Diagnosis and management of Duchenne muscular dystrophy, part 2: respiratory, cardiac, bone health, and orthopaedic management

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A coordinated, multidisciplinary approach to care is essential for optimum management of the primary manifestations and secondary complications of Duchenne muscular dystrophy (DMD). Contemporary care has been shaped by the availability of more sensitive diagnostic techniques and the earlier use of therapeutic interventions, which have the potential to improve patients’ duration and quality of life. In part 2 of this update of the DMD care considerations, we present the latest recommendations for respiratory, cardiac, bone health and osteoporosis, and orthopaedic and surgical management for boys and men with DMD. Additionally, we provide guidance on cardiac management for female carriers of a disease-causing mutation. The new care considerations acknowledge the effects of long-term glucocorticoid use on the natural history of DMD, and the need for care guidance across the lifespan as patients live longer. The management of DMD looks set to change substantially as new genetic and molecular therapies become available.

Introduction

The 2010 care considerations for Duchenne muscular dystrophy (DMD)1,2 advocated a multidisciplinary approach to the management of this severe, progressive neuromuscular disease. This three-part update was necessitated by a number of themes that characterise contemporary DMD care: the increasing complexity of subspecialty care and the need for a multidisciplinary clinical team; the use of more sensitive diagnostic techniques and earlier therapeutic interventions; the expectation of prolonged survival, prompting the need for care guidance across the lifespan; and the recognition that the natural history of DMD has been altered by the long-term use of glucocorticoids.3 The new care considerations have also been shaped by the expectation that emerging genetic and molecular therapies will substantially change the nature of DMD management in the near future.

In 2014, the DMD Care Considerations Working Group steering committee, comprising experts from a wide range of disciplines, identified 11 topics to be included in this update. Part 2 contains the latest care considerations for respiratory, cardiac, bone health and osteoporosis, and orthopaedic and surgical management. Large-scale, randomised controlled trials (RCTs) are rare in this field, so guidance was developed using a method that queries a group of experts on the appropriateness and necessity of specific assessments and interventions, using clinical scenarios. This methodology was designed to produce an essential toolkit for DMD care; only assessments and interventions that have been deemed both appropriate and necessary are recommended. A complete description of the methods is provided in part 1 and the appendix.

Figure 1 in part 1 of this Review provides a brief overview of assessments and interventions across all topics, organised by stage of disease. It is intended to serve as a pocket guide to overall disease management.

Respiratory management

Respiratory complications are a major cause of morbidity and mortality in people with DMD. Complications include respiratory muscle fatigue, mucus plugging, atelectasis, pneumonia, and respiratory failure. If left untreated, patients are at risk of severe dyspnoea, lengthy hospital admissions due to atelectasis or pneumonia, and death due to respiratory arrest or respiratory-induced cardiac arrhythmias.4,5

An anticipatory approach to management includes monitoring of respiratory muscle function and the timely use of lung volume recruitment, assisted coughing, nocturnally assisted ventilation, and subsequent daytime ventilation. These core therapies can decrease respiratory complications, improve quality of life, and prolong survival.6,7 Patients should typically be using most or all of these core therapies by the age of 18–21 years, before their transition from paediatric to adult respiratory care providers.

Implementation of respiratory care considerations and guidelines7,8,9 requires a multidisciplinary team—including physicians, respiratory therapists (or physical therapists in some health-care systems), and home caregivers—to perform pulmonary function testing and sleep studies to initiate and manage lung volume recruitment,10 manual and mechanically assisted coughing,1,12,14 non-invasive ventilation, and invasive ventilation via tracheostomy. Decisions for optimum respiratory management need to be made with awareness of the patient’s other body systems, especially the cardiac system.9,14

In this update, we endorse higher pulmonary function thresholds (ie, milder levels of respiratory impairment)
for initiation of assisted coughing and assisted ventilation than were recommended in the 2010 care considerations. The new criteria are intended to result in more anticipatory use of these interventions, with the possibility that therapy will be initiated in slightly younger patients.

**Ambulatory stage**

Figure 1 shows respiratory diagnostic tests and therapies for individuals with DMD, by stage of disease. Spirometry should be initiated when the patient is 5–6 years of age. Serial monitoring of pulmonary function is critical for respiratory management. Typically, forced vital capacity (FVC) rises with growth, until an individual becomes non-ambulatory. FVC reaches a peak, followed by a plateau, and then deteriorates over time.19–21 Deteriorating FVC can occur in the absence of dyspnoea and remain unrecognised unless pulmonary function is measured regularly. In a large cohort study in boys who had not been treated with corticosteroids, the age at loss of ambulation was predictive of the age at which peak FVC was realised, the absolute peak FVC, and the rate of subsequent decline.19 For example, earlier loss of ambulation was associated with an earlier and lower peak FVC as well as a more rapid decline in FVC than was later loss of ambulation. However, because the rate of change in FVC over time can vary greatly among individuals, serial measurement of FVC is necessary to characterise each individual's respiratory phenotype or trajectory.

Sleep studies with capnography might be necessary during the ambulatory stage, especially for individuals with weight gain due to glucocorticoid therapy and for individuals with symptoms of sleep-disordered breathing. Sleep studies can also be used as an alternative method to monitor respiratory status among individuals who cannot cooperate with pulmonary function testing.

Individuals with DMD should receive yearly immunisation with the inactivated influenza vaccine (ie, the injectable vaccine, not the live, attenuated nasal vaccine) and pneumococcal vaccines (including PCV13 and

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<td>Addition of assisted daytime ventilation when, despite nocturnal ventilation,§ daytime SpO₂ &lt;95%, pCO₂ &gt;45 mm Hg, or symptoms of awake dyspnoea are present</td>
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*See text for definitions of sleep study results. †All specified threshold values of PCF, MEP, and MIP apply to older teenage and adult patients. §Fatigue, dyspnoea, morning or continuous headaches, frequent nocturnal awakenings or difficult arousal, hypersomnia, difficulty concentrating, awakenings with dyspnoea and tachycardia, or frequent nightmares. We strongly endorse the use of non-invasive methods of assisted ventilation instead of tracheostomy to optimise patient quality of life; indications for tracheostomy include patient preference, inability of patient to use non-invasive ventilation successfully, three failed extubation attempts during a critical illness despite optimum use of non-invasive ventilation and mechanically assisted coughing, or failure of non-invasive methods of cough assistance to prevent aspiration of secretions into the lungs due to weak bulbar muscles.
PPSV23), according to guidelines available from the US Centers for Disease Control and Prevention, other public health entities such as the Immunization Action Coalition, and Parent Project Muscular Dystrophy.

