

Muscular Dystrophy UK

Fighting muscle-wasting conditions



go bright
for the fight!

Friday 7 February 2020

A young boy with blonde hair, dressed as a superhero, is the central focus. He is wearing a blue long-sleeved shirt, a bright orange cape, and a matching orange eye mask. He has a determined expression with his mouth open and his right fist raised in a power salute. The background is a bright, orange-toned sky with white clouds and three orange stars of varying sizes.



Our priorities



Research to accelerate the steps to treatments and cures



Improved care and support



Quality of life for independent living

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Every Day Counts


Everyday we're searching for treatments and cures to improve the lives of people today and transform those of future generations.





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**Every day scientists are
finding out more about
muscular dystrophy to
bring us closer to new
treatments.**

EVERY DAY COUNTS



Every Day Counts

Every day people with progressive muscle wasting conditions live with an uncertain future. Currently there is no cure and it gets worse over time. It's the hardest challenge they and the people close to them have to face.

We know how time is precious for people with muscular dystrophy.





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**Together, we're
driving change so
that people can get
out and about when
and where they want.
Count me in.**

EVERY DAY COUNTS

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Every Day Counts

Everyday we're reaching out with information and support to help people live well with their condition and driving change to see the best care and support.



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Every day brilliant people like you are helping to bring forward the day when we beat muscular dystrophy. Make today count.

EVERY DAY COUNTS





For over 50 years, we've been here for families. This is only possible thanks to donations, gifts in Wills and family fundraising.

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Every Day Counts

**THANK YOU
FOR JOINING
IN WITH
GO BRIGHT
FOR THE
FIGHT**

