INTRODUCTION

Neurologists and cardiologists are frequently involved in coordinating the care of patients with a variety of conditions, the most common being stroke. This article reviews the diagnosis and management of conditions that have both cardiac and
neurologic manifestations. The review is organized by cardiac disorders that predispose to neurological complications and neurological conditions that can involve the heart.

**NEUROLOGICAL COMPLICATIONS OF CARDIAC DISEASE**

Cardiac diseases can be complicated by stroke, cognitive impairment, and brain infections. Cardiac conditions including atrial fibrillation (AF), cardiomyopathies, valvular heart disease and interatrial septal anomalies account for 20%-30% of all ischemic strokes. The mechanism of cardiogenic stroke is often embolic, but hypoperfusion may also occur, particularly in those with cerebral steno-occlusive disease. Cognitive decline can be associated with congestive heart failure (CHF) and coronary artery bypass graft procedures, whereas meningitis and brain abscesses are possible complications of infective endocarditis (IE).

**Atrial Fibrillation**

Nonvalvular AF is the most common cause of cardiogenic stroke, accounting for 50% of cardiogenic emboli and 10% of ischemic strokes. It affects about 1% of the general population and is the most common cardiac arrhythmia in the elderly, having a prevalence of nearly 10% in those older than 80 years. AF is associated with a 4- to 5-fold increased risk of stroke across all age groups; those who also had a prior stroke or transient ischemic attack (TIA) have an additional 2.5-fold increase in stroke risk.1,2 The mechanism of cardiogenic stroke is often embolic, but hypoperfusion may also occur, particularly in those with cerebral steno-occlusive disease. Cognitive decline can be associated with congestive heart failure (CHF) and coronary artery bypass graft procedures, whereas meningitis and brain abscesses are possible complications of infective endocarditis (IE).

### Abbreviations
- ASA: atrial septal aneurysm
- CHF: congestive heart failure
- ECG: electrocardiogram
- IE: infective endocarditis
- MD: muscular dystrophy
- MI: myocardial infarction
- PFO: patent foramen ovale
- TIA: transient ischemic attack
general population has a PFO. Patients with a PFO may also have an atrial septal aneurysm (ASA), defined as >10 mm septal excursion with a base of 1.5 cm or greater. Potential mechanisms of stroke in patients with PFO with or without an ASA include: a) paradoxical embolization; b) embolization of thrombus from the rim of the ASA; c) paroxysmal AF related to damage to the cardiac conduction system; d) passage of unmeasured vasoactive humoral substances that escape pulmonary degradation; and/or e) causes not related to the defect. A potential source of paradoxical embolization is identified in 10% to 57% of patients with PFO and cryptogenic strokes.

A meta-analysis of case-control studies established a higher prevalence of PFO and ASA in stroke patients overall and in those who had cryptogenic strokes. The combination was more frequent in those with stroke compared to nonstroke control subjects (odds ratio [OR] = 5.25; 95% CI, 2.91-9.45), in those with cryptogenic stroke versus known stroke cause (OR=20.74; 95% CI, 4.14-103.90), and cryptogenic stroke versus nonstroke controls (OR=23.9; 95% CI, 3.09-185.42). The association of PFO/ASA abnormalities with stroke was higher in patients who were younger than 55 years. These studies, however, are largely based on hospitalized patients, possibly reflecting a referral bias. Population-based studies do not find such a relationship. For example, PFO was present in 140 of 585 (24.3%) and ASA in 11 (1.9%) randomly sampled subjects older than 45 years from Olmstead County, Minnesota. The combination of PFO and ASA was present in 6 (4.3%) and was not associated with an increased stroke risk over 5 years.

