

About motor neurone disease (MND):

- MND is a fatal, rapidly progressing disease that affects the brain and spinal cord.
- It attacks the nerves that control movement so muscles no longer work. MND does not usually affect the senses such as sight, sound and feeling etc.
- It can leave people locked in a failing body, unable to move, talk and eventually breathe.
- Over 80% of people with MND will have communication difficulties, including for some, a complete loss of voice.
- It affects people from all communities.
- Around 35% of people with MND experience mild cognitive change, in other words, changes in thinking and behaviour. A further 15% of people show signs of frontotemporal dementia which results in more pronounced behavioural change.
- It kills a third of people within a year and more than half within two years of diagnosis.
- A person's lifetime risk of developing MND is around 1 in 300.
- Six people per day are diagnosed with MND in the UK.
- It affects up to 5,000 adults in the UK at any one time.
- It kills six people per day in the UK, just under 2,200 per year.
- It has no cure.

About the Motor Neurone Disease Association:

The MND Association was founded in 1979 by a group of volunteers with experience of living with or caring for someone with MND. Since then, we have grown significantly, with an ever-increasing community of volunteers, supporters and staff, all sharing the same goal – to support people with MND and everyone who cares for them, both now and in the future.

We are the only national charity in England, Wales and Northern Ireland focused on MND care, research and campaigning.

Our mission

We improve care and support for people with MND, their families and carers.

We fund and promote research that leads to new understanding and treatments, and brings us closer to a cure for MND.

We campaign and raise awareness so the needs of people with MND and everyone who cares for them are recognised and addressed by wider society.

Our vision

A world free from MND

PLEASE NOTE: motor neurone disease is lower case and then abbreviated to MND. The Motor Neurone Disease Association can be shortened to the MND Association or the Association, but never the MNDA.

Media Team:

We offer journalists, influencers and broadcasters' access to expert information about MND and the MND Association.

We can organise interviews with people affected by the disease, and regional and national spokespeople covering England, Wales and Northern Ireland, on a wide range of topics including:

- Living with MND
- The latest research into possible causes of MND
- The search for treatments and a cure
- Specialist social and health care affecting people with MND
- Our current campaigns
- Fundraising

For interview, comment or personal story requests, as well as imagery and logo files please:

Email: communications@mndassociation.org

Phone: **01604 611838** (during office hours) or

Mobile: **07831 349409** (for **urgent** out of office media queries)

Further information:

Videos: A collection of short films are available on our website

<https://www.mndassociation.org/videos/> under the following sections:

- Awareness campaigns
- Care for people living with MND
- Campaigning
- Conferences
- Research
- For professionals
- From our supporters

The below provide a starting point:

The Ride

'The Ride' shows the devastating effects of motor neurone disease portrayed as a terrifying rollercoaster ride. https://www.youtube.com/watch?v=2668TvHs_dw&feature=youtu.be

What is MND? A short animation

This animated video provides a simple overview about motor neurone disease.

<https://youtu.be/tq0MO2x31NA>

How can the MND Association help?

This animated video explains how the Association can help families affected by motor neurone disease and explains our services

<https://youtu.be/Rblkgq7jvPw>

