# SUPPLEMENTARY MATERIAL

## **Table S1:** Classification of neuromuscular disease used in study analysis

|  |  |  |
| --- | --- | --- |
| Group | Sub-group | Sub-group used in analysis |
| Motor Neuron Disorders | Post-Polio syndrome  Spinal muscular atrophy |  |
| Muscle disease | Acquired myopathies  Hereditary myopathies  Mitochondrial disease  Muscle channelopathies  Myotonic disorders (unspecified)1 | Inflammatory myopathies  Muscular dystrophies, Myotonic dystrophy (Type 1) |
| Neuropathy | Hereditary Neuropathies Inflammatory & autoimmune neuropathies | Charcot-Marie Tooth disease  Guillain-Barré syndrome |
| Neuromuscular Junction Disorder | Lambert-Eaton syndrome  Myasthenia gravis  Other | Myasthenia gravis |
| Muscular or neuromuscular disease unspecified2 |  |  |

1 - This category only applies if they cannot be assigned within “Muscular Dystrophy” or “Muscle channelopathies”  
2 - This category only applies if they cannot be assigned to any of the above categories

For Read codes used in the definition please see <https://doi.org/10.24376/rd.sgul.21878271> and also Carey, I. M., et al. (2021). "Prevalence and incidence of neuromuscular conditions in the UK between 2000 and 2019: A retrospective study using primary care data." PloS One 16(12).