A middle-aged man with progressive fatigue

A 61-YEAR-OLD WHITE MAN presents with progressive fatigue, which began several months ago and has accelerated in severity over the past week. He says he has had no shortness of breath, chest pain, or symptoms of heart failure, but he has noticed a decrease in exertional capacity and now has trouble completing his daily 5-mile walk.

He saw his primary physician, who obtained an electrocardiogram that showed a new left bundle branch block. Transthoracic echocardiography indicated that his left ventricular ejection fraction, which was 60% a year earlier, was now 35%.

He has hypertension, dyslipidemia, type 2 diabetes, and chronic kidney disease. Although he was previously morbidly obese, he has lost more than 100 pounds with diet and exercise over the past 10 years. He also used to smoke; in fact, he has a 30-pack-year history, but he quit in 1987. He has a family history of premature coronary artery disease.

Physical examination. His heart rate is 75 beats per minute, blood pressure 142/85 mm Hg, and blood oxygen saturation 96% while breathing room air. He weighs 169 pounds (76.6 kg) and he is 6 feet tall (182.9 cm), so his body mass index is 22.9 kg/m².

He is awake and in no acute distress. His breath sounds are normal, without crackles or wheezes. His heart has a normal rate and regular rhythm; he has normal first and second heart sounds and no extra sounds or murmurs; the apical impulse is not displaced. His abdomen is soft and nontender, with no hepatosplenomegaly or hepatojugular reflex. His extremities are warm and well perfused, with normal peripheral pulses and no edema. He has no gross neurologic defects.

Initial laboratory analysis (TABLE 1) shows evidence of anemia and renal insufficiency and a slightly elevated serum level of glucose. His cardiac biomarkers are within normal limits, but his B-type natriuretic peptide level is 483 pg/mL (reference range < 100 pg/mL). His thyroid-stimulating hormone level is in the normal range.

Electrocardiography reveals sinus rhythm with a left bundle branch block and left axis deviation (FIGURE 1), which were not present 1 year ago.

Chest roentgenography is normal.

A WORRISOME PICTURE

Which of the following is associated with left bundle branch block?

- Myocardial injury
- Hypertension
- Aortic stenosis
- Intrinsic conduction system disease
- All of the above

All of the above are true. For left bundle branch block to be diagnosed, the rhythm must be supraventricular and the QRS duration must be 120 ms or more. There should be a QS or RS complex in V₁ and a monophasic R wave in I and V₆. Also, the T wave should be deflected opposite the terminal deflection of the QRS complex. This is known as appropriate T-wave discordance with bundle branch block. A concordant T wave is nonspecific but suggests ischemia or myocardial infarction.
Potential causes of a new left bundle branch block include hypertension, acute myocardial infarction, aortic stenosis, and conduction system disease. A new left bundle branch block with a concomitant decrease in ejection fraction, especially in a patient with cardiac risk factors, is very worrisome, raising the possibility of ischemic heart disease.

A gated rubidium-82 dipyridamole stress test yields no evidence of a fixed or reversible perfusion defect.

Left heart catheterization reveals angiographically normal coronary arteries.

Magnetic resonance imaging (MRI) shows a moderately hypertrophied left ventricle with moderately to severely depressed systolic function (left ventricular ejection fraction 27%). The left ventricle appears dysynchronous. Delayed-enhancement imaging reveals patchy delayed enhancement in the basal septum and the basal inferolateral walls. These findings suggest cardiac sarcoidosis, with a sensitivity of nearly 100% and a specificity of approximately 78%.

Sarcoidosis is a multisystem disease characterized by noncaseating granulomas. Almost any organ can be affected, but it most commonly involves the respiratory and lymphatic systems. Although infectious, environmental, and genetic factors have been implicated, the cause remains unknown. The prevalence is approximately 20 per 100,000, being higher in black and Japanese populations.

Cardiac involvement is seen in 20% to 30% of patients with sarcoidosis. However, most cases are subclinical, and symptomatic cardiac involvement is rare.

### MORE CARDIAC TESTING

The patient undergoes more cardiac testing.

