CONDITIONS WE SUPPORT AT MDNSW

Welcome to Muscular Dystrophy NSW. The support community for people with neuromuscular conditions.

With a name like MDNSW, you'd be forgiven for thinking that we only support people with Muscular Dystrophy.

The good news is - we're actually here for people with **all** neuromuscular conditions.

Which means you can feel confident that we understand your condition, and that we've supported others with your condition before.

After 60 years of serving the neuromuscular community, we're here to support you with the tailored services and community you deserve.

Here's how to find your condition on this list, in 3 easy steps:

1. Click **CTRL + F**, and type in the name of your condition (or just scroll through the list)
2. Find your condition
3. Get in contact with us, to find out how we can support you today

Can't find your condition listed? We'd still love to chat. Just get in contact and we'll help you find tailored support for your condition.

NEUROMUSCULAR CONDITIONS WE SUPPORT AT MDNSW

Anterior Horn Cell Disease
Becker MD
Carnitine Deficiency (CD)
Carrier DMD Gene
Central Core Disease (CCD)
Centro-Neuclear Myopathy
Charcot-Marie-Tooth Disease (CMT) ▪ Type IV - X-linked Charcot-Marie-Tooth Disease ▪ Congenital Hypomyelinating Neuropathy ▪ Dejerine-Sottas disease ▪ Hereditary Motor and Sensory Neuropathy
CIDP (Chronic Inflammatory Demyelinating Peripheral Neuropathy)
Congenital Muscular Dystrophy ▪ Bethlem myopathy ▪ Fukuyama CMD (Dystroglycanopathy) ▪ Laminin a2 related dystrophy ▪ Merosin-deficient CMD ▪ Muscle-eye-brain disease (Dystroglycanopathy) ▪ SEPN1 related myopathy or rigid spine muscular dystrophy ▪ Ullrich congenital muscular dystrophy ▪ Walker-Warburg Syndrome (Dystroglycanopathy)
Congenital Myasthenic Syndrome (CMS)
Distal MD (DD)

NEUROMUSCULAR CONDITIONS WE SUPPORT AT MDNSW

Duchenne MD
Emery-Dreifuss MD
FacioScapuloHumeral MD
Fibre-Type Disproportion
Friedreich's Ataxia (FA)
Glycogen Storage Disease: Type VI - Phosphfructokinase Deficiency (Tarui's Disease)
Idiopathic Inflammatory Myopathies (IIM) ▪ Dermatomyositis ▪ Polymyositis
Inclusion Body Myositis (IBM)
Kearns Sayer Syndrome
Limb Girdle Muscular Dystrophy (LGMD) ▪ LGMD1A ▪ LGMD1B ▪ LGMD1C ▪ LGMD2A
Mini / Multi Core Disease
Mitochondrial Myopathy (MITO)
Miyoshi Distal Myopathy
Multi Core Myopathy
Muscular Dystrophy of Late Onset

NEUROMUSCULAR CONDITIONS WE SUPPORT AT MDNSW

Myasthenia Gravis (MG) Juvenile & Adult
Myotonia Congenita (MC)
Myotonic Muscular Dystrophy
Myotubular (Centronuclear) Myopathy
Occular Myasthasenia
Oculopharyngeal MD (OPMD)
Paramytonia Congenita (PC)
Polymyositis (PM)
Pompe Disease
Spinal Muscular Atrophy ▪ Spinal Muscular Atrophy Type I ▪ Spinal Muscular Atrophy Type 2 ▪ Spinal Muscular Atrophy Type 3 ▪ Spinal Muscular Atrophy Type 4
Spino Cerebellar Ataxia
Ullrich Congenital MD
Unknown/Undiagnosed MD Type
Other

Helping you get the most out of your NDIS plan!

MUSCULAR DYSTROPHY NSW

YOUR NEUROMUSCULAR SUPPORT COMMUNITY

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Phone: **(02) 9888 5711**