Patients and their caregivers should be educated about respiratory complications during the ambulatory stage of DMD to prepare them for future medical complications and therapies.

**Early non-ambulatory stage**

The need for respiratory interventions occurs mainly after the loss of ambulation (figure 1). Seated FVC (expressed both as an absolute value and as a percentage predicted on the basis of arm span or ulnar length), maximum inspiratory and expiratory pressures, peak cough flow, and blood oxygen saturation by pulse oximetry (SpO2) should be measured at least every 6 months in all non-ambulatory individuals. Additionally, end-tidal or transcutaneous partial pressure of carbon dioxide in the blood (p_CO2 or ____ ptcCO2, respectively) should be measured every 6 months or any time SpO2 is 95% or lower on room air, when the necessary equipment is available.

As their vital capacity decreases, patients with DMD develop stiff, non-compliant chest walls and lung volume restriction. To preserve lung compliance, lung volume recruitment is indicated when FVC is 60% predicted or less, achieved with a self-inflating manual ventilation bag or mechanical insufflation–exsufflation device to provide deep lung inflation once or twice daily.25–27

During the early non-ambulatory stage, some individuals with DMD need surgery for progressive scoliosis.28 Previously published guidelines address respiratory management of patients undergoing surgery, including indications for preoperative training in the use of assisted cough devices and non-invasive ventilators.11 For patients who are cognitively impaired and unable to reliably perform pulmonary function testing, preoperative polysomnography might be helpful.

**Late non-ambulatory stage**

As they progress through the non-ambulatory stage, individuals with DMD develop weak cough efforts, placing them at risk of atelectasis, pneumonia, ventilation–perfusion mismatch, and progression to respiratory failure, especially during respiratory tract infections. Treatment consists of manual and mechanically assisted coughing, which are indicated when FVC is less than 50% predicted, when peak cough flow is less than 270 L/min, or when maximum expiratory pressure is less than 60 cm H2O (figure 1).29–33

We advise having a home pulse oximeter for individuals treated with assisted coughing during respiratory infections. When SpO2 is less than 95% on room air, the frequency of assisted coughing should be increased to prevent and treat mucus plugging, atelectasis, and pneumonia. We also recommend initiation of antibiotic therapy during acute respiratory illnesses when individuals have three of the following five signs of pneumonia: fever, elevated white blood count or C-reactive protein concentration, sputum production, a pulmonary infiltrate on chest radiograph, or hypoxaemia or respiratory distress.

In the late non-ambulatory stage, individuals with DMD need assisted ventilation to prolong survival.32 Ventilation devices should incorporate a back-up rate of breathing to avoid apnoea. Indications for nocturnally assisted ventilation include signs or symptoms of hypoventilation or sleep-disordered breathing, irrespective of the level of pulmonary function; relevant symptoms include fatigue, dyspnoea, morning or continuous headaches, frequent nocturnal awakenings or difficult arousal, hypersomnolence, difficulty concentrating, awakenings with dyspnoea and tachycardia, and frequent nightmares. However, some individuals remain asymptomatic despite the presence of hypoventilation.33 Thus, nocturnally assisted ventilation should be initiated when a patient’s FVC is less than 50% predicted, or when the absolute value of maximum inspiratory pressure is less than 60 cm H2O. It should also be initiated when the individual is awake and, because of daytime hypoventilation, any of the following is true: p_CO2 or ____ ptcCO2 is more than 45 mm Hg; arterial, venous, or capillary blood pCO2 is more than 45 mm Hg; or baseline SpO2 is less than 95% on room air (figure 1).31–33

Nocturnal ventilation is also indicated for individuals with abnormal sleep studies, including overnight oximetry, combination oximetry–capnography, and polysomnography with capnography. Non-ambulatory individuals with symptoms of sleep-disordered breathing should have sleep studies as often as annually, if possible. Sleep study results that indicate the need for assisted ventilation include p_CO2 or ____ ptcCO2 of more than 50 mm Hg for at least 2% of sleep time, a sleep-related increase in p_CO2 or ____ ptcCO2 of 10 mm Hg above the awake baseline for at least 2% of sleep time, an SpO2 of 88% or less for at least 2% of sleep time or for at least 5 min continuously, or an apnoea–hypopnoea index of five events per h or more.33–35 Because patients with DMD inevitably need assisted ventilation to treat hypoventilation, nocturnal non-invasively assisted ventilation (rather than continuous positive airway pressure at a constant level) is first-line therapy for individuals with DMD with obstructive sleep apnoea.

Non-invasive ventilation can also be used during and after procedures involving anaesthesia or sedation and, in conjunction with assisted coughing, to extubate individuals who are mechanically ventilated for respiratory infections.41 In DMD, hypoxaemia is usually due to hypoventilation, atelectasis, or pneumonia. Therefore, supplemental oxygen therapy should not be used alone. In conjunction with assisted ventilation and

For more on the Immunization Action Coalition see http://immunize.org/ For more on Parent Project Muscular Dystrophy see http://www.parentprojectmd.org/
assisted coughing, oxygen therapy can be safe, especially when blood CO₂ levels are monitored.

With declining pulmonary function, patients develop symptoms of hypoventilation such as dyspnoea, fatigue, and difficulty concentrating, despite their use of assisted ventilation during sleep; those with very low FVCs (<680 mL in one study) are at particular risk. Thus, patients often self-extend their use of assisted ventilation into the daytime, ultimately up to 24 h/day. The indications for daytime-assisted ventilation are listed in figure 1. Options for continuous non-invasive ventilation include mouthpiece or so-called sip ventilation with a portable volume ventilator during the day, changing to nasal ventilation with a bi-level pressure device overnight. Alternatively, 24 h/day nasal ventilation with a bi-level pressure device can be effective and well tolerated.³²,⁴² These devices should have an internal battery for safety and portability.