Transthoracic echocardiography is done again. The left ventricle is normal in size, but the ejection fraction is 35%. In addition, stage 1 diastolic dysfunction (abnormal relaxation) and evidence of mechanical dyssynchrony (disruption in the normal sequence of activation and contraction of segments of the left ventricular wall) are seen. The right ventricle is normal in size and function. There is trivial mitral regurgitation and mild tricuspid regurgitation with normal right-sided pressures.

**TABLE 1**

<table>
<thead>
<tr>
<th>STUDY</th>
<th>VALUE</th>
<th>REFERENCE RANGE</th>
</tr>
</thead>
<tbody>
<tr>
<td>White blood cell count</td>
<td>5.83</td>
<td>4.0–11.0 k/uL</td>
</tr>
<tr>
<td>Hemoglobin</td>
<td>13.0</td>
<td>13.5–17.5 g/dL</td>
</tr>
<tr>
<td>Hematocrit</td>
<td>36.8</td>
<td>40%–52%</td>
</tr>
<tr>
<td>Mean corpuscular volume</td>
<td>89.3</td>
<td>80–100 fl</td>
</tr>
<tr>
<td>Platelet count</td>
<td>257</td>
<td>150–400 k/uL</td>
</tr>
<tr>
<td>Protein, total</td>
<td>5.9</td>
<td>6.0–8.4 g/dL</td>
</tr>
<tr>
<td>Albumin</td>
<td>3.3</td>
<td>3.5–5.0 g/dL</td>
</tr>
<tr>
<td>Calcium</td>
<td>8.6</td>
<td>8.5–10.5 mg/dL</td>
</tr>
<tr>
<td>Bilirubin, total</td>
<td>0.7</td>
<td>0.0–1.5 mg/dL</td>
</tr>
<tr>
<td>Alkaline phosphatase</td>
<td>557</td>
<td>40–150 U/L</td>
</tr>
<tr>
<td>Aspartate aminotransferase</td>
<td>59</td>
<td>7–40 U/L</td>
</tr>
<tr>
<td>Glucose</td>
<td>95</td>
<td>65–100 mg/dL</td>
</tr>
<tr>
<td>Blood urea nitrogen</td>
<td>31</td>
<td>10–25 mg/dL</td>
</tr>
<tr>
<td>Creatinine</td>
<td>1.64</td>
<td>0.70–1.40 mg/dL</td>
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<tr>
<td>Sodium</td>
<td>138</td>
<td>135–146 mmol/L</td>
</tr>
<tr>
<td>Potassium</td>
<td>4.6</td>
<td>3.5–5.0 mmol/L</td>
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<tr>
<td>Chloride</td>
<td>105</td>
<td>98–110 mmol/L</td>
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<tr>
<td>CO₂</td>
<td>25</td>
<td>23–32 mmol/L</td>
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<tr>
<td>Anion gap</td>
<td>8</td>
<td>0–15 mmol/L</td>
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<tr>
<td>Alanine aminotransferase</td>
<td>47</td>
<td>5–50 U/L</td>
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<tr>
<td>Thyroid-stimulating hormone</td>
<td>1.870</td>
<td>0.5–5.0 mIU/L</td>
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<tr>
<td>Troponin T</td>
<td>&lt; 0.01</td>
<td>0.00–0.10 ng/mL</td>
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<tr>
<td>Creatine kinase</td>
<td>83</td>
<td>30–220 U/L</td>
</tr>
<tr>
<td>Creatine kinase MB</td>
<td>3.2</td>
<td>0.0–8.8 ng/mL</td>
</tr>
<tr>
<td>B-type natriuretic peptide</td>
<td>483</td>
<td>&lt; 100 pg/mL</td>
</tr>
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</table>
involvement is present in only about 5% of patients with systemic sarcoidosis. Isolated cardiac sarcoidosis has been described in case reports but is rare.

The clinical manifestations of cardiac sarcoidosis depend on the location and extent of granulomatous inflammation. In a necropsy study of 113 patients with cardiac sarcoidosis, the left ventricular free wall was the most common location, followed by the interventricular septum.

