Long-term management of children with neuromuscular disorders
Eugen-Matthias Strehle*

Abstract

Objective: Duchenne muscular dystrophy is the commonest genetic myopathy but there exist a large number of inherited neuromuscular diseases which individually are very rare and where clinical information is not widely available. This review is based on the author’s experience in a pediatric muscle clinic and provides practical guidance and treatment plans for frequently encountered problems.

Sources: A MEDLINE search was conducted to retrieve recent articles relevant to the management of children with inherited myopathies and neuropathies. A patient cohort (n = 200) was evaluated using descriptive statistics.

Summary of the findings: Duchenne muscular dystrophy accounted for almost half of the diagnoses, followed by spinal muscular atrophy (12%), Becker muscular dystrophy and myotonic dystrophy (7% each). Sixteen patients (9%) had an unknown myopathy.

Conclusions: As with other chronic illnesses, these patients should be regularly reviewed by health professionals from an early age to increase life expectancy and improve quality of life. It is useful for physicians to take a structured approach when looking after children with neuromuscular disorders and to monitor all affected organ systems.


Introduction

Inherited neuromuscular disorders (NMDs) affect approximately one in 3,500 children worldwide, and X-linked Duchenne muscular dystrophy (DMD) has the highest incidence among them. Many conditions present with muscle weakness and wasting and can be diagnosed by genetic mutation analysis alone, but muscle, skin or nerve biopsy still remain important diagnostic tools. Although various strategies for genetic therapies of these diseases are being explored, the cornerstones of current treatment are symptomatic and supportive healthcare.\(^1\)\(^,\)\(^2\) Comprehensive reviews on specific NMDs have been published but it can be a challenge for the novice or the busy clinician to keep up to date with the latest literature when looking after children suffering from a wide range of rare disorders.\(^3\)\(^-\)\(^5\) However, when monitoring these chronically ill patients over a prolonged period, certain recurring themes can be observed that are pertinent to most NMDs. These themes can be ascribed to the specific muscle or nerve pathology and its effect on other systems of the body. Here, the author describes his experience of following up 200 children presenting to a pediatric muscle clinic during a 3-year period.

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Clinics

The young patients were reviewed on average twice a year by a multidisciplinary team consisting of three to five doctors (at least one of them senior), one to three specialist physiotherapists, two pediatric nurses, one genetic counselor and one social care advisor. A psychologist would have been desirable but was not routinely available. To reduce travelling for the families, one of the appointments was scheduled at the regional genetics center and the other at a district general hospital close to their homes. This meant that the pediatric muscle team held regular outreach clinics covering on average 100 miles in a day. The age of the predominantly male patients ranged from 1.9 to 19.8 years with a mean age of 11.5 years (Figure 1). The children comprised 44% of all young patients reviewed during that period, and only a small proportion was seen by the author twice. Almost half of the patients (45%) had DMD, followed by spinal muscular atrophy (SMA), Becker muscular dystrophy (BMD) and congenital myotonic dystrophy (MD). A relatively large proportion of children (9%) had an unknown myopathy where a specific diagnosis had not been reached despite extensive investigations. In some cases, a diagnosis was made after several years in the light of new research findings. Hospital database entries were incomplete for 30 patients (15%).

The remaining 170 patients presented with 18 different diagnoses (Table 1).

Systems

Muscular

At each 1-hour assessment a detailed medical history with emphasis on mobility covering the preceding months was obtained, and any concerns raised by the parent/caregiver or patient were addressed. Muscle strength was measured using the Medical Research Council grading: 1 = flicker or trace of contraction, 2 = active movement with gravity.