Whether individuals with DMD should be ventilated via tracheostomy or non-invasively is a controversial question. Some centres use time on the ventilator (eg, 16 h/day or more) as an indication for tracheostomy.³³–³⁵ However, clinical experience supports the use of non-invasively assisted ventilation for up to 24 h/day.³²,⁴² We strongly endorse the use of non-invasive ventilation in most clinical situations. Our indications for tracheostomy are listed in figure 1, and include patient preference, inability to use non-invasive ventilation, three failed extubation attempts during a critical illness despite optimum use of non-invasive ventilation and mechanically assisted coughing, or failure of non-invasive methods of cough assistance to prevent aspiration of secretions into the lungs due to weak bulbar muscles. Overall, the decision is highly dependent on each individual’s preference and clinical course, the skills and usual practices of the individual’s clinicians, the local standard of care, and the availability of home resources, such as overnight nursing.³⁶ The use of non-invasive respiratory aids is especially challenging when individuals with very advanced DMD have acute respiratory illnesses and when they have chronic difficulty swallowing their saliva.

Continuous ventilation provides life support, so a back-up ventilator and a manual resuscitator should be available in case the primary ventilator malfunctions. Batteries or a generator should be available for use during a power outage. The ventilation device and battery should attach to the individual’s wheelchair for mobility and quality of life. When practical, the presence of a night nurse can greatly decrease the risk of potentially catastrophic medical events, such as mucus plugging of the trachea.

Cardiac management

Cardiovascular complications are a leading cause of disease-related morbidity and mortality among individuals with DMD.⁴⁶ Dystrophin deficiency in the heart manifests as a cardiomyopathy. As the disease progresses, the myocardium fails to meet physiological demands and clinical heart failure develops. The failing myocardium is also at risk of life-threatening rhythm abnormalities.⁴⁷

Historically, individuals with DMD have not been referred to a cardiac specialist until late in the disease, contributing to poor clinical outcomes. Furthermore, cardiac management has been challenging because the New York Heart Association classification of heart failure³⁹ relies on reduced exercise tolerance, a feature that in DMD arises from skeletal muscle and cardiac disease combined. The signs and symptoms of heart failure in the non-ambulatory individual are frequently subtle and overlooked. A proactive strategy of early diagnosis and treatment is essential to maximise duration and quality of life. Involvement of a cardiologist who is integrated into a multidisciplinary care team is recommended, given the complex decision making involved in managing DMD cardiomyopathy. Ideally, the cardiologist should have clinical expertise in diagnosing and treating heart failure and the cardiomyopathy associated with neuromuscular disease, and have readily available access to state-of-the-art expertise in non-invasive imaging. A National Heart, Lung, and Blood Institute (NHLBI) expert working group was convened and recently published updated comprehensive DMD cardiac care considerations, including important areas for future research.³⁷ The specific core care considerations are detailed below and summarised in figure 2.

Ambulatory stage and early non-ambulatory stage

The baseline cardiac assessment includes past and present cardiac medical history, family history, and a physical examination. Electrocardiogram and non-invasive imaging are advised to establish baseline cardiac function and to screen for underlying anatomical abnormalities that could affect long-term cardiovascular health. Cardiovascular MRI (CMR) is the non-invasive imaging modality of choice; however, young individuals might not be able to cooperate for the procedure. Thus, echocardiography is recommended until at least age 6–7 years, when CMR can typically be done without anaesthesia. Until the age of 10 years, individuals should have an annual cardiac assessment, including electrocardiogram and non-invasive imaging. After the age of 10 years, asymptomatic individuals should have a cardiac assessment at least annually because of the increased risk of left ventricular dysfunction. With the onset of heart failure symptoms or when abnormalities are first seen on cardiac imaging—eg, myocardial fibrosis, left ventricular enlargement, or left ventricular dysfunction—the frequency of assessment should increase at the discretion of the cardiologist. An electrocardiogram and non-invasive cardiac imaging should be done before major surgical procedures, such as scoliosis correction. DMD is associated with a

For more on the National Heart, Lung, and Blood Institute see https://www.nhlbi.nih.gov/
particular set of anaesthesia risks, and the anaesthetist should be made aware of the patient’s cardiac history.52

Traditionally, angiotensin-converting enzyme (ACE) inhibitors or angiotensin receptor blockers (ARBs) were used as first-line therapy for the treatment of heart disease associated with DMD. Opinion differs on the use of ACE inhibitors in very young (<10 years) asymptomatic individuals without evidence of abnormality on CMR or echocardiogram. After discussing potential benefits and risks with the family, the cardiologist could initiate therapy in this group of individuals. Some evidence suggests that initiation of ACE inhibitors in asymptomatic boys with normal left ventricular systolic function as they approach 10 years of age can improve long-term cardiac outcomes, and the 2014 NHLBI working group recommended use of ACE inhibitors or ARBs by the age of 10 years in boys with DMD.51 Dosing and ACE inhibitor selection are left to the discretion of the cardiologist.53

Irrespective of age, pharmacological therapy should be initiated with the onset of heart failure symptoms or when abnormalities such as depressed left ventricular ejection fraction, abnormal chamber dimensions, or the presence of myocardial fibrosis are noted on imaging studies (CMR or echocardiogram). After discussing potential benefits and risks with the family, the cardiologist could initiate therapy in this group of individuals. Some evidence suggests that initiation of ACE inhibitors in asymptomatic boys with normal left ventricular systolic function as they approach 10 years of age can improve long-term cardiac outcomes, and the 2014 NHLBI working group recommended use of ACE inhibitors or ARBs by the age of 10 years in boys with DMD.51 Dosing and ACE inhibitor selection are left to the discretion of the cardiologist.53

Irrespective of age, pharmacological therapy should be initiated with the onset of heart failure symptoms or when abnormalities such as depressed left ventricular ejection fraction, abnormal chamber dimensions, or the presence of myocardial fibrosis are noted on imaging studies (CMR or echocardiogram). Given the absence of dystrophin-specific targeted cardiac therapies, traditional treatment strategies for heart failure should be used. β-adrenergic blockade is typically started with evidence of ventricular dysfunction. In a prospective, randomised double-blind placebo-controlled trial in patients aged 7–25 years with DMD, the mineralocorticoid receptor antagonist eplerenone attenuated the decline in cardiac function, as measured by circumferential strain.54 This benefit was supported by findings from a 2 year, open-label extension trial.55 However, although eplerenone might prove to be a useful adjunctive therapy to other heart failure medications, further investigations are needed to establish effectiveness.54,55

Late non-ambulatory stage
Progressive myocardial fibrosis leads to ventricular dysfunction. More frequent cardiac monitoring, as determined by the patient’s cardiologist, is advised in the late, non-ambulatory stage to reduce disease-related morbidity and mortality. The cardiologist should work closely with the multidisciplinary care team to ensure that respiratory care has been optimised, because abnormal pulmonary mechanics affect cardiac function.56,57 Specifically, there is evidence that non-invasive nocturnal ventilation increases long-term survival.8 The NHLBI working group suggested that early initiation of nocturnal ventilation be considered because of potential long-term benefit.51

Symptomatic heart failure can be particularly difficult to diagnose in non-ambulatory patients with DMD. Clinical manifestations of heart failure—fatigue, weight loss, vomiting, abdominal pain, sleep disturbance, and inability to tolerate daily activities—are often unrecognised until late in the disease because of musculoskeletal limitations. The cardiologist should maximise medical therapy for heart failure. Consideration should also be given to thromboembolism prevention in individuals with severe left ventricular dysfunction.