Despite this uncertainty, clinical trials have been conducted or are in progress evaluating treatments of patients with stroke in the setting of PFO with or without an ASA. The Patent Foramen Ovale in Cryptogenic Stroke Study (PICSS) was a substudy of the Warfarin-Aspirin Recurrent Stroke Study (WARSS) that randomized patients with otherwise cryptogenic stroke to acetylsalicylic acid as compared to warfarin. There was no difference in the rate of stroke or death between those with cryptogenic stroke based on the presence of a PFO and no overall difference in the effect of acetylsalicylic acid versus warfarin on recurrent stroke risk. The results of the prospective Spanish multicenter study (CODICIA) were similar. In contrast, the Patent Foramen Ovale and Atrial Septal Aneurysm Study found that the combination of PFO and ASA was associated with an increased risk of recurrent strokes (HR 4.17; 95% CI, 1.47-11.84) among patients aged 18 to 55 years with cryptogenic stroke who were receiving acetylsalicylic acid. The differing results may be due to differences in patient populations (younger cryptogenic stroke patients and only those taking acetylsalicylic acid in the Patent Foramen Ovale and Atrial Septal Aneurysm Study). Currently, there are insufficient data to recommend anticoagulation over acetylsalicylic acid for secondary stroke prevention in patients with PFO or PFO/ASA, and either choice may be reasonable.

The role of endovascular PFO closure for secondary stroke prevention is also uncertain. An open-treatment nonrandomized study comparing percutaneous PFO closure to medical therapy (Vitamin K antagonist or antiplatelet therapy) in patients with cryptogenic stroke did not find an overall benefit of PFO closure after 4 years. An initial report of the CLOSURE 1 trial, comparing endovascular PFO closure with medical therapy in patients with stroke or TIA, found no benefit from PFO closure, which was associated with several periprocedural complications (presented at AHA Scientific Sessions, November 2010). Other randomized trials of PFO closure for secondary stroke prevention are in progress.

**Acute Myocardial Infarction and Left Ventricular Thrombus**

There is a short- and long-term increased risk of stroke following MI. Although the exact cause of MI-associated stroke is often unclear, embolization from left ventricular thrombus is a possible mechanism. Left ventricular thrombus complicating acute myocardial infarction (AMI) results from turbulent blood flow and stasis related to an akinetic left ventricular wall segment or aneurysm. The incidence of left ventricular thrombus was 6.2% in a study of 642 patients with anterior wall AMI followed for 6 years, and did not differ between patients treated conservatively, with thrombolysis, or with a primary percutaneous intervention. Predictors of left ventricular thrombus were low ejection fraction and severe mitral regurgitation.

The incidence of stroke in the acute phase following MI is approximately 1%. Risk factors include large MI, anterior wall involvement, prior stroke, and increasing age. The incidence of stroke was 0.7% at 30 days in patients with AMI without persistent ST-segment elevation in the Platelet Glycoprotein IIb/IIIa in Unstable Angina: Receptor Suppression Using Integrilin Therapy (PURSUIT) trial that enrolled 10 948 patients; 84% of all strokes were ischemic. Strokes occurring after MI resulted in high morbidity and mortality and were predicted by a higher baseline heart rate. The long-term risk of stroke following MI is about 6%. Strokes are mainly ischemic. Risk factors include advancing age, diabetes mellitus, previous history of stroke, history of hypertension, and smoking.

The risk of post-MI stroke may be highest over the first 3 months. Some data suggest that treatment with warfarin (target INR, 3.0-4.0) or the combination of acetylsalicylic acid and warfarin (INR, 2.0-2.5) may be more efficacious than acetylsalicylic acid alone in preventing recurrent cardiovascular events after MI, including strokes. Warfarin, however, is not generally recommended after MI for stroke prevention unless a stroke or TIA occurs in association with a left ventricular mural thrombus. In this setting, oral anticoagulation with a target INR of 2.5 for at least 3 months is suggested. The American College of Chest Physicians guidelines recommend considering moderate intensity oral anticoagulation (INR, 2.0-3.0) in addition to low-dose acetylsalicylic acid (≤100 mg/d) for high-risk post-MI patients, including those with a large anterior MI, significant heart failure, intracardiac thrombus, AF, or a history of a thromboembolic event.

