Anti-3-Hydroxy-3-Methylglutaryl Coenzyme A Reductase (Anti-HMG CoA) Myopathy With Cardiac Involvement: Presentation, Diagnosis, and Management

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Abstract

Immune-mediated necrotizing myopathy (IMNM) is categorized into three groups: anti-3-hydroxy-3-methylglutaryl coenzyme A reductase (HMGCR) IMNM, anti-signal recognition particle (SRP) IMNM, and seronegative IMNM. Cardiac involvement has been reported in a significant segment of patients with IMNM of the anti-SRP type. Emerging evidence now suggests that cardiac involvement is also implicated in the anti-HMGCR subgroup. In this report, we present a case of anti-HMGCR IMNM with cardiac involvement demonstrated by elevated troponin levels, a low ejection fraction of 40%, and regional wall motion abnormalities in the inferior, inferolateral, anteroseptal, inferoseptal, and anterolateral myocardial walls, as visualized on echocardiography. These findings markedly improved after treatment with intravenous immunoglobulin (IVIG) and prednisone. This case and other recent reports highlight the need for a cardiac workup in patients diagnosed with anti-HMGCR IMNM.

Categories: Cardiology, Allergy/Immunology, Rheumatology
Keywords: autoantibodies, immune-mediated necrotizing myopathy, statin, cardiac involvement, myopathy

Introduction

Statins are commonly prescribed as lipid-lowering medications, and they function by interfering with cholesterol synthesis. Their mechanism of action involves the inhibition of the rate-limiting enzyme 3-hydroxy-3-methylglutaryl coenzyme A reductase (HMGCR). The expression of HMGCR is upregulated in muscle cells following exposure to statins. In addition, regenerating muscle cells express high levels of HMGCR necessary for normal muscle cell differentiation [1]. However, in rare cases, the use of statins can lead to an immune-mediated necrotizing myopathy (IMNM), a recently recognized subtype of idiopathic inflammatory myopathies [2-4]. While the original categorization of idiopathic inflammatory myopathies included polymyositis and dermatomyositis, a new classification system was developed by the European Neuromuscular Centre in 2005 and later expanded in 2016 to include anti-HMGCR IMNM, anti-signal recognition particle (SRP) IMNM, and seronegative IMNM [5,6].

The pathophysiology of statin-induced-myopathy is associated with the binding of anti-HMGCR antibodies to the surface of myofibers, resulting in complement activation, recruitment of macrophages, and muscle cell necrosis [7]. Furthermore, these antibodies impair muscle regeneration by disrupting myoblast differentiation [8]. Anti-HMGCR IMNM clinically presents with progressive proximal muscle weakness that correlates with elevated creatinine kinase (CK) levels (>1,000–10,000 U/L), indicating skeletal muscle damage [7]. While the anti-HMGCR subgroup was previously thought to lack cardiac involvement [8,9], a recent retrospective study found that 11 out of 36 patients with anti-HMGCR IMNM had echocardiographic abnormalities [10]. In this report, we discuss a case of anti-HMGCR IMNM showing cardiac involvement with elevated cardiac troponins, a low ejection fraction, and extensive regional wall motion abnormalities.

Case Presentation

A 68-year-old female with a history of childhood epilepsy, hypothyroidism, hyperlipidemia, and prior falls was brought to the emergency department (ED) after falling twice. The first fall had occurred while the patient had been getting out of bed in the morning and the second while the patient had bent over to grab a newspaper. The patient denied losing consciousness or injury to the head. She also denied muscle pain, lightheadedness, or dizziness. The patient had had four ED visits in the past year due to falls. Her medications included atorvastatin 40 mg, which had been started five years prior. The physical exam was unremarkable except for proximal weakness (3/5) in the shoulders and hip flexors. The patient’s fine motor coordination skills were intact. She was able to follow commands, conversed without difficulty, made eye contact, and was oriented to person, place, time, and year. The patient did not have any tremors, seizures, facial asymmetry, or speech difficulty, and the sensation was intact. Her vital signs were as follows: blood pressure of 143/61 mmHg, a pulse rate of 92 beats/minute, temperature of 99.5 °F, respiratory rate of 18
breaths per minute, oxygen saturation of 96% on room air, and a body mass index of 29 kg/m². The complete blood count and differentials were within normal limits except for elevated neutrophils at 88% (reference range: 40-60%) and decreased lymphocytes at 9% (reference range: 20-40%). A metabolic panel revealed an elevated alanine transaminase level of 123 U/L (reference range: 7-55 U/L) and an aspartate aminotransferase level of 308 U/L (reference range: 8-33 U/L). Vitamin B12 levels were less than 60 pg/mL (reference range: 100-900 pg/mL). Serum CK levels were obtained and found to be elevated at 3575 U/L (reference range: 26-192 U/L). The CK MB isoenzyme (CK-MB) index was 3.0% (reference range: 0.0-2.5%), indicating the involvement of skeletal and cardiac muscles. Troponin levels were not checked during this hospital stay. An electrocardiogram performed on the day of admission showed a normal sinus rhythm with non-specific ST and T wave abnormalities in the lateral leads. The patient was started on B12 injections of 1000 μg intramuscular daily. She was also recommended to follow up with neurology for an electromyogram and was discharged to a short-term rehabilitation facility for physical therapy.

