

# Natural History Study for Patients With Nemaline Myopathy in Spain

NCT07488806

---

**Status** RECRUITING  
**Sponsor** Hospital Universitari Vall d'Hebron Research Institute  
**Enrollment** 100 participants

## Key Eligibility Criteria

---

### Inclusion (2)

- Patients with a confirmed clinical and genetic diagnosis of MN (mutations in ACTA1, NEB, TPM2, TPM3, KBTBD13, CFL2, KLHL40, KLHL41, LMOD3, MYPN, TNNT1, TNNT3), or under discussion if they only have a compatible biopsy.
- Signed informed consent by the patient or Legal Authority Responsible, and/or assent by the subject (in pediatric population).

## Locations (1 total)

---

University Hospital Vall d'Hebron, Barcelona, Spain