

National Registry of Myotonic Dystrophy and Facioscapulohumeral Muscular Dystrophy Patients and Family Members

The Registry collects information from myotonic dystrophy and facioscapulohumeral muscular dystrophy (FSHD) patients and family members interested in joining the Registry. The Registry reports information on the progression of disease symptoms and helps scientists recruit subjects to participate in research trials.

Once you are enrolled, you will be informed by mail about studies for which you might be eligible. If you are interested in participating in a particular study, you simply contact the researcher listed in the study announcement. The Registry will not release your name to an investigator and you are under no obligation to volunteer for any study.

The Purpose of the Registry is to:

- Help people with Myotonic Dystrophy or FSHD participate in research and learn more about their disease;
- Help scientist accomplish research on Myotonic Dystrophy or FSHD;
- Encourage more research on these diseases

**For more information or to enroll
please contact us:**

Toll Free #: (888) 925-4302

Fax #: (585) 273-1255

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Web site: www.dystrophyregistry.org

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RSRB-University of Rochester-Approval
RSRB No. 12163
Expires September 26, 2010
- jsl 8/30/09 -