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**Figure 2:** Cardiac monitoring, diagnosis, and treatment algorithm for patients with Duchenne muscular dystrophy
Various antithrombotic drugs are available, and should be initiated after discussion with the cardiologist. People with DMD are at risk of rhythm abnormalities—including atrial fibrillation or flutter, ventricular tachycardia, and ventricular fibrillation—that can be treated with standard antiarrhythmic medications or device management, when indicated. Surveillance should include periodic Holter monitoring. In most circumstances, 24 h Holter monitoring will be sufficient. Event monitors could also be indicated when individuals complain of episodic, non-sustained rhythm disturbances. The optimum frequency of monitoring has not been established and should be directed by the cardiologist, depending on the patient’s clinical course. It is reasonable to initiate annual Holter monitor screening with the onset of abnormal left ventricular function or development of myocardial fibrosis. The benefit of implantable cardioverter defibrillators as primary prevention for ventricular tachycardia or ventricular fibrillation is unknown. These devices can be used for secondary prevention in patients who have had ventricular tachycardia or ventricular fibrillation. At present, placement for primary arrhythmia prevention is on the basis of established adult heart failure guidelines. Among adults with heart failure, placement of implantable cardioverter defibrillators is advised for individuals with an ejection fraction of less than 35%. Clear patients with DMD have unique issues (eg, chest wall deformities and sedation risk), which might affect this recommendation.

In individuals in whom maximal medical management has failed, the use of mechanical circulatory support is a therapeutic consideration, as illustrated by relevant case reports. A left ventricular assist device could be used as a destination therapy—ie, in individuals for whom a heart transplant is not considered appropriate. The decision to proceed with a ventricular assist device is complex and involves a deep understanding of all the inherent risks and potential benefits. Risks include, but are not limited to, thromboembolism, bleeding, infection, device malfunction, and right heart failure. In an ideal situation, the device has the potential to improve duration and quality of life. Cardiac transplantation is also a theoretical option, but given the small number of available donors, it needs to be considered on a case-by-case basis.

Female carriers

In this update, we acknowledge that female carriers of a disease-causing mutation are at risk of not only skeletal muscle disease, but also cardiomyopathy. The natural history and incidence of cardiomyopathy in girls and women is not well characterised, but in a 2016 study, 47% of carriers had at least one positive finding on CMR. We recommend a baseline cardiac assessment in early adulthood that includes an electrocardiogram and non-invasive imaging, preferably CMR, when available. Ongoing surveillance will be required, on the basis of guidance for individuals with cardiomyopathy. The optimum frequency has not been established in the DMD carrier population, but our current guidance is for assessment every 3–5 years, on the basis of screening recommendations for other genetic cardiomyopathies.

Bone health and osteoporosis management

Boys with glucocorticoid-treated DMD frequently develop osteoporosis, which manifests as low-trauma vertebral or long-bone fractures. This outcome is not surprising given the potent osteotoxicity of glucocorticoid therapy combined with progressive myopathy, both of which are key risk factors for reduced bone strength. 20–60% of boys with DMD have low-trauma extremity fractures (usually the distal femur, tibia, or fibula), while up to 30% develop symptomatic vertebral fractures. Vertebral fractures are frequently asymptomatic when identified in children with glucocorticoid-treated illnesses through a monitoring programme that includes a lateral spine radiograph, so the true prevalence is probably higher than existing reports suggest. Left untreated, vertebral fractures can lead to chronic back pain and spine deformity, while leg fractures can cause premature, permanent loss of ambulation. Death due to fat embolism syndrome after long-bone fractures has also been reported in boys with DMD.

The notion that some glucocorticoid agents and dosing regimens are bone-sparing compared with others has arisen from studies of deflazacort versus prednisolone or meprednisone (also known as methylprednisone) in children after renal transplant and in those with chronic juvenile arthritis. The steroid dose equivalences used in these studies were variable, making comparisons difficult; however, disease outcomes were favourable in the deflazacort-treated children, with associated improvements in bone density outcomes, linear growth, weight–height ratios, and lean body mass. By contrast, recent publications cast doubt on the bone-sparing properties of deflazacort, showing that bone fragility (including vertebral fractures) is frequent in deflazacort-treated boys with DMD, probably related partly to the large doses that are used in this condition. Comparative studies of different steroid regimens in DMD are underway, assessing the effect on final adult height, body composition, and fractures.

Despite the high prevalence of fractures, no published studies of DMD or any osteoporotic condition of childhood have assessed the safety and efficacy of medical therapy in preventing the first-ever fracture. Therefore, the current standard is to identify and treat early indications of bone fragility (eg, vertebral fractures) in individuals with chronic illnesses who have little possibility of recovery. This secondary prevention approach has the goal of mitigating osteoporosis progression and promoting recovery among patients presenting with early, rather than late, indications of osteoporosis and in those with little potential for
medication-unassisted recovery because of persistent risk factors.