Table 1 - Spectrum of neuromuscular diseases encountered in a pediatric muscle clinic (n = 170)

<table>
<thead>
<tr>
<th>Disease</th>
<th>n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duchenne muscular dystrophy</td>
<td>77 (45)</td>
</tr>
<tr>
<td>Spinal muscular atrophy</td>
<td>21 (12)</td>
</tr>
<tr>
<td>Unknown myopathy</td>
<td>16 (9)</td>
</tr>
<tr>
<td>Becker muscular dystrophy</td>
<td>12 (7)</td>
</tr>
<tr>
<td>Myotonic dystrophy</td>
<td>11 (7)</td>
</tr>
<tr>
<td>Merosin-negative congenital muscular dystrophy</td>
<td>5 (3)</td>
</tr>
<tr>
<td>Bethlem myopathy</td>
<td>4 (2)</td>
</tr>
<tr>
<td>Central core disease</td>
<td>4 (2)</td>
</tr>
<tr>
<td>Charcot-Marie-Tooth disease</td>
<td>4 (2)</td>
</tr>
<tr>
<td>Nemaline myopathy</td>
<td>3 (2)</td>
</tr>
<tr>
<td>Ullrich congenital muscular dystrophy</td>
<td>3 (2)</td>
</tr>
<tr>
<td>Facioscapulohumeral muscular dystrophy</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Fukuyama congenital muscular dystrophy</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Walker-Warburg syndrome</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Autosomal recessive limb girdle muscular dystrophy</td>
<td>1 (1)</td>
</tr>
<tr>
<td>Myotubular myopathy</td>
<td>1 (1)</td>
</tr>
<tr>
<td>Rigid spine muscular dystrophy</td>
<td>1 (1)</td>
</tr>
<tr>
<td>X-linked myopathy with excessive autophagy</td>
<td>1 (1)</td>
</tr>
</tbody>
</table>
eliminated, 3 = active movement against gravity, 4 = active movement against gravity and resistance, 5 = normal power. As a minimum timed tests for rising from the floor, running 10 m and climbing four stairs up and down were performed on all ambulant children. Boys with DMD also underwent a North Star Ambulatory Assessment which measures levels of activity and gives a maximum score of 34.6 Non-ambulant children were assessed using the Egen Classification Scale Version 2.7 Most children received regular physiotherapy, and their parents were reminded of the importance of daily home stretching exercises to prevent contractures. Physical activity that was highly competitive or carried an increased risk of injury was not recommended. If ankle dorsiflexion was not possible beyond the neutral position, night splints were prescribed for ambulant children and day splints for those in wheelchairs. Manual and electric wheelchairs were regularly maintained to ensure a comfortable sitting position. Some young people with DMD benefitted from a standing frame, swivel walker or knee-ankle-foot orthosis (KAFO). A recent Cochrane review provided evidence that oral steroids improve muscle strength and function in DMD for up to 5 years.9,10 Accordingly, the patients reviewed here were treated with prednisolone 0.75 mg/kg/day or deflazacort 0.9 mg/kg/day, once their gross motor skills had reached a plateau or started to decline, which usually coincided with admission to primary school (age 4-6 years). The doses were starting doses that were not regularly adjusted with changes in body weight but were altered depending on the clinical picture. The maximum daily dose for either drug was 40 mg. The steroid treatment was generally well tolerated, and the observed adverse effects included increased appetite, weight gain, behavioral problems, high blood pressure, stunted growth, osteoporosis, and cataracts. Prior to commencing treatment, all patients had a blood screen including full blood count, biochemical profile, vitamin D level, and varicella antibodies. If appropriate, children were immunized against chickenpox. A family history of active tuberculosis was a contraindication for steroid treatment of DMD. A steroid alert card was issued to every child reminding health professionals of the possibility of drug-induced adrenal insufficiency. At each clinic visit, weight, height, blood pressure, and forced vital capacity (FVC) were measured, and urine was tested for glucose, protein, and blood. Although no patient with a myasthenic syndrome was among this cohort, the therapeutic options for these rare disorders affecting the neuromuscular junction should be mentioned. Pyridostigmine bromide, 3,4-diaminopyridine and ephedrine sulphate are effective in non-immune congenital myasthenic syndromes, whereas acetyl cholinesterase inhibitors, immunosuppressant drugs, and thymectomy are the treatment of choice for juvenile autoimmune myasthenia gravis.11,12 Recent studies suggest a positive effect of oral salbutamol 2 mg thrice daily on muscle power in patients with SMA but the drug was not routinely used in this group.13,14