**Congestive Heart Failure**

Congestive heart failure can lead to cerebral embolism and hypoperfusion-related ischemia, causing both stroke and cognitive impairments. The risks and mechanisms of cognitive decline with CHF are less well understood than the risk for stroke. Severe CHF can be associated with impaired alertness, behavioral change and cognitive impairment similar to metabolic encephalopathy. There is not a consistent improvement in neuropsychological functioning in patients with CHF-associated cognitive impairments after cardiac transplantation. Accounting for 9% of strokes, CHF is second after AF as a cause of cardiogenic stroke and CHF and AF coexist in about 2% of stroke patients. In community studies, CHF is associated with more than a 2-fold increased stroke risk, but much less in clinical trials. Some studies suggest that the risk may increase with decreasing ejection fraction.

The optimal strategy for decreasing stroke risk in the CHF setting is uncertain. Two large prospective, randomized, controlled studies assessed whether warfarin is superior to acetylsalicylic acid
in patients with a low ejection fraction (ie, <35%). The Warfarin and Antiplatelet Therapy in Chronic Heart Failure (WATCH) trial compared warfarin to acetylsalicylic acid and clopidogrel to acetylsalicylic acid.54,55 Although stopped early due to poor enrollment, WATCH did not find a difference in the composite endpoint of death, MI or nonfatal stroke, but did find that nonfatal stroke, a secondary outcome, was lower in patients receiving warfarin compared to those receiving acetylsalicylic acid or clopidogrel.56 The Warfarin versus Aspirin in Patients With Reduced Cardiac Ejection Fraction (WARCEF) trial is ongoing and compares warfarin (INR, 2.5-3) versus acetylsalicylic acid (325 mg).57 The WATCH and WARCEF investigators have planned a combined analysis. Given the low overall stroke rate in CHF, it may be important to develop a risk stratification scheme similar to the CHADS2 score to identify those at highest stroke risk in order to guide future treatments and enrich clinical trials.50

**Native Valvular Heart Disease and Prosthetic Heart Valves**

Aortic and mitral valve diseases are associated with an increased stroke risk. A population based study found that patients with mitral stenosis, mitral regurgitation, aortic stenosis, and aortic regurgitation had a higher than expected number of cerebrovascular events. Predictors included advancing age, AF, and severe aortic stenosis.58 Mitral valve prolapse and mitral annular calcification have also been associated with increased stroke risk in some studies.59–61 Mitral annular calcification, an uncommon nonrheumatic cause of mitral stenosis predominantly occurring in women, may be associated with stroke because of concomitant aortic atheromas and the risk of embolization of fibrocalcific material. The stroke risk in patients with mitral annular calcification increases with age.2,62

Antithrombotic medications reduce the risk of stroke and systemic embolism in patients with native valvular heart disease, mechanical and bioprosthetic valves, but their use is associated with increased bleeding risk.2 The AHA/American Stroke Association recommends long-term warfarin (INR target, 2.5) in patients with rheumatic mitral valve disease based on the high risk of stroke recurrence, because mitral valvuloplasty does not eliminate embolic risk, and because observational studies indicate that treatment is associated with a lower risk of recurrent embolism.63–65 In the absence of randomized trials, long-term antiplatelet therapy may be considered in patients with native aortic or nonrheumatic mitral valve disease who have ischemic stroke or TIA.2 For secondary stroke prevention in patients with mechanical prosthetic valves, warfarin (INR target, 3.0; range 2.5–3.5) is recommended over antiplatelet agents.2,66 Acetylsalicylic acid (75 to 100 mg/day) may be added to warfarin in patients who have strokes despite adequate anticoagulation and who are not at high risk for bleeding.2,67

