AD), or autosomal recessive (AR). X-linked EDMD (EDMD1) is caused by a mutation in *STA* located in chromosome xq28 encoding for emerin. Mutations in the *FHL1* gene can also present with EDMD phenotype (EDMD6). Mutations in *LMNA* can lead to both AD (EDMD2) and AR form of EDMD (EDMD3). Rarely, sporadic cases and AD-EDMD have been reported in mutations of *SYNE1* encoding

nesprin-1 (EDMD4), *SYNE2* encoding nesprin-2 (EDMD5), and *TMEM 43* encoding LUMA (EDMD7).<sup>3</sup>

AD-EDMD and LGMD 1B are overlapping syndromes and allelic disorders caused by mutations in *LMNA*. However, in AD-EDMD, muscle wasting occurs in a humeroperoneal distribution and in LGMD 1B contractures are minimal or absent.<sup>4</sup> Cardiac involvement in AD-EDMD is age-independent and can occur in isolation without detectable muscle manifestations. On the other hand, the severity of atrioventricular (AV) conduction disturbances in LGMD 1B is usually age-related and neuromuscular involvement almost always precedes cardiac involvement. Cardioembolic strokes have been reported in both disorders but more frequently in EDMD.<sup>5,6</sup>

Bethlem myopathy is an AD disorder caused by mutations in *COL6A1*, *COL6A2*, and *COL6A3*. Contractures at the elbow and ankle may be evident, occasionally even before the manifestation of any weakness. However, some patients may have proximal hip and shoulder girdle weakness without contractures. Contractures can be preceded by joint laxity. About 10% of cases have cardiac involvement including arrhythmia, intraventricular conduction delay, and echocardiogram abnormalities.<sup>7</sup>

Clinically, this patient's phenotype is most consistent with EDMD but a definitive diagnosis cannot be made solely based on clinical features.

### Question for consideration:

1. What is the next best diagnostic test?

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### Section 3

Genetic testing would be the next best step. EMG or a muscle biopsy would not discriminate among the disorders under consideration.

Focused genetic testing was performed, which revealed heterozygous mutations in the *LMNA* gene on c.122G>A (p. Arg41His) and in the *TMEM43* gene at c.796C>T (p. Arg266Trp). No mutations or exonic deletion/duplications were noted in *COL6A1*, *COL6A2*, *COL6A3*, *EMD*, *FHL1*, or *SYNE1*; thus X-linked EDMD and Bethlem myopathy were ruled out. *TMEM43* mutation at c.796CC>T has previously been reported in arrhythmogenic right ventricular cardiomyopathy type 5.<sup>3</sup>

Though previously not reported, the *LMNA* mutation is most likely pathogenic in this patient. The *LMNA* gene located in chromosome 1q22 is composed of 12 exons and encodes 4 lamins by alternative splicing. The lamin A and C main isoforms are both nuclear envelope proteins. *LMNA* can be involved in both EDMD2 and LGMD 1B. They were the first 2 disorders reported to be associated with *LMNA* mutations. Subsequently, over 400 *LMNA* mutations have been reported in more than 10 distinct genetic diseases, including dilated cardiomyopathy associated with conduction system disease, congenital muscular dystrophy (L-CMD), AR form of

Charcot-Marie-Tooth type 2B1, Dunnigan type familial partial lipodystrophy, Hutchinson-Gilford progeria syndrome, and mandibuloacral dysplasia.<sup>3</sup>

Different mutations in *LMNA* can lead to AD-EDMD. Missense mutations are common and 65% of patients have de novo mutations. Initially, mutations in the rod domain were thought to be solely associated with cardiomyopathy and conduction defects, sparing skeletal muscles; however, various rod domain mutations are now known to cause the full spectrum of EDMD.<sup>8,9</sup> Usually, patients with missense mutations have early skeletal muscle involvement, whereas frameshift mutations have later onset of muscle symptoms.<sup>3,8,9</sup>

For our patient, this unique mutation in the rod domain was initially considered as a variant of unknown significance. Parental genetic testing did not show the same mutation, confirming the de novo origin and pathogenic nature of the mutation. This exact mutation has not been reported earlier in EDMD; however, another missense mutation (c.122G>T) at the same location has been reported in a patient with classic EDMD with atrial standstill.<sup>9</sup>

#### Questions for consideration:

1. What is the most serious complication of this disease?
2. How would you manage this patient?

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## Section 4

Based on the clinical presentation and genetic testing, our patient has EDMD2. Development of joint contractures can be delayed by a few years in EDMD2 (usually between 3 and 6 years) compared to EDMD1, where the contractures can be the earliest symptom. Weakness in EDMD is generally not profound but cardiac involvement is common and leads to serious complications, including heart failure and sudden cardiac death (SCD).<sup>6,8</sup>

Generally, cardiac manifestations develop with the progression of muscle weakness and after the second decade of life. Cardiac conduction defects are common and range from sinus bradycardia, prolongation of the PR interval, AV conduction block, to complete heart block. Atrial flutter or fibrillation can develop later and may evolve into atrial paralysis. Junctional escape rhythm (40–50 beats/min) without obvious p-waves is the most characteristic ECG finding and atrial paralysis is almost pathognomonic.<sup>6,10</sup>

Frequent cardiac monitoring is essential in EDMD. ECG, Holter monitoring, and 2D echocardiogram every 6–12 months is recommended in asymptomatic patients. National guidelines endorse permanent pacing for patients with EDMD with any degree of AV block, even if asymptomatic, as the progression of AV conduction disease is unpredictable. SCD can occur even with pacemaker placement. In the absence of guidelines, some authors have suggested prophylactic implantable cardioverter–defibrillator (ICD) placement to prevent possible lethal tachyarrhythmias.<sup>6,10</sup>

Our patient had atrial fibrillation with heart rate in 30s and attempted DC cardioversion failed. A pacemaker and a single chamber ICD were placed. At the time of his stroke, he was started on anticoagulation. He continues to be asymptomatic in terms of cardiac symptoms although episodes of ventricular tachycardia (with cycle length >300 ms) have been noted on ICD interrogation.

Cardioembolic stroke in a young patient with signs of skeletal muscle involvement should raise the suspicion of an underlying muscular dystrophy. This patient carried a lifelong diagnosis of SMA type 3, although the cardiac conduction abnormalities combined with prominent elbow and heel cord

contractures and a humeroperoneal pattern of weakness suggested an alternative diagnosis. Genetic testing confirmed EDMD2 with a novel mutation in *LMNA*. Early recognition of clinical features and diagnosis of EDMD is critical so that close monitoring and appropriate treatment of cardiac conduction abnormalities may be undertaken to prevent further cardiac complications, including SCD.

## Author contributions

Bhaskar Roy: study concept and design, acquisition of data, analysis and interpretation of data, drafted the manuscript. Elizabeth Raynor: study concept and design, critical revision of manuscript for intellectual content, study supervision.

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## Disclosure

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