Help make a difference!

Join the National Registry of Myotonic Dystrophy and FSHD Patients and Family Members







Why join the Registry?

- Receive information about research studies and how to participate
- ➤ Get connected with myotonic dystrophy and FSHD researchers:
 - o Participate in studies
 - Receive newsletters and research updates through mail, email, or social media
 - o Talk to knowledgeable staff
- ➤ Help contribute to knowledge about DM and FSHD by providing yearly updates on your health



About the Registry:

- ➤ Over 2,000 members & growing
- Advancing research for over 14 years
- Free to join- funded by the National Institutes of Health
- > Privacy secured and protected
- ➤ Led by Dr. Rabi Tawil and other experts at the University of Rochester

Call to join!

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