**Infective Endocarditis**

Stroke, mycotic aneurysms, meningitis, and intracerebral abscesses can complicate IE.58 Approximately two-thirds of embolic events in patients with endocarditis involve the central nervous system; 20–40% of left-sided IE cases are complicated by stroke, most commonly in the middle cerebral artery distribution.66,67 Stroke can also be the initial manifestation of IE.15 Intracerebral hemorrhage (ICH) occurs in about 5% of IE due to hemorrhagic infarction or mycotic aneurysmal rupture and is associated with more than 50% mortality.70 Clinically silent cerebral embolism occurs in 30% of patients with mitral or aortic valve endocarditis.71 A multicenter prospective incidence cohort study of 1437 patients with left-sided IE reported a stroke incident rate of 4.82/1000 patient-days over the first week of therapy that fell to 1.71/1000 patient-days in the second week, with a further decline with continued antimicrobial therapy. Stroke risk factors included infection with *Staphylococcus aureus*, mitral valve vegetations, and myocardial abscess. Other risk factors include mobile vegetations or vegetations > 10 mm in diameter occurring on the anterior mitral leaflet.70,72,73

Surgery to prevent further embolic complications remains controversial because stroke risk falls dramatically with effective antimicrobial therapy.70 Moreover, the safety of cardiopulmonary bypass following stroke is uncertain due to concerns for intracranial hemorrhage related to anticoagulation during the procedure and potential hemodynamic worsening of the ischemic infarction. Despite these concerns, surgery may still be considered if performed early (ie, within 72 h) when embolic rates are highest and when there are other predictors of poor outcome (eg, recurrent embolization, CHF, prosthetic valve IE).74,75 Furthermore, the outcome of stroke patients following cardio-pulmonary bypass may be better than previously thought, with the exception of those with ICH.70,71,76 Large prospective studies quantifying the risk of cardiac surgery after IE-related ICH are lacking, but delaying for 4 weeks before cardiac surgery is often recommended.77 The AHA recommends stopping anticoagulation for at least the first 2 weeks of antibiotic therapy in the case of recent embolic stroke in the setting of *Staphylococcus aureus* prosthetic valve IE.74

**Neurologic Complications of Cardiac Interventions**

Stroke can complicate many cardiac procedures, including percutaneous coronary intervention (PCI), coronary artery bypass operations, valvuloplasty, and catheter ablation for AF. Stroke infrequently complicates PCI (about 0.3%), with the risk increasing with age, advanced renal failure, diabetes mellitus, hypertension, previous stroke, and emergency use of an intra-aortic balloon pump.78,79 Stroke complicating PCI portends high in-hospital and 1-year mortality.78,79

There are no established guidelines for stroke prevention in the setting of cardiac interventions. Furthermore, the management of stroke in these settings is not straightforward. For example, in the case of stroke complicating catheterization, intravenous tissue plasminogen activator (IV-tPA) has been contraindicated in patients with MI within the 3 months prior to stroke, previously excluding many of these patients from treatment; recent MI is, however, no longer an absolute contraindication for IV-tPA.80 Other concerns include tPA-related bleeding due to the invasive procedure, concurrent use of antithrombotics, and conjectures about the composition of clots that may be less amenable to lysis.81 There is, however, some evidence that thrombolysis, whether intra-arterial or intravenous, might improve early outcomes after percutaneous coronary intervention strokes.81 Coronary artery surgery, especially conventional on-pump coronary artery bypass graft, carries a higher rate of stroke and cerebral dysfunction than PCI.5,82 Because of the bleeding risk, IV-tPA is contraindicated for periprocedural strokes. Case series suggest reasonable safety and efficacy for intra-arterial thrombolytic therapy; however, adequate controlled studies with appropriate follow-up are lacking.82-84 Although coronary artery bypass graft has been associated with an increased risk of cognitive impairments, studies comparing the cognitive status of patients after coronary artery bypass graft with an appropriate control group of otherwise similar patients with
cardiac disease not undergoing coronary artery bypass graft do not find greater decline after the operation.85-87 Furthermore, it does not seem to be a sustainable difference in cognitive function in patients who had on-pump versus off-pump coronary artery bypass graft.88