We present care considerations for monitoring that will enable timely diagnosis and treatment of osteoporosis in boys and men with DMD (figure 3). We also review specific diagnostic criteria for osteoporosis, along with care considerations for prescription of osteoporosis therapy, including agents, dose, and duration of therapy. Comprehensive reviews of all issues relating to management of osteoporosis therapy (including contra-indications and monitoring of safety and efficacy) have been published elsewhere.81,82

**Bone health monitoring and diagnosis of osteoporosis**

An important development that distinguishes the current guidance from the 2010 care considerations is that bone health monitoring and diagnosis in children no longer focus on bone mineral density (BMD); rather, BMD serves as an adjuvant in an approach that focuses on bone mineral density (BMD) rather than BMD.88

**Monitoring and diagnosis**

At each clinical visit
- Presence of back pain or fractures

At baseline only (follow up as appropriate)
- Serum calcium
- Phosphate
- Magnesium
- Alkaline phosphatase
- Parathyroid hormone

At baseline and annually
- Calcium/vitamin D intake
- Spine BMD by DXA
- Serum 25-hydroxyvitamin D

At baseline and follow-up
- Lateral thoracolumbar spine radiograph
- On steroids, every 1–2 years
- Not on steroids, every 2–3 years

If back pain or ≥0.5 SD decline in spine BMD Z score over 12-month period
- Lateral thoracolumbar spine radiograph

Continue monitoring until signs of bone fragility

**Treatment: stabilisation phase**

Before initiating intravenous bisphosphonate therapy
- Treat calcium/vitamin D deficiency
- Verify normal renal function

When starting intravenous bisphosphonate therapy
- Follow published regimen
- Treat until clinically stable

For monitoring of safety and efficacy of treatment
- Obtain thoracolumbar spine radiograph annually and monitor the following every 6 months:
  - Spine BMD by DXA
  - Serum hydroxyvitamin D
  - Patient-reported back pain
  - Calcium/vitamin D intake
  - Biochemical markers of bone and mineral ion metabolism

**Clinical stable†**

**Treatment: maintenance phase**

Once clinically stable
- Consider continuing intravenous bisphosphonate therapy with titration to a lower dose to preserve gains realised during stabilisation phase
- Vary duration of maintenance therapy depending on bone health status and whether steroid therapy is ongoing

Monitor safety and efficacy of maintenance therapy

**Clinically significant bone fragility**

**No longer clinically stable**

Vertebral fractures can occur in children who have BMD Z scores higher than –2 SD, an observation that invalidates the use of a BMD Z score threshold to define osteoporosis in children with low-trauma vertebral fractures.86 This observation prompted the International Society for Clinical Densitometry to revise the definition of osteoporosis in a child with a low-tramua vertebral fracture so that cutoff criteria based on BMD Z scores are no longer required to make the diagnosis of osteoporosis.88 Similarly, 15% of children with neuromuscular disorders and extremity fractures will have BMD Z scores for the distal femur higher than –2 SD,89 which again challenges the use of a BMD Z score threshold to define osteoporosis in a child with extremity fractures. Finally, findings from a recent study87 showed that spine BMD Z scores can vary by as much as

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**Figure 3:** Osteoporosis monitoring, diagnosis, and treatment algorithm for patients with Duchenne muscular dystrophy

BMD = bone mineral density. DMD = Duchenne muscular dystrophy.DXA = dual-energy x-ray absorptiometry. *Signs of clinically significant bone fragility are low-trauma fractures of long bones or vertebra. †Clinical stability refers to absence of non-vertebral fractures, stable healed vertebral fractures, absence of new vertebral fractures in previously normal vertebral bodies, absence of bone and back pain, and BMD Z score appropriate for height Z score or higher than –2 SD.
2 SD depending on the normative database that is used to generate the \( Z \) scores. In view of these findings, the diagnosis of osteoporosis in at-risk children now rests on the presence of evident bone fragility, often manifesting as vertebral fractures, and a BMD \( Z \) score above −2 SD does not preclude the diagnosis of osteoporosis.\(^8\) Although BMD \( Z \) scores are no longer at the forefront of diagnosis, they remain useful to determine the overall trajectory of bone health in an individual child and thereby guide frequency of lateral spine radiographs during the monitoring phase.

The 2010 care considerations recommended a spine radiograph for vertebral fracture detection in patients with a history of back pain or spine deformity on physical examination. In the current care considerations, a baseline spine radiograph for vertebral fracture detection is recommended in all patients, with intermittent follow-up radiographs to assess changes in spine morphology in the face of persistent (ie, glucocorticoid therapy) or permanent (ie, myopathy) risk factors. Given the need for serial spine radiographs, assessment by dual-energy x-ray absorptiometry is an emerging method for at-risk populations, and a validation study in children has shown that this technique compares favourably with detection of Genant-defined vertebral fractures on spine radiographs.\(^9\) Overall, spine radiographs should be prioritised over BMD in view of the need to detect the earliest signs of bone fragility.

**Treatment of osteoporosis**

Indications for treatment with intravenous bisphosphonate—the presence of low-trauma vertebral fractures or long-bone fractures—generally remain unchanged, but with notable differences in the timing of treatment initiation. Previously, only back pain or spine deformity prompted a radiograph to identify vertebral fractures necessitating bisphosphonate therapy. The current call for routine spine radiographs for all patients with DMD will lead to diagnosis of symptomatic vertebral fractures (mild, moderate, and severe) and asymptomatic moderate and severe vertebral fractures, all of which should prompt referral to an osteoporosis expert for treatment. Because even mild and asymptomatic vertebral fractures are predictive of future fractures in both children\(^7\) and adults,\(^9\) treatment of asymptomatic moderate (Genant grade 2) and severe (Genant grade 3) vertebral fractures is now recommended. Treatment with intravenous bisphosphonate therapy had a protective effect on spine BMD and vertebral morphology in controlled trials of osteogenesis imperfecta\(^9,10\) and DMD.\(^11,12\) Additional support for treatment of asymptomatic but nevertheless advanced (ie, moderate and severe) vertebral fractures stems from the fact that no cases of spontaneous (ie, medication unassisted) reshaping of previously fractured vertebral bodies have been reported in boys with DMD;\(^13\) however, reshaping has been observed after intravenous bisphosphonate therapy in this population.\(^14\) For children with glucocorticoid-treated diseases, such as DMD, including those with minimally symptomatic or asymptomatic mild (grade 1) vertebral fractures, controlled trials are underway to investigate the efficacy of antiresorptive therapy (ClinicalTrials.gov identifiers NCT00799266 and NCT02632916); for now, mild asymptomatic fractures should be closely monitored for symptomatology or progressive height loss that would prompt treatment.