How does cardiac sarcoidosis most commonly present?

- Conduction abnormalities
- Ventricular tachycardia
- Cardiomyopathy
- Sudden death
- None of the above

Common presentations of cardiac sarcoidosis include conduction system disease and arrhythmias (which can sometimes result in sudden death), and heart failure.

Conduction abnormalities due to granulomas (in the active phase of sarcoidosis) and fibrosis (in the fibrotic phase) in the atrioventricular node or bundle of His are the most common presentation of cardiac sarcoidosis. These lesions may result in relatively benign first-degree heart block or may be as potentially devastating as complete heart block.

In almost all patients with conduction abnormalities, the basal interventricular septum is involved. Patients who develop complete heart block from sarcoidosis tend to be younger than those with idiopathic heart block. Therefore, complete heart block in a young patient should raise the possibility of this diagnosis.

Ventricular tachycardia (sustained or non-sustained) and ventricular premature beats are the second most common presentation. Up to 22% of patients with sarcoidosis have electrocardiographic evidence of ventricular arrhythmias. The cause is believed to be myocardial scar tissue resulting from the sarcoid granulomas, leading to electrical reentry. Sudden death due to ventricular tachyarrhythmias or conduction blocks accounts for 25% to 65% of deaths from cardiac sarcoidosis.

Heart failure may result from sarcoidosis when there is extensive granulomatous disease.
in the myocardium. Depending on the location of the granulomas, both systolic and diastolic dysfunction can occur. In severe cases, extensive granulomas can cause left ventricular aneurysms.

The diagnosis of cardiac sarcoidosis as the cause of heart failure can be difficult to establish, especially in patients without evidence of sarcoidosis elsewhere. Such patients are often given a diagnosis of idiopathic dilated cardiomyopathy. However, compared with patients with idiopathic dilated cardiomyopathy, those with cardiac sarcoidosis have a greater incidence of advanced atrioventricular block, abnormal wall thickness, focal wall motion abnormalities, and perfusion defects of the anteroseptal and apical regions.17

Progressive heart failure is the second most frequent cause of death (after sudden death) and accounts for 25% to 75% of sarcoid-related cardiac deaths.9,18,19

### How is cardiac sarcoidosis diagnosed?

Electrocardiography
Echocardiography
Computed tomography
Endomyocardial biopsy
There are no guidelines for diagnosis

Given the variable extent and location of granulomas in sarcoidosis, the diagnosis of cardiac sarcoidosis is often challenging.

To make the diagnosis of sarcoidosis in general, the American Thoracic Society2 says that the clinical picture should be compatible with this diagnosis, noncaseating granulomas should be histologically confirmed, and other diseases capable of producing a similar clinical or histologic picture should be excluded.

A newer diagnostic tool, the Sarcoidosis Three-Dimensional Assessment Instrument,20 incorporates two earlier tools.20,21 It assesses three axes: organ involvement, sarcoidosis severity, and sarcoidosis activity and categorizes the diagnosis of sarcoidosis as “definite,” “probable,” or “possible.”20

In Japan, where sarcoidosis is more common, the Ministry of Health and Welfare22 says that cardiac sarcoidosis can be diagnosed histologically if operative or endomyocardial biopsy specimens contain noncaseating granuloma. In addition, the diagnosis can be suspected in patients who have a histologic diagnosis of extracardiac sarcoidosis if the first item in the list below and one or more of the rest are present:

- Complete right bundle branch block, left axis deviation, atrioventricular block, ventricular tachycardia, premature ventricular contractions (> grade 2 of the Lown classification), or Q or ST-T wave abnormalities
- Abnormal wall motion, regional wall thinning, or dilation of the left ventricle on echocardiography
- Perfusion defects on thallium 201 myocardial scintigraphy or abnormal accumulation of gallium citrate Ga 67 or technetium 99m on myocardial scintigraphy
- Abnormal intracardiac pressure, low cardiac output, or abnormal wall motion or depressed left ventricular ejection fraction on cardiac catheterization
- Nonspecific interstitial fibrosis or cellular infiltration on myocardial biopsy.

