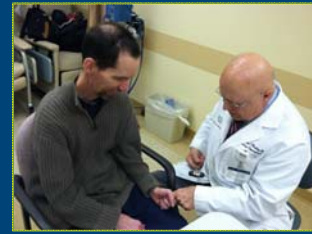


# Help us make a difference!

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



## Why join the Registry?

- Find out about research studies you can participate in!
- Get connected with experts in myotonic dystrophy (DM) and facioscapulohumeral muscular dystrophy (FSHD)!
- Help contribute to knowledge about DM and FSHD by providing yearly updates on your health!
- Help speed up the development of new treatments and prepare for clinical trials!

## About the Registry:

- Over 2,000 members & growing!
- Advancing research for over 12 years.
- Free to join! Funded by the National Institutes of Health.
- Privacy secured and protected.
- Led by Dr. Richard T. Moxley and other experts at the University of Rochester.



## Call to learn more and join!

Jim Hilbert and Liz Luebbe  
Web: [dystrophyregistry.org](http://dystrophyregistry.org)  
Phone (toll-free): 1-888-925-4302  
E-mail: [dystrophy\\_registry@urmc.rochester.edu](mailto:dystrophy_registry@urmc.rochester.edu)

The National Registry  
University of Rochester Medical Center  
601 Elmwood Avenue, Box 673  
Rochester, NY 14642

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- nf 5/1/13-

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