Three weeks later, the patient returned to the ED following another fall. She reported diffuse muscle weakness that limited her ability to ambulate without assistance, get out of a car, walk up steps, and comb her hair. On physical exam, she had difficulty rising from the chair. Grip and biceps strength were relatively preserved with the exception of the left hand, which was weaker. The patient had significantly decreased bilateral triceps and deltoid strength. The CK level was 4805 U/L and CK-MB index was 3.7%. The troponin I level was normal although this was only checked once. Vitals were stable and other lab values were similar to those from the previous admission. CT of the head showed no acute abnormality and the chest X-ray was unremarkable. Atorvastatin was discontinued due to a concern for rhabdomyolysis and elevated liver enzymes. The patient received aggressive intravenous fluid hydration due to rhabdomyolysis.

A myositis serologic panel was negative for ribonucleoprotein (RNP), Mi-2, Ku, SRP, threonyl (PL-7), alanyl (PL-12), glycyl (E), isoleucyl (O), and histidyl (Jo-1). An electromyogram of the right upper and lower extremities later showed a generalized myopathic process with membrane instability, axonal polyneuropathy, and moderate right carpal tunnel syndrome. Anti-acetylcholine receptor (AChR) antibody and anti-HMGCR antibody tests, as well as a muscle biopsy, were obtained. The patient’s AChR binding antibody, AChR blocking antibody, and AChR modulating were negative. However, the anti-HMGCR antibody was positive with a level of 94.5 CU (reference level: <20.0 CU). The muscle biopsy revealed scattered regenerating and necrotic muscle fibers with a limited inflammatory infiltrate (Figure 1).

Two weeks after the muscle biopsy, the patient presented again to the ED with generalized weakness after yet another fall. Physical exam this time revealed mild lower extremity edema. The CK was 3097 U/L and CK-MB index was 7.8%. Troponin I levels were elevated and peaked at 2.36 ng/mL (reference range: <0.04 ng/mL).

An echocardiogram showed a left ventricular ejection fraction of 40% in addition to mid to apical regional wall motion abnormalities in the inferior, inferolateral, anteroseptal, inferoseptal, and anterolateral
myocardial walls. A prior echocardiogram performed six years ago had shown an ejection fraction of 65% with normal left ventricular chamber size, normal systolic and diastolic function, and no regional wall motion abnormalities. The patient did not want to receive a left heart catheterization. She received intravenous immunoglobulin (IVIG) of five cycles and was started on prednisone 60 mg daily for the IMNM. Ten days after admission, the CK levels decreased to 937 U/L and the CK-MB index decreased to 6.0%.

On follow-up two months later, the patient showed an improvement in symptoms and had not sustained any additional falls. She was started on azathioprine 25 mg twice daily and was to be slowly tapered off prednisone. A repeat echocardiogram showed the resolution of the cardiomyopathy with an ejection fraction of 60% and no regional wall motion abnormalities. Images of the patient’s echocardiogram comparing systole and diastole before the treatment with those after the treatment are illustrated in Figure 2. Videos of the patient’s echocardiogram before and after the treatment are shown in Videos 1, 2, respectively. Notably, there was a decreased movement of the heart apex before the treatment compared with after the treatment.