CARDIOVASCULAR DISEASE

Cerebrovascular Disease

Just as stroke may complicate a variety of cardiac disorders, cerebrovascular disease can affect the heart. Electrocardiographic (ECG) changes suggesting MI occur in patients with intracerebral hemorrhage, subarachnoid hemorrhage, and 15% to 40% of those with ischemic stroke.89 Stroke can cause large inverted T-waves in anterior ECG leads and a variety of cardiac arrhythmias. Severe strokes can cause myocardial cytolysis. Monitoring for cardiac arrhythmias and tests for myocardial injury are part of the standard evaluation of stroke patients.83

Neuromuscular Disorders

Cardiac disorders accompany a wide variety of neuromuscular diseases (Table 1).89-92 For example, left ventricular hypertrophy (LVHT), a cardiac disorder characterized by prominent trabeculations and intertrabecular recesses within the left ventricle, is associated with neuromuscular disorders in up to 80% of cases.93 Cardiac involvement is frequent in mitochondrial disorders and can be the primary determinant of prognosis. Cardiomyopathy may also be the presenting and main feature of Becker muscular dystrophy (BMD) and Emery-Dreifuss muscular dystrophy (EDMD).94 Depending on the underlying neuromuscular disorder, management of cardiac involvement should be aimed at determining the degree and type of abnormalities, preventing complications and alleviating symptoms.90

Mitochondrial Disorders

Mitochondrial disorders result either from mutations affecting mitochondrial DNA encoding protein subunits of the respiratory chain complexes or from mitochondrial defects originating from nuclear DNA.95,96 Cardiac involvement usually manifests in childhood or early adulthood, can precede the diagnosis of mitochondrial disease, and is associated with increased mortality.95 Cardiac involvement varies between different mitochondrial disorders and can present as hypertrophic cardiomyopathy, dilated cardiomyopathy, and conduction abnormalities. Symptoms include CHF, syncope, and sudden death.96

For example, Kearns-Sayre syndrome is characterized by atrioventricular conduction abnormalities that can progress rapidly to complete atrioventricular block, often warranting prophylactic pacemaker implantation.89,97 In contrast, myoclonus epilepsy and ragged red fibers (MERRF) and mitochondrial myopathy, encephalopathy, lactic acidosis, and stroke-like episodes (MELAS) are characterized by left ventricular hypertrophy and dilated cardiomyopathy, with asymmetric septal hypertrophy further characterizing MERRF.90 Left ventricular hypertrophy is also the main cardiac manifestation of Friederich ataxia.90

The diagnosis can often be made by mitochondrial enzymatic analysis in skeletal or cardiac muscles biopsies.96 Regardless of cardiac symptoms, cardiac evaluation is recommended for patients with suspected mitochondrial disorders because of the strong association with cardiac disorders and risk for sudden death.