The updated guidance represents a fundamental change in the goals of therapy. The aim is to identify and treat the earliest signs of bone fragility to better preserve the heights of the vertebral bodies.\(^15\) We endorse the use of intravenous (and not oral) bisphosphonates as first-line therapy for the treatment of osteoporosis in patients with DMD,\(^16,17\) on the basis of an extrapolation from results of controlled trials in osteogenesis imperfecta. Such studies have shown increased vertebral heights in growing patients with osteogenesis imperfecta treated with intravenous bisphosphonate therapy.\(^18,19\) By contrast, no controlled studies of oral bisphosphonates in osteogenesis imperfecta have shown an effect on vertebral height.\(^20,21\) These data are particularly relevant to patients with glucocorticoid-treated DMD, who have a high frequency of vertebral fractures.\(^22\) Recent reviews on the management of children with fractures due to osteoporosis concur with the view that intravenous rather than oral bisphosphonates should be used as first-line therapy.\(^23,24\) Because bisphosphonates remain off label for children in most countries and they require judicious prescription, patients with a low-trauma fracture should be referred to an expert in osteoporosis management to ensure proper bisphosphonate dosing, dose titration on longer-term therapy, timing of treatment cessation, and monitoring of treatment safety and efficacy.

**Orthopaedic and surgical management**

The overall aim of musculoskeletal care is to maintain motor function for as long as possible, minimise joint contractures, maintain a straight spine, and promote bone health. The assessment and treatment of musculoskeletal complications should involve an interdisciplinary team that might include a physical and occupational therapist, rehabilitation physician, neurologist, orthopaedic surgeon, and social worker. When a surgical intervention is recommended, it is crucial to involve the respiratory physician and cardiologist.

Figure 4 outlines the care considerations for orthopaedic and surgical care related to contracture, spine, and fracture management. Figure 5 provides general guidance for patients and families about fracture prevention. In the absence of RCTs comparing different therapeutic and surgical approaches, this guidance is based on the expert consensus of orthopaedic and rehabilitation specialists, using the methods described in part 1. Care considerations on stretching, orthoses, and adaptive equipment for contracture management are provided in the rehabilitation management section of part 1.
Ambulatory stage

Children in the ambulatory stage might benefit most from surgery, but it is recommended less frequently than in the past. Although the 2010 care considerations included recommendations for multilevel surgeries, the consensus of the current panel is that surgery on the foot to improve the varus positioning and on the Achilles tendon to improve dorsiflexion range might be sufficient to improve gait in patients with clinically significant ankle contracture and good quadriceps and hip extensor strength. Interventions related to the hips and knees are not recommended.

Assessment for scoliosis should be done at least annually, although onset is unusual in the ambulatory stage. Visual assessment is appropriate, with radiographic assessment only if a curve is observed on examination or if visual inspection alone is inadequate, such as in children with obesity. The preceding section on bone health and osteoporosis management provides information regarding monitoring and treatment of spinal compression fractures. Use of spinal orthoses is not generally recommended in the setting of a compression fracture.

Anticipatory guidance during routine clinic visits is an important part of a fracture prevention programme throughout all disease stages (figure 5). As noted, corticosteroids have been associated with osteoporosis and subsequent vertebral fractures in DMD. In a study of 143 boys with DMD, the long-bone fracture rate in those treated with corticosteroids was 2.6 times greater than in those who had never received steroids. A lower-limb

<table>
<thead>
<tr>
<th>Ambulatory stage</th>
<th>Early non-ambulatory stage</th>
<th>Late non-ambulatory stage</th>
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<tbody>
<tr>
<td><strong>Assessments</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Assess range of motion at least every 6 months</td>
<td>Conduct visual inspection of the spine annually</td>
<td>Conduct visual inspection of the spine every 6 months</td>
</tr>
<tr>
<td>Conduct visual inspection of the spine annually</td>
<td>Obtain radiographic assessment if curve observed or visual inspection difficult</td>
<td>Obtain spine radiograph when patients become non-ambulatory, if curve present, obtain radiograph every 6 months to 1 year, depending on skeletal maturity, refer to orthopaedic surgeon for curve &gt;20°</td>
</tr>
<tr>
<td>Obtain radiographic assessment if curve observed or visual inspection difficult</td>
<td>Obtain annual anteroposterior upright spinal radiographs for patients with known progressive scoliosis</td>
<td>Obtain anteroposterior upright spinal radiographs for patients with known progressive scoliosis</td>
</tr>
<tr>
<td><strong>Interventions</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>With physical therapy guidance, implement home stretching programme focusing on ankles, knees, and hips</td>
<td>With occupational therapy guidance, add focus on upper extremities</td>
<td>Use custom-molded daytime ankle-foot orthoses to delay worsening of equinovarus contracture</td>
</tr>
<tr>
<td>When passive dorsiflexion &lt;10°, use custom-molded nighttime ankle-foot orthoses set in neutral position</td>
<td>Use custom-molded nighttime ankle-foot orthoses to delay worsening of equinovarus contracture</td>
<td>Continue use of lower-extremity braces; fabrication of custom wrist and hand splints may be appropriate</td>
</tr>
<tr>
<td>Refer for surgery on foot and Achilles tendon to improve gait if substantial ankle contracture with good quadriceps and hip extensor strength</td>
<td>Refer for surgery on foot and Achilles tendon to improve gait if substantial ankle contracture with good quadriceps and hip extensor strength</td>
<td>Initiate standing programme using standing device or wheelchair with upright positioning</td>
</tr>
<tr>
<td>Avoid use of spinal orthoses</td>
<td>Refer for foot and ankle surgery to improve foot positioning only if advocated by patient</td>
<td>Use standing programmes with caution</td>
</tr>
<tr>
<td>Provide anticipatory fracture prevention guidance to families</td>
<td>Consult with cardiology and respiratory specialists before any surgical intervention</td>
<td>Refer for physical therapy after surgery</td>
</tr>
<tr>
<td>Refer for physical therapy after surgery</td>
<td>Refer for posterior spinal instrumentation and fusion if spinal curve &gt;20–30° in prepubertal individuals who are not on corticosteroids; provide preoperative and postoperative evaluation with physical therapy</td>
<td>Refer for posterior spinal instrumentation and fusion if curve is progressive</td>
</tr>
<tr>
<td>Ensure families and medical team are aware of fat embolism syndrome</td>
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*Figure 4: Considerations for orthopaedic and surgical care of patients with Duchenne muscular dystrophy by stage of disease*
Common considerations or possible modifications

