



Muscular Dystrophy Ireland

Membership Application Form



General Details

Name			Date	
Address				
Telephone Numbers	<i>Home</i>	<i>Mobile</i>	<i>Work</i>	
	From time to time MDI would like to send you text messages about fundraising, social and other events which you may be interested in. If you would like to receive these text messages please tick this box: <input type="checkbox"/>			
Email				

Type of Membership & Annual Subscription Rates

Please tick whichever of the following applies to you

Person with MD
€15.00

Family Membership
€20.00

Friend of MDI
€15.00

If you would like to make a donation additional to the membership fee, please indicate the amount here:	€
Total amount enclosed with this form:	€
Would you like a receipt?	Yes <input type="checkbox"/> No <input type="checkbox"/>

MDI Database

To enable us to update our records and maintain accurate information, we would appreciate it if you would provide the following information regarding your family if applicable:

Note: When specifying “Type of MD” please see list overleaf and tick appropriate. Thank you.

Name(s) of Person(s) with MD	Date(s) of Birth	Type of MD

All cheques and postal orders should be made payable to Muscular Dystrophy Ireland.

It is also possible to pay your subscription by Standing Order to our account;
Muscular Dystrophy Ireland Savings Account at Bank of Ireland, 2 College Green, Dublin 2.

BIC: BOFIE2D **IBAN:** IE11BOFI90001775261619

Please complete this form and return it, with the relevant membership fee (see above) to:
Muscular Dystrophy Ireland, 75 Lucan Road, Chapelizod, Dublin 20. Thank You.

Types of Muscular Dystrophy

Each year MDI organises Information Days on specific types of muscular dystrophy. If you have a form of muscular dystrophy, we would be grateful if you could **specify the exact type by ticking the appropriate box below**. This will insure that you are invited to appropriate MDI Information Days and provided with relevant available updates and news on research and treatments for your specific condition. The conditions listed below are covered by MDI. For more information on a specific condition, visit: <http://www.mdi.ie/types-of-muscular-dystrophy.html>. If you are in any doubt over whether MDI covers the condition you or someone you know has, please do not hesitate to contact our Information Officer by emailing info@mdi.ie or phoning (01) 6236414.

Please tick appropriate box which applies to you or your family member(s):

1. Muscular Dystrophies

Becker muscular dystrophy	<input type="checkbox"/>	Facioscapulohumeral muscular dystrophy	<input type="checkbox"/>
Duchenne muscular dystrophy	<input type="checkbox"/>	Limb-girdle types of muscular dystrophy (LGMD) - General	<input type="checkbox"/>
Manifesting carrier of Duchenne	<input type="checkbox"/>	• LGMD 1B (also known as Laminopathy)	<input type="checkbox"/>
Congenital muscular dystrophy - General	<input type="checkbox"/>	• LGMD 1C (also known as Caveolinopathy)	<input type="checkbox"/>
• MDC1A (merosin-deficient congenital muscular dystrophy)	<input type="checkbox"/>	• LGMD 2A (also known as Calpainopathy)	<input type="checkbox"/>
• Rigid spine syndrome (RSS)	<input type="checkbox"/>	• LGMD 2B (also known as Dysferlinopathy)	<input type="checkbox"/>
• Ullrich congenital muscular dystrophies	<input type="checkbox"/>	• LGMD 2I	<input type="checkbox"/>
• Bethlem myopathy	<input type="checkbox"/>	Ocular myopathies including ocularopharangeal muscular dystrophy	<input type="checkbox"/>
Emery-Dreifuss muscular dystrophy	<input type="checkbox"/>		

2. Myotonic Disorders

Congenital Myotonic Dystrophy	<input type="checkbox"/>
Myotonia	<input type="checkbox"/>
Myotonic Dystrophy	<input type="checkbox"/>

3. Congenital Myopathies

Central Core Myopathy	<input type="checkbox"/>
Congenital fibre-type disproportion myopathy	<input type="checkbox"/>
Minicore (Multicore) myopathy	<input type="checkbox"/>
Myotubular or Centronuclear myopathy	<input type="checkbox"/>
Nemaline myopathy	<input type="checkbox"/>

4. Mitochondrial Myopathies

	<input type="checkbox"/>
--	--------------------------

5. Metabolic Disorders

Metabolic disorders (general)	<input type="checkbox"/>
McArdle's Disease	<input type="checkbox"/>
Pompe's Disease	<input type="checkbox"/>

6. Periodic Paralyses

	<input type="checkbox"/>
--	--------------------------

7. Autoimmune Myositides

Polymyositis, Dermatomyositis and Sarcoid myopathy	<input type="checkbox"/>
Juvenile dermatomyositis	<input type="checkbox"/>
Inclusion body myositis	<input type="checkbox"/>

8. Spinal Muscular Atrophies

Severe (Type I)	<input type="checkbox"/>
Intermediate (Type II)	<input type="checkbox"/>
Mild (Type III)	<input type="checkbox"/>
Adult spinal muscular atrophy	<input type="checkbox"/>

9. Hereditary Motor and Sensory Neuropathies

(Also known as Charcot-Marie-Tooth or Peroneal muscular atrophy)	<input type="checkbox"/>
--	--------------------------

10. Disorders of the Neuromuscular Junction

Congenital myasthenic syndromes	<input type="checkbox"/>
Myasthenia Gravis	<input type="checkbox"/>

11. Friedreich's Ataxia

	<input type="checkbox"/>
--	--------------------------

12. Other (Please Specify)

--