Muscular Dystrophies

Duchenne and Becker muscular dystrophies. Duchenne muscular dystrophy (DMD) and BMD result from X-linked mutations in the dystrophin gene on chromosome Xp21. They initially present with proximal weakness that gradually progresses to involve other muscles, including respiratory muscles.94 The dystrophin protein is located on the inner surface of the sarcolemma. It is completely absent or dysfunctional in DMD, but only reduced in BMD.94 Children with DMD typically present with muscle weakness and difficulty reaching motor milestones by the age of 5, whereas patients with BMD may present later and have less muscle weakness. Respiratory and cardiac failures are the leading causes of death in DMD, which usually occurs between the second and third decade.94 In contrast, patients with BMD have a slower progression of weakness and typically survive into adulthood, but heart failure remains an important cause of death and can be the presenting symptom that brings the patient to medical attention.98 Both DMD and BMD female carriers may develop dilated cardiomyopathy without apparent weakness.98 Cardiomyopathy in DMD and BMD is thought to be due to absent or diminished dystrophin in cardiomyocytes.99 Dystrophin deficiency may further increase the susceptibility to myocarditis that may play an additional role in the progression to heart failure.100 In DMD, focal areas of myocardial injury may appear as areas of regional wall abnormalities on cardiac imaging.95 Symptoms of cardiac involvement can be masked by reduced physical activity due to skeletal muscle weakness.99 Left ventricular dysfunction with predilection for wall motion defects in the posterobasal and posterolateral segments can be diagnosed with echocardiography and cardiac magnetic resonance imaging.99,101 Mitral regurgitation can also develop as a result of papillary muscle dysfunction.99 Arrhythmias, including sinus node dysfunction, atrioventricular node dysfunction, AF, and ventricular tachycardia/fibrillation may develop later in the course of the disease.99 Echocardiography and ECG is recommended starting at age 10 years in children with DMD.99,102 Treatment of cardiomyopathy includes β blockers, angiotensin-converting enzyme inhibitors, and management of cardiac arrhythmias.99 The angiotensin-converting enzyme inhibitor perindopril preserves systolic function in patients with DMD and decreases mortality.103,104

Emery Dreifuss muscular dystrophy. EDMD can be X-linked or autosomal. The X-linked form is caused by mutations of the STA gene at chromosome Xq28 encoding emerin, a nuclear membrane protein. The autosomal type is caused by a mutation of the LMNA gene at chromosome 1q21 encoding the nuclear lamina proteins, lamin A and C.105 The LMNA mutations cause other pathologies such as dilated cardiomyopathy and conduction system disease, limb-girdle muscular dystrophy with arrhythmogenic conduction disturbances, an autosomal form of axonal neuropathy, and a rare form of familial partial lipodystrophy.105 Early elbow, ankle and cervical spine contractures characterize EDMD.105 Proximal upper limb and distal lower limb weakness is usually slowly progressive, with later involvement of the proximal limb-girdle muscles.105 Cardiac involvement is characterized by conduction defects, usually evident by age 30 years and may present without muscle weakness.105,106 X-linked EDMD is characterized by AF/flutter, bradyarrhythmia requiring pacemakers, prolongation of PR interval, complete heart block and atrial standstill. Absent P waves due to atrial paralysis is a typical ECG finding.94 Embolic strokes can result from cardiac arrhythmias.105,107 Heart failure occurs mainly in the autosomal forms of EDMD, with symptoms appearing in the...
third or fourth decades of life. Holter monitoring may help early detection of arrhythmias in this population and an ECG should be obtained annually. A pacemaker is recommended after sinus node or atrioventricular conduction is apparent. Sudden death may still occur after pacemaker placement, possibly due to ventricular tachycardia or fibrillation. An intracardiac defibrillator may be warranted in those with documented symptomatic ventricular arrhythmias.