The current diagnostic guidelines from the American Thoracic Society2 and the Japanese Ministry of Health and Welfare22 and the Sarcoidosis Three-Dimensional Assessment Instrument,20 however, do not incorporate newer imaging studies as part of their criteria.

### A DEFINITIVE DIAGNOSIS

Endomyocardial biopsy often provides the definitive diagnosis of cardiac sarcoidosis.

True
False

False. Endomyocardial biopsy often fails to reveal noncaseating granulomas, which have a patchy distribution.13 TABLE 2 summarizes the accuracy of tests for cardiac sarcoidosis.

**Electrocardiography** is abnormal in up to 50% of patients with sarcoidosis,23 reflecting the conduction disease or arrhythmias commonly seen in cardiac involvement.

**Chest radiography** classically shows hilar lymphadenopathy and interstitial disease, and may show cardiomegaly, pericardial effusion, or left ventricular aneurysm.
Echocardiography is nonspecific for sarcoid disease, but granulomatous involvement and scar tissue of cardiac tissue may appear hyperechogenic, particularly in the ventricular septum or left ventricular free wall.24

Angiography. Primary sarcoidosis rarely involves the coronary arteries,25 so angiography is most useful in excluding the diagnosis of atherosclerotic coronary artery disease.

Radionuclide imaging with thallium 201 in patients with suspected cardiac sarcoidosis may be useful to suggest myocardial involvement and to exclude cardiac dysfunction secondary to coronary artery disease. Segmental areas with defective thallium 201 uptake correspond to fibrogranulomatous tissue. In resting images, the pattern may be similar to that seen in coronary artery disease. However, during exercise, perfusion defects increase in patients who have ischemia but actually decrease in those with cardiac sarcoidosis.26

Nevertheless, some conclude that thallium scanning is too nonspecific for it to be used as a diagnostic or screening test.27,28 The combined use of thallium 201 and gallium 67 may better detect cardiac sarcoidosis, as gallium is taken up in areas of active inflammation.

Positron-emission tomography (PET) with fluorodeoxyglucose F 18 (FDG), with the patient fasting, appears to be useful in detecting the early inflammation of cardiac sarcoidosis29–34 and monitoring disease activity.30,31 FDG is a glucose analogue that is taken up by granulomatous tissue in the myocardium.34 The uptake in cardiac sarcoidosis is in a focal distribution.30,31,34 The abnormal FDG uptake resolves with steroid treatment.31,32

MRI has promise for diagnosing cardiac sarcoidosis. With gadolinium contrast, MRI has superior image resolution and can detect cardiac involvement early in its course.27,29,35–44

Inflammation of the myocardium due to sarcoid involvement appears as focal zones of increased signal intensity on both T2-weighted and early gadolinium T1-weighted images. Late myocardial enhancement after gadolinium infusion is the most typical finding of cardiac sarcoidosis on MRI, and likely represents fibrogranulomatous tissue.27 Delayed gadolinium enhancement is also seen in myocardial infarction but differs in its distribution.1,13,45

Cardiac sarcoidosis most commonly affects the basal and lateral segments. In one study, the finding of delayed enhancement had a sensitivity of 100% and a specificity of 78%,1,27 though it may not sufficiently differentiate active inflammation from scar.30

Like FDG-PET, MRI has also been shown to be useful for monitoring treatment.33,46 However, PET is more useful for follow-up in patients who receive a pacemaker or implantable cardioverter-defibrillator, in whom MRI is contraindicated. One case report29 described using both delayed-enhancement MRI and FDG-PET to diagnose cardiac sarcoidosis.

TREATMENT

How is cardiac sarcoidosis currently treated?