**Skeletal**

Despite regular physiotherapy and preventative splinting, joint contractures were frequently observed in children with NMDs, and in some they were present from birth (e.g. congenital muscular dystrophy). The commonest type was an ankle-foot contracture which was treated with serial casting or Achilles tendon release by an orthopedic surgeon.15,16 At each appointment, the patient’s spine was examined for abnormal curvatures (hyperlordosis, scoliosis, and kyphosis) and vertebral fractures. There was a low threshold for requesting spinal X rays or referring to a spinal surgeon for a baseline assessment. Conservative treatment of a scoliosis included a spinal brace, and in severe cases spinal fixation and fusion surgery were performed.17-19 Osteopenia was common in boys with DMD, and the added effect of oral steroids increased the incidence of osteoporosis and vertebral and other fractures.20,21 The bone mineral density of these children was measured by dual-energy X-ray absorptiometry (DEXA scan) every 2 years. Lately, patients commenced on prednisolone or deflazacort were also prescribed the oral bisphosphonate risedronate at a dose of approximately 1 mg/kg once weekly (maximum dose 35 mg once weekly) and a combined calcium and vitamin D preparation on 6 days per week.22,23 DMD patients on steroids diagnosed with a fracture were treated as inpatients with intravenous infusions of pamidronate. Twice-yearly dental checkups were recommended as osteonecrosis of the jaw (ONJ) is a recognized complication of bisphosphonate therapy.24,25 In children with joint hypermobility, heritable disorders of connective tissue, such as Ehlers-Danlos syndrome, were considered.

**Nervous**

Cognitive impairment of varying severity and neuropsychiatric disorders were frequently seen in patients with dystrophinopathies, MD, and other inherited muscle diseases.26-28 When assessing young infants, it was important to differentiate between those who had poor truncal but good limb tone (central hypotonia) and those who were floppy and weak (peripheral hypotonia). In children with unknown neuropathy or myopathy, neuroradiological studies and peripheral neurophysiological tests were sometimes helpful. Associated seizures were controlled with standard anti-epileptic drugs (e.g. sodium valproate, carbamazepine). Therapeutic sleep systems and melatonin 3-6 mg at night were prescribed for patients with sleeping difficulties. Occasionally, adolescents with MD and excessive daytime sleepiness were treated with modafinil 100 mg once or twice daily.29,30 Children with MD and those on long-term steroids were screened for small cataracts annually by an optician or ophthalmologist. Sensorineural deafness was rare but has been reported in MD, facioscapulohumeral muscular dystrophy, and Charcot-Marie-Tooth disease.31 Malignant hyperthermia is a serious condition associated
with general anesthesia and characterized by muscle rigidity, rhabdomyolysis, acidosis, myoglobinuria, and hyperthermia. Patients with a wide range of myopathies can be affected and they were made aware of this risk prior to undergoing surgery. When patients complained of pain, their symptoms were analyzed according to main site, radiation, character, severity, duration, and frequency. In young children, a simple pain rating scale was used. If an underlying pathology could be identified, specific treatment was initiated. Leg pain and cramps were frequently seen in children with NMDs and often responded to massage, paracetamol, ibuprofen, a mechanical compression device (Flowtron®) or transcutaneous electrical nerve stimulation (TENS). A significant number of children had non-organic or psychosomatic pain.

Cardiovascular

As cardiac involvement is common in children with NMDs, all patients had a baseline cardiovascular assessment at the time of diagnosis. Boys with DMD and teenagers with BMD had annual echocardiograms to check for early signs of cardiomyopathy. Regular electrocardiograms were recommended for patients with MD and Emery-Dreifuss muscular dystrophy to detect conduction defects that may necessitate anti-arrhythmic drug therapy or a pacemaker. Once the left ventricular ejection fraction started to decrease significantly in DMD patients, treatment with a beta-adrenoceptor blocker (e.g. bisoprolol) and an angiotensin-converting enzyme inhibitor (e.g. perindopril) was commenced. Occasionally, long-term use of oral steroids led to moderate hypertension. If this was suspected following one abnormal reading, the blood pressure was measured repeatedly on different days using a sphygmomanometer with a cuff covering at least 2/3 of the child's upper arm. Antihypertensive therapy was rarely required and usually initiated by a pediatric nephrologist without stopping the steroids.

