Impact of Childhood and Congenital DM on Quality of Life
Nicholas Johnson, MD, and fellow researchers at the University of Rochester recently published an article in The Journal of Child Neurology that describes the impact of childhood and congenital myotonic dystrophy on quality of life. Click here to read a summary

Would You Like to Tell Us Your DM Story?
Sharing the unique stories of our community members and how they are meeting the challenge of living with DM is one of MDF’s most important communication goals. If you’d be willing to tell us your personal story, we’d love to hear it. We may contact you about sharing it with the community via the MDF Dispatch, Facebook or our website. Please write to us at info@myotonic.org.

Meet the MDF Team!
Have you ever wondered who’s who at MDF? We’ve recently added people to our team to better support the DM community and help advance research. Here’s a look at who’s doing what!

Registry now over 900!
We’re very close to our goal of registering 1,000 people in the Myotonic Dystrophy Family Registry. The more individuals we have registered and reporting their personal experiences, the more accurate and helpful the Registry will be to researchers and the DM community as a whole. If you or someone you know has DM1 or DM2 and hasn’t yet registered, please click here to do so today!

Like us on Facebook and follow us on Twitter!

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