<table>
<thead>
<tr>
<th>Safety of home environment</th>
<th>Avoidance of falls from wheelchair or mobility device</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Remove obstacles such as rugs, toys, cords, and clutter</td>
<td>• Use seatbelt at all times</td>
</tr>
<tr>
<td>• Use anti-tippers on wheelchairs</td>
<td>• Use anti-tippers on wheelchairs</td>
</tr>
<tr>
<td>Safety on uneven or slippery surfaces</td>
<td>Safe transfer in and out of wheelchair</td>
</tr>
<tr>
<td>• Take special care when outdoors because of uneven surfaces</td>
<td>• Consider adaptive equipment and patient lift systems early for use in all settings to provide safe support and minimise risk of falls or injury during transfers, toileting, and bathing or showering</td>
</tr>
<tr>
<td>• Wear pool shoes for protection against falls when walking on slippery surfaces around water</td>
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</tr>
<tr>
<td>• Use non-slip treads on ankle-foot orthoses at night to decrease fall risk when walking to and from bathroom</td>
<td></td>
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Potential home modifications

• Non-slip mats in shower or bathtub
• Grab bars for shower or bathtub
• Bath seat or other adaptive equipment for bathing
• Non-slip treads for bare-wood steps
• Handrails on both sides of stairways

Figure 5: General guidance on fracture prevention for patients with Duchenne muscular dystrophy and their families

A fracture during the ambulatory stage might need aggressive management to maintain ambulation. Internal or external fixation allows for early mobilisation compared with casting or splinting.31

There have been case reports of fat embolism syndrome in boys with DMD and acute fracture or trauma to the lower extremities.47,25 Boys with fat embolism syndrome present with altered mental status, respiratory distress, and tachycardia, which should prompt immediate medical attention due to high morbidity and mortality associated with this condition. Current treatment focuses on supportive respiratory care and high-dose corticosteroids.35

**Early non-ambulatory stage**

Foot and ankle surgery to improve equinovarus foot might help with foot positioning in the wheelchair or for shoe wear, but is typically done only if a patient requests the procedure. After foot and ankle surgery, use of ankle-foot orthoses will be needed during the daytime to prevent a recurrence of the contractures.

Inspection of the spine should be part of every clinical examination. Experienced clinicians should be able to monitor the spine in non-ambulatory boys by visual inspection alone; however, less experienced clinicians should obtain a spine radiograph when a child first becomes non-ambulatory. A spine radiograph is also useful when inspection is unhelpful, such as in children with obesity. Once a curve has been detected with radiography, further surveillance depends on the skeletal maturity of the individual: skeletally immature individuals should undergo radiographs once every 6 months, and skeletally mature individuals should undergo radiographs at least once a year. A curve of 20° or more should warrant involvement of an orthopaedic surgeon. The use of spinal orthoses is not recommended. In contrast to the typical clinical course in untreated boys, patients treated with corticosteroids have milder spinal curvatures and less frequent need for spinal surgeries.8,7,10,31

Despite an absence of RCTs, we advise posterior spinal fusion in young men with DMD, given the positive effect on function, sitting balance and tolerance, pain, and quality of life observed in non-randomised, prospective cohort studies.28,30,35 Posterior spinal instrumentation and fusion are recommended in non-ambulatory individuals who have a spinal curve in the sitting position greater than 20–30°, who have not yet reached puberty, and who have not been treated with corticosteroids because the curve is expected to progress. Although patients treated with corticosteroids can still develop scoliosis, the progression might be less predictable, so observation for clear evidence of progression is a reasonable approach before intervening. An anterior spinal fusion approach is not required as the fusion is generally done in the second decade when little additional longitudinal spine growth is anticipated.

When surgical correction for scoliosis is done, stabilisation into the pelvis and fusion are advised in those with a pelvic obliquity of greater than 15° to assist with seating and positioning. In those without a severe pelvic obliquity, fusion to the lower lumbar vertebra is sufficient. The goal of surgical intervention for the spine is to prevent further progression of scoliosis, improve sitting tolerance, and reduce pain.38

Anticipatory fracture prevention guidance should continue through the non-ambulatory stages (figure 5). A more conservative approach to management of lower-limb fractures is advised in non-ambulatory children because the goal is no longer to bear weight. Internal fixation might be necessary for an unstable fracture, but splinting might be sufficient for bone healing and pain control. Pain management is important for all children, but special monitoring could be necessary in the setting of pulmonary and cardiac compromise. Health-care providers and families should be aware of fat embolism syndrome, as described above.

**Late non-ambulatory stage**

Surgical interventions to manage contractures involving the upper or lower extremities are not recommended during the late non-ambulatory stage of DMD unless pain, positioning, or skin integrity is the concern.
Clinicians should examine the spine at every clinical visit. Individuals with known scoliosis should have yearly anteroposterior upright spinal radiographs when there is any concern about progression. Posterior spinal fusion is recommended during the late non-ambulatory stage for those with a progressive curve. It is essential to consult with the patient’s respiratory physician and cardiologist to ensure that lung and heart function are sufficient to proceed with this surgical intervention. Some studies indicate that spinal fusion slows the progression of respiratory decline, whereas others show no significant difference in the rate of decline postoperatively.26–30

The treatment of an acute fracture during the late non-ambulatory stage is similar to that in the early non-ambulatory stage, with the goals of fracture stabilisation and pain control. Cast or splint management is usually sufficient in the setting of a distal femoral metaphyseal fracture. In the case of a proximal femur fracture, operative stabilisation is necessary. As with any fracture, providers and families should be aware of fat embolism syndrome.

**Surgical considerations**

Important surgical considerations for individuals with DMD are detailed in figure 6. Young men with DMD are at risk of potentially fatal rhabdomyolysis and hyperkalaemia when exposed to inhalational anaesthetics or when given suxamethonium chloride (succinylcholine).109 A cardiologist and respiratory physician should be consulted before all surgical procedures, and anaesthetists should be aware that individuals with DMD are at risk of cardiac and respiratory decompensation during and after surgery.110 A detailed discussion of surgical considerations is provided in the appendix.

**Conclusions and future directions**

Improved approaches to respiratory, cardiac, bone health and osteoporosis, and orthopaedic and surgical management can now be offered to children and adults with DMD. However, despite advances in our knowledge and understanding of best approaches to management, progress is needed across these subspecialties to meet the needs of patients.

For respiratory management, diagnostic tools and measures that might have clinical relevance but need further study include assisted cough peak flow, maximum insufflation capacity, the difference between maximum insufflation capacity and FVC, supine FVC, highest flow generated during an inspiratory FVC manoeuvre, the rapid shallow breathing index, and sniff nasal inspiratory pressure. Therapies for which research is needed to establish efficacy and optimum use include high-frequency chest oscillation, intrapulmonary percussive ventilation, and negative-pressure ventilation. Improved understanding of pulmonary phenotypic variability and of the effect of cardiac function and nutritional status on the respiratory system is needed to optimise care and to develop pulmonary outcome measures to assess the efficacy of current and emerging DMD therapies. Prospective studies are needed to assess the criteria recommended in this document for initiation of cough assistance and non-invasive ventilation, with use of clinically relevant outcome measures to develop evidence-based guidelines.