Respiratory

Children with NMDs, particularly those with DMD and SMA, are prone to develop chest infections, which is due to their weak respiratory muscles and their reduced mobility. It was therefore recommended that they should have the pneumococcal and the seasonal influenza vaccines. Lower respiratory tract infections were treated aggressively with oral or intravenous antibiotics. At each clinic visit, the young patients' lung function was measured using an electronic spirometer with a printer. If symptoms of nocturnal hypoxia and hypercapnia were reported, for instance headaches and tiredness in the morning, an overnight pulse oximetry study was arranged which allowed continuous recording of oxygen saturations and heart rate, and early recognition of desaturations. When a patient's coughing capacity was significantly reduced, a mechanical insufflation/exsufflation device (CoughAssist®) was prescribed to help removing secretions from the lungs. A steady decline in the FVC (absolute and percent predicted) indicated the need for a referral to the home ventilation team and, usually in the second decade of life, initiation of non-invasive positive pressure ventilation (NIPPV) via a nasal mask, a face mask or a mouthpiece to prevent respiratory failure. A tracheostomy was rarely required.

Gastrointestinal

At each consultation, a dietary history was obtained including appetite, length of meal times and difficulties with chewing and swallowing. If recurrent choking episodes were reported, the patient was referred to a specialist speech and language therapist for a formal feeding assessment and a videofluoroscopy study. Parents were advised to take their children for regular dental appointments to recognize caries and gingivitis early. The importance of a balanced, high-fiber diet was emphasized to improve general well-being and facilitate regular bowel function. Constipation was fairly common and usually responded to lactulose, macrogols, senna or docusate sodium. Gastroesophageal reflux was treated with aluminum hydroxide and magnesium carbonate or ranitidine. Excessive weight gain was frequently observed in children receiving oral steroids, particularly those on prednisolone, who developed cushingoid features. Some young patients with a NMD did not show adequate weight gain despite input from a dietician and therefore required nasogastric tube feeding or a gastrostomy.

Urogenital

Children with NMDs are prone to dehydration due to their restricted mobility, which can lead to renal impairment. Myoglobinuria is another complication of inherited myopathies and can result in renal failure, if untreated. The protein myoglobin is released from damaged muscle fibers and causes brown discoloration of the urine which tests positive for blood on dipstick analysis. Older children were advised to avoid excessive exercise and encouraged to drink plenty of fluids, particularly during the summer months. Nocturnal and diurnal enuresis was treated with desmopressin or oxybutynin in children who did not have associated global developmental delay. Symptoms of a urinary tract infection were investigated promptly and managed according to current guidelines.

Conclusion

Although the timely management of medical problems had priority in the children followed up here, there were other issues such as education and social integration that contributed to the quality of life of the patients and their families. Most children with NMDs attended a nursery or...
(special) school during the day and had a statement of special educational needs. Their treatment and any adaptations made to the family homes were funded by the National Health Service and local social services departments. Some children were offered respite care in a hospice. The transition period from adolescence to adulthood often lasts longer in young people with a chronic disease or disability, and a number of patients were seen in the pediatric muscle clinic beyond the British upper age limit of 16 years. In a recent survey, adult physicians in the United States expressed concern about their lack of training in childhood-onset conditions which may hinder a smooth transition of care for affected adolescents. This problem will not occur, if children and adults with NMDs receive their specialist care by the same multidisciplinary team throughout their lives as is practice in some centers in Great Britain. Where this solution is not feasible, additional postgraduate training for doctors is required specifically addressing transition care. Finally, there remains hope that a few of the novel therapeutic agents that are currently tested in multicenter trials will become a successful treatment for one or more neuromuscular diseases in the future.59

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References


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