Participants: Patients and their family members diagnosed with a wide variety of neuromuscular disease such as ALS, IBM, CMT.

Background: Many neuromuscular diseases are caused by, or strongly influenced by genetic mutations. In most conditions, fewer than half of causative genes have been identified, limiting our understanding of the disease.

What takes place in study: Participants provide information about their general health, family history, and neurological disease. They may undergo a neurological examination. Participants also provide 2 tubes of blood, which is made into DNA for storage and use in ongoing and future genetics studies.

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