![Echocardiograms before and after treatment](https://youtu.be/SdYX3W78T68)
Echocardiogram after the treatment

View video here: https://youtu.be/WNjRYi3_ygU

The patient has remained asymptomatic and without any hospitalizations for additional falls so far. The patient’s clinical course is illustrated in Table 1.

| March 2014 | September 2014 | April 2015 | January 2020 | February 2020 | March 2020 | April 2020 | May 2020 | June 2020 | July 2020 | August 2020 | September 2020 | October 2020 | November 2020 |
|------------|----------------|------------|--------------|---------------|-------------|------------|-----------|-----------|-----------|--------------|----------------|----------------|---------------|
| Statin use | Started on simvastatin | Switched to atorvastatin | | | | | | | | | | | |
| CK (U/L)   | ↑1507 | ↑14805 | 13,097 | 19,37 | 19,02 | | | | | | | |
| INR        | 1.06 | 1.24 | | | | | | | | | | | |
| Troponin I (ng/mL) | <0.02 | 11.92* | 11.53** | | | | | | | | | | |
| Echo: EF   | 65% | ↓40% | ↓48% | 60% | | | | | | | | | |
| ED visit due to fall | X | X | X | X | X | | | | | | | | |
| EMG        | | | | | | | | | | | X | | |
| HMGCR, AChR Abs test | | | | | | | | | | | | X | |
| Muscle biopsy | | | | | | | | | | | | X | |
| Initiated on IVIG | | | | | | | | | | | X | | |
| Initiated on prednisone | | | | | | | | | | | | X | |
| Initiated on azathioprine | | | | | | | | | | | | | X |

**TABLE 1: Timeline of patient’s treatment course, laboratory values, and imaging results**

Reference values: CK: 26-192 U/L; troponin I: ≤0.04 ng/mL; INR: 0.84-1.19; EF: 50-75%

↑ represents values above the reference range, and ↓ represents values below the reference range. An empty column represents a gap in the timeline

*Average of four values. **Average of three values, at a different visit

Ab: antibody; AChR: acetylcholine receptor; CK: creatine kinase; disc: discontinued; Echo: echocardiogram; ED: emergency department; EF: ejection fraction; HMGCR: 3-hydroxy-3-methylglutaryl coenzyme A reductase; IVIG: intravenous immunoglobulin

**Discussion**

Multiple factors suggest that this patient’s cardiac involvement was secondary to IMNM. Firstly, the
Anti-HMGCR IMNM typically presents with proximal muscle weakness and elevated CK levels [7]. Cardiac involvement in IMNM is not typical and has been limited to small case series so far in the literature [8,9]. Recently, a retrospective study investigating 109 patients with IMNM, who were evaluated for echocardiogram abnormalities, found that of the 36 patients positive for anti-HMGCR antibody, 11 showed left ventricular diastolic dysfunction, and six showed systolic dysfunction [10]. Another study documented cardiac abnormalities in patients with IMNM, although the study did not specify the IMNM subgroups [11].

Our patient showed regional wall motion abnormalities in the inferior, inferolateral, anteroseptal, inferoseptal, and anterolateral myocardial walls on echocardiography, which were reversible following the treatment of IMNM.

A diagnosis of anti-HMGCR IMNM is confirmed when the following three criteria are met: (1) elevated serum CK levels, (2) proximal muscle weakness, and (3) anti-HMGCR antibodies [6]. Initial treatment includes intravenous and/or oral steroids along with the addition of IVIG and/or methotrexate concurrently or within one month [6]. After at least two years of well-controlled disease with minimal or no steroids, other agents should be stopped or tapered off, although many patients do require prolonged IVIG.

This case of cardiac involvement in the anti-HMGCR subgroup and the emerging evidence reviewed herein suggest that a cardiac evaluation should be considered in patients diagnosed with anti-HMGCR IMNM.

Conclusions
Cardiac involvement is a rare finding in anti-HMGCR IMNM. We reported the case of a patient with anti-HMGCR IMNM with cardiac involvement demonstrated by elevated troponin levels, a low ejection fraction, and regional wall motion abnormalities in the inferior, inferolateral, anteroseptal, inferoseptal, and anterolateral walls on echocardiography. The patient’s condition markedly improved after treatment with IVIG and prednisone. This case underlines the need for a cardiac workup in patients diagnosed with anti-HMGCR IMNM.

Additional Information
Disclosures
Human subjects: Consent was obtained or waived by all participants in this study. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

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