Cardiac outcomes should be included in clinical trials because survival will not improve if emerging therapies do not effectively treat DMD cardiomyopathy. Biomarkers

<table>
<thead>
<tr>
<th>Cardiac care</th>
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<tbody>
<tr>
<td>A cardiologist should be consulted before all surgical procedures</td>
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<tr>
<td>Anaesthetists should be aware that patients with DMD are at risk of cardiac decompensation during surgery</td>
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<tr>
<td>Major surgical procedures</td>
</tr>
<tr>
<td>• Patients with DMD are at particular risk of cardiac compromise during major procedures</td>
</tr>
<tr>
<td>• Echocardiogram and electrocardiogram should be done in close proximity to any planned surgery</td>
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<tr>
<td>Minor surgical procedures</td>
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<tr>
<td>• In patients with normal cardiac function, a cardiac assessment is suggested if last investigation was &gt;1 year before</td>
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<table>
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<tr>
<th>Respiratory care</th>
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<tbody>
<tr>
<td>Preoperative training in and postoperative use of assisted cough techniques</td>
</tr>
<tr>
<td>• Cough techniques are necessary for patients with baseline peak cough flow &lt;270 L/min or baseline maximum expiratory pressure &lt;60 cm H2O*</td>
</tr>
<tr>
<td>Preoperative training in and postoperative use of non-invasive ventilation</td>
</tr>
<tr>
<td>• Non-invasive ventilation is necessary for patients with baseline FVC &lt;30% predicted</td>
</tr>
<tr>
<td>• Non-invasive ventilation is strongly recommended for patients with FVC &lt;50% predicted</td>
</tr>
<tr>
<td>Extubation to supplemental oxygen alone without concomitant use of non-invasive ventilation should be avoided</td>
</tr>
<tr>
<td>Incentive spirometry is not indicated because it is potentially ineffective in patients with respiratory muscle weakness, and preferred alternatives are available</td>
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<tr>
<th>Anaesthesia</th>
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<tbody>
<tr>
<td>Total intravenous anaesthesia is strongly recommended</td>
</tr>
<tr>
<td>Degalvanising muscle relaxants, such as suxamethonium chloride, are absolutely contraindicated because of risk of fatal reactions</td>
</tr>
<tr>
<td>Risk of rhabdomyolysis and hyperkalaemia</td>
</tr>
<tr>
<td>• Patients with DMD are at risk of developing rhabdomyolysis with inhalational anaesthetics or when given suxamethonium chloride</td>
</tr>
<tr>
<td>• Rhabdomyolysis complications are frequently confused with malignant hyperthermia</td>
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<tr>
<th>Blood loss</th>
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</thead>
<tbody>
<tr>
<td>Hypotensive anaesthetics to minimise blood loss are not recommended because of haemodynamic risk with cardiomyopathy in patients with DMD</td>
</tr>
<tr>
<td>Cell-saver technology, along with use of aminoacaric acid or tannic acid, can be considered to help manage intraoperative blood loss</td>
</tr>
<tr>
<td>Postoperative antiocoagulation with heparin or aspirin is not appropriate for patients with DMD</td>
</tr>
<tr>
<td>Compression stockings of sequential compression might be indicated for prevention of deep-vein thrombosis</td>
</tr>
</tbody>
</table>

Figure 6: Surgical considerations for patients with Duchenne muscular dystrophy DMD=Duchenne muscular dystrophy. FVC=forced vital capacity. *Guidance applies to older teenage and adult patients.
that indicate short-term attenuation of disease-related progression need to be identified. Novel dystrophin-specific cardiac treatments are needed to improve patient outcomes. The natural history of cardiomyopathy in female carriers of a disease-causing mutation needs to be clarified, and studies are needed to identify the best diagnostic and therapeutic strategies for affected girls and women.

Because vertebral fractures are an early manifestation of bone fragility and the adverse effects of glucocorticoids occur rapidly, longitudinal trials addressing osteoporosis prevention should originate with young patients, with vertebral fractures as a key outcome measure. Further studies are also needed to assess the potential of growth-promoting therapies to prevent bone fragility and of anabolic agents, such as parathyroid hormone or alendronate, to treat osteoporosis.

Controlled trials of surgical techniques for orthopaedic management, when appropriate, are needed, as is a better understanding of musculoskeletal complications and of the best outcome measures to assess musculoskeletal effects of available and emerging DMD therapies. More studies of patient-reported and family-reported outcomes would help to guide decision making about lower-extremity surgeries and spinal fusion.

Contributors
DJB, KB, CM, BAA, SDA, AB, LEC, IC, SH, AKO, DWS, JB, DRW, and LMW provided intellectual expertise in the study design, generation and interpretation of data, review of the literature, writing of the article, and the decision to publish. DJB, aided by CM, drafted and edited the article and approved the final version.

Declaration of Interests
DJB was a paid consultant for Hill-Rom Corporation and has US patents (8651107, 8844510, and 9795728) for respiratory devices, as well as related international patents and patent applications. KB was a consultant for Solid Ventures, Catabasis, LGC Ltd, Bristol Myers Squibb, PTC therapeutics, GLC Research, Eli Lilly, and Publicis Life Brands Resolute; she has received grant support from PTC Therapeutics. SDA is a principal investigator for multicentre clinical trials sponsored by PTC Therapeutics and Sarepta Pharmaceuticals. LEC has received personal fees for speaking and participating in research supported by Genzyme Corporation of Sanofi; she has participated in research with CNIRG (Cooperative International Neuromuscular Research Group), Enobia Pharma Inc/Alexion, Robertson Foundation, GlassSmithKline, Eli Lilly, Valeron, Pfizer, Prosensa, BioMarin, Ionis, Ultrasynex, Roivant Sciences, Therapeutic Research in Neuromuscular Disorders Solutions, NS Pharma, and the Marcus Foundation. DRW is a paid consultant for Health Research Inc and Marathon Pharmaceuticals. LMW has received grant support and honoraria from Novartis and Amgen. All other authors declare no competing interests.

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