### Table 1

<table>
<thead>
<tr>
<th>Neuromuscular disorders</th>
<th>Mode of inheritance</th>
<th>Clinical manifestations</th>
<th>Cardiac manifestations</th>
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<tbody>
<tr>
<td><strong>Mitochondrial disorders</strong></td>
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<tr>
<td>Kearns-Sayre</td>
<td>Mitochondrial</td>
<td>Progressive external ophthalmoplegia, pigmentary degeneration of the retina, short stature, cerebellar ataxia, dementia</td>
<td>AV conduction abnormalities</td>
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<td></td>
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<td>AV conduction block</td>
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<tr>
<td>MELAS</td>
<td>Mitochondrial</td>
<td>Stroke like episodes, lactic acidosis, seizures, dementia</td>
<td>LV hypertrophy</td>
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<tr>
<td>MERFF</td>
<td>Mitochondrial</td>
<td>Myoclonic seizures, cerebellar ataxia, progressive muscular weakness</td>
<td>LV hypertrophy</td>
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<tr>
<td></td>
<td>AR</td>
<td></td>
<td>Dilated cardiomyopathy</td>
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<tr>
<td>Friedreich’s ataxia</td>
<td>AR</td>
<td>Cerebellar ataxia, dysthria, loss of proprioception, lower extremities muscle weakness, extensor plantar response, areflexia, diabetes mellitus</td>
<td>LV hypertrophy</td>
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<tr>
<td></td>
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<td>Dilated cardiomyopathy</td>
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<td>Sudden cardiac death</td>
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<td><strong>Muscular dystrophies</strong></td>
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<tr>
<td>Duchenne and Becker muscular dystrophy</td>
<td>X-linked</td>
<td>Gradually progressive proximal weakness with later involvement of the majority of muscles including respiratory muscles, slower progression in Becker muscular dystrophy</td>
<td>Subclinical LV dilatation, reduction in LV ejection fraction, severe LV dysfunction, dilated cardiomyopathy and clinical heart failure, conduction blocks, sinus node dysfunction, Q waves on ECG mainly in the lateral leads</td>
</tr>
<tr>
<td>Emery Dreifuss muscular dystrophy</td>
<td>X-linked, AD, AR</td>
<td>Early contractures of the elbows, ankles and cervical spine, slowly progressive weakness that is proximal in the upper limbs and distal in the lower limbs with later involvement of the proximal limb-girdle muscles</td>
<td>Severe AV conduction abnormalities including conduction block, bradycardia, atrial fibrillation/flutter, non-sustained ventricular tachycardia, LV dilation, dilated cardiomyopathy, sudden cardiac death</td>
</tr>
<tr>
<td>Limb girdle muscular dystrophy</td>
<td>AD,AR</td>
<td>Proximal upper and lower limb-girdle weakness</td>
<td>Dilated cardiomyopathy, Q waves on ECG mainly in the lateral leads, conduction blocks, atrial fibrillation, ventricular tachycardia, sudden cardiac death</td>
</tr>
<tr>
<td>Facioscapulohumeral muscular dystrophy</td>
<td>AD</td>
<td>Facial, shoulder and upper arm weakness, foot dorsiflexors weakness, hip and pelvis weakness</td>
<td>Conduction blocks, possibly stress-induced perfusion abnormalities</td>
</tr>
<tr>
<td>Myotonic dystrophy 1</td>
<td>AD</td>
<td>Ptosis, weakness of the face, jaw and anterior neck muscles, progressive muscle weakness, myotonia</td>
<td>Sinus bradycardia, conduction blocks, atrial fibrillation/flutter, LV dysfunction, left atrial dilatation</td>
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<tr>
<td>Diseases of metabolism</td>
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<tr>
<td>Fabry’s disease</td>
<td>X-linked</td>
<td>Multisystemic disease involving the kidneys, nervous system, heart and gastrointestinal system, stroke and neuropathic pain, angiokeratoma, corneal dystrophy</td>
<td>LV hypertrophy</td>
</tr>
<tr>
<td>Pompe disease</td>
<td>AR</td>
<td>Multisystemic disease, muscle weakness, respiratory failure, macroglossia</td>
<td>LV hypertrophy</td>
</tr>
</tbody>
</table>

AD, autosomal dominant; AR, autosomal recessive; AV, atrioventricular; ECG, electrocardiogram; LV, left ventricular; MELAS, mitochondrial encephalopathy with lactic acidosis and stroke-like episodes; MERFF, myoclonic epilepsy with ragged red fibers.
CONFLICTS OF INTEREST

common neuromuscular disorders, although not impossible. 103,104

CONCLUSIONS

Cardiac and neurologic diseases frequently overlap. Randomized trials are ongoing to elucidate the best management strategies in many of these disorders, and have been successful in providing certain management guidelines such as for stroke prevention in AF. Similar studies are more difficult for the less common neuromuscular disorders, although not impossible.103,104

FUNCTIONS OF INTEREST

None declared.

REFERENCES


