



The Myotubular and Centronuclear Myopathy Patient Registry

Why join the registry?

- ✓ **Help** identify patients for research studies or clinical trials.
- ✓ **Support** research projects by providing MTM and CNM data
- ✓ **Encourage** further research into the conditions thanks to the existence of data
- ✓ **Provide** information on living with MTM and CNM to develop standards of care
- ✓ **Receive** updates on relevant research, studies or clinical trials



We welcome the registration of:

- ✓ Patients with a diagnosis of Myotubular or Centronuclear Myopathy, confirmed via genetic testing or muscle biopsy.
- ✓ Female carriers of X-linked MTM.
- ✓ Deceased patients who had a confirmed diagnosis.

If you do not fit these descriptions but would like to receive updates and newsletters, please email us and we will add you to our mailing list.

This registry is looked after by the John Walton Muscular Dystrophy Research Centre at Newcastle University, as part of the TREAT-NMD Neuromuscular Network. Any enquiries can be emailed to the Registry Curator, or visit our website for more information.

STRENGTH IN NUMBERS - REGISTER ONLINE TODAY



www.mtmcnmregistry.org



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