- Implantable cardioverter-defibrillator
- Corticosteroids
- Heart transplantation
- All of the above
- None of the above

Corticosteroids

Corticosteroids are the mainstay of treatment of cardiac sarcoidosis, as they attenuate the characteristic inflammation and fibrosis of sarcoid granulomas. The goal is to prevent compromise of cardiac structure or function.47 Although most of the supporting data are anecdotal, steroids have been shown to improve ventricular contractility,38 arrhythmias,49 and conduction abnormalities.50 MRI and FDG-
The patient’s cardiac MRI

**FIGURE 2.** Magnetic resonance imaging of the patient’s heart. The long-axis phase-sensitive image shows delayed enhancement in the basal septum and basal inferolateral walls (arrows), strongly suggesting sarcoidosis. End-systolic and end-diastolic steady-state free precession images in the same plane show a moderately hypertrophied but contractile left ventricle, which argues against ischemia.

PET studies have shown cardiac lesions resolving after steroids were started.31,45,46

The optimal dosage remains unknown. Initial doses of 30 to 60 mg daily, gradually tapered over 6 to 12 months to maintenance doses of 5 to 10 mg daily, have been effective.45,51

Relapses are common and require vigilant monitoring.

Alternative agents such as cyclophosphamide (Cytoxan),52 methotrexate (Rheumatrex),53 and cyclosporine (Sandimmune)54 can be given to patients whose disease does not respond to corticosteroids or who cannot tolerate their side effects.

**Implantable cardioverter-defibrillator**

Sudden death due to ventricular tachyarrhythmias or conduction block accounts for 30% to 65% of deaths in patients with cardiac sarcoidosis.10 The rates of recurrent ventricular tachycardia and sudden death are high, even with antiarrhythmic drug therapy.55

Although experience with implantable cardiac defibrillators is limited in patients with cardiac sarcoidosis,55-58 some have argued that they be strongly considered to prevent sudden cardiac death in this high-risk group.57,58

**Heart transplantation**

The largest body of data on transplantation comes from the United Network for Organ Sharing database. In the 65 patients with cardiac sarcoidosis who underwent cardiac transplantation in the 18 years from October 1987 to September 2005, the 1-year post-transplant survival rate was 88%, which was better than in patients with all other diagnoses (85%). The 5-year survival rate was 80%,59,60

Recurrence of sarcoidosis within the cardiac allograft and transmission of sarcoidosis from donor to recipient have both been documented after heart transplantation.61,62
CAUSES OF DEATH

What is the most common cause of death in patients with cardiac sarcoidosis?

☐ Respiratory failure
☐ Conduction disturbances
☐ Progressive heart failure
☐ Ventricular tachyarrhythmias
☐ None of the above

The prognosis of symptomatic cardiac sarcoidosis is not well defined, owing to the variable extent and severity of the disease. The mortality rate in sarcoidosis without cardiac involvement is about 1% to 5% per year.63,64 Cardiac involvement portends a worse prognosis, with a 55% survival rate at 5 years and 44% at 10 years.17,65 Most patients in the reported series ultimately died of cardiac complications of sarcoidosis, including ventricular tachyarrhythmias, conduction disturbances, and progressive cardiomyopathy.10,17

Since corticosteroids, pacemakers, and implantable cardioverter-defibrillators have begun to be used, the cause of death has shifted from sudden death to progressive heart failure.66

REFERENCES


CASE CONTINUED

While hospitalized, our patient had two episodes of nonsustained ventricular tachycardia (7 and 12 beats) on telemetry. Cardiac MRI showed a lesion in the basal septum most likely involving the left bundle and an area of lateral basilar involvement near the mitral annulus (FIGURE 2). Ventricular dys synchrony was clearly evident on both echocardiography and MRI, with depressed left ventricular function (ejection fraction 28% on MRI, 35% on echocardiography).

Electrophysiologic testing revealed inducible monomorphic sustained ventricular tachycardia. The patient subsequently had a biventricular cardioverter-defibrillator implanted. He was started on an angiotensin-converting enzyme inhibitor and a beta-blocker for his heart failure. Further imaging of his chest and abdomen revealed lesions in his thyroid and liver. As of this writing, he is undergoing further workup. Because of active infection with Clostridium difficile, steroid therapy was deferred.