**Part 1—Eligible neurological conditions**

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| Category 1 | CONGENITAL MORPHOLOGICAL DISORDERS |
|  | Agenesis of Corpus Callosum |
|  | Anorectal Malformation |
|  | Apert Syndrome |
|  | Arnold-Chiari Syndrome |
|  | Arthrogryposis |
|  | Bladder Exstrophy |
|  | Caudal agenesis |
|  | Caudal Regression Syndrome |
|  | Cerebral Neuronal Migration Disorders |
|  | Charge Syndrome |
|  | Cloacal Exstrophy |
|  | Congenital Epispadias |
|  | Congenital Hydrocephalus |
|  | Dandy-Walker malformation |
|  | Developmental Cord Disorder |
|  | Hirschsprung’s Disease |
|  | Holoprosencephaly |
|  | Imperforate Anus |
|  | Incomplete Corpus Callosum/Aicardi Syndrome |
|  | Lissencephaly |
|  | Megalencephaly |
|  | Microcephaly |
|  | Neural tube defect |
|  | Polymicrogyria |
|  | Pontocerebellar Hypoplasia |
|  | Posterior Urethral Valve Syndrome |
|  | Prune Belly Syndrome |
|  | Sacral Agenesis |
|  | Schizencephaly |
|  | Spinal Agenesis |
|  | Spinal Dysraphism |
|  | Spinal Hemangioma |
|  | Syringobulbia |
|  | Syringomyelia |
|  | Tethered spinal cord |
|  | Vater Syndrome/Vacterl Syndrome |
|  | Velocardiofacial Syndrome |
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| Category 2 | CEREBRAL PALSY |
|  | Dystonic Cerebral Palsy |
|  | Hereditary Spastic Paralysis |
|  | Spastic Quadriplegia |
|  | Mixed cerebral palsy |
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| Category 3 | SYNDROMES ASSOCIATED WITH INTELLECTUAL IMPAIRMENT |
|  | 2-Hydroxyglutaric Aciduria |
|  | Alpers Disease |
|  | Angelman Syndrome |
|  | Alpha Thalassaemia X-linked intellectual disability Syndrome |
|  | Bardet Biedl Syndrome |
|  | Beare-Stevenson Syndrome |
|  | Cyclin Dependent Kinase-Like 5 Gene Mutation |
|  | Chime Syndrome |
|  | Chromosome 1 Deletion |
|  | Chromosome 5q deletion (Cri Du Chat Syndrome) |
|  | Chromosome 13q Deletion Syndrome |
|  | Chromosome 15q Duplication Syndrome |
|  | Chromosome 18q Deletion Syndrome |
|  | Chromosome 1p36 Deletion Syndrome/Mono 1p36 |
|  | Chromosome 22 Ring |
|  | Chromosome 2q Deletion Syndrome |
|  | Chromosome 6 Ring Syndrome |
|  | Chromosome 8  Inversion or Duplication |
|  | Chromosome 9p Deletion Syndrome |
|  | Chromosome 9q Deletion Syndrome |
|  | Chromosome 11q (Jacobsen Syndrome) |
|  | Chromosome Xp Duplication |
|  | Cockayne Syndrome |
|  | Coffin-Lowry Syndrome |
|  | Cognitive Impairment |
|  | Cohen Syndrome |
|  | Congenital disorders of glycosylation |
|  | Congenital Neurological Infections |
|  | Cornelia de Lange Syndrome |
|  | Costello Syndrome |
|  | Cowden Disease |
|  | Developmental Delay |
|  | Developmental Delay associated with Autism, Autism Spectrum Disorder and Aspergers Syndrome |
|  | Dravet Syndrome |
|  | Fragile X Syndrome |
|  | Fumarase Deficiency |
|  | GLUT1-Deficiency Syndrome |
|  | Glutaric Aciduria Type 1 |
|  | Goldenhar’s Syndrome |
|  | Hunter Syndrome |
|  | Hurler-Scheie Syndrome |
|  | Hypomyelination disorders |
|  | Joubert Syndrome |
|  | Kabuki Syndrome |
|  | Langer-Gideon Syndrome |
|  | Lawrence Moon Biedel Syndrome |
|  | Lennox-Gastaut Syndrome |
|  | Lesch-Nyhan Syndrome |
|  | Lowe Syndrome |
|  | Mannosidosis |
|  | Maple Syrup Urine Disease |
|  | Meningitis |
|  | Menkes Syndrome |
|  | Mitochondrial Dieases |
|  | Molybdenum Cofactor Deficiency |
|  | Mowat-Wilson Syndrome |
|  | Mucolipidosis IV |
|  | Myotonic Dystrophy (Type 1) |
|  | Neonatal Hypoxic ischaemic encephalopathy |
|  | Neonatal Onset Multisystem Inflammatory Disease |
|  | Neuronal ceroid lipofuscinosis |
|  | Normal Pressure Hydrocephalus |
|  | OHDO Syndrome |
|  | Opitz Trigonocephaly Syndrome |
|  | Ohtahara Syndrome |
|  | Ouvrier Syndrome |
|  | Pallister-Killian Mosaic Syndrome |
|  | Peroxisome Biogenesis Disorder |
|  | Phelan McDermid Syndrome/22q 13 Deletion Syndrome |
|  | Phenylketonuria |
|  | Prader-Willi Syndrome |
|  | Pyruvate Dehydrogenase Deficiency/Leigh’s Disease |
|  | Rasmussen’s Disease |
|  | Rett Syndrome |
|  | Rubinstein-Taybi Syndrome |
|  | Sensory Integration Disorder/Dysfunction |
|  | Smith-Lemli-Opitz Syndrome |
|  | Smith-Magenis Syndrome |
|  | Sotos Syndrome |
|  | Sturge-Weber Syndrome |
|  | Subcortical Band Heterotopia |
|  | Translocation of Chromosome 2 |
|  | Translocation Trisomy 5/18 |
|  | Trichothiodystrophy |
|  | Triploidy |
|  | Trisomy 10 |
|  | Trisomy 13 (Patau syndrome) |
|  | Trisomy 18 (Edward Syndrome) |
|  | Trisomy 20p |
|  | Trisomy 21 (Down Syndrome) |
|  | Trisomy 47 |
|  | Trisomy 4p |
|  | Trisomy 9 |
|  | Tuberous Sclerosis |
|  | Turner Syndrome |
|  | Urea Cycle Defect |
|  | Valproate Embryopathy |
|  | West Syndrome |
|  | Williams Syndrome |
|  | Wolf-Hirschhorn Syndrome |
|  | X-Linked Adrenoleukodystrophy |
|  | Young-Simpson Syndrome |
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| Category 4 | PARAPLEGIA and QUADRIPLEGIA |
|  | Paraparesis |
|  | Spinal Cord Compression |
|  | Spinal Cord Infarction |
|  | Spinal Cord Damage |
|  | Tetraplegia |
|  | Transverse Myelitis |
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| Category 5 | ACQUIRED NEUROLOGICAL CONDITIONS |
|  | Acquired Brain Injury |
|  | Acute disseminated encephalomyelitis |
|  | Adhesive Arachnoiditis |
|  | Alcoholic Encephalopathy |
|  | Alzheimer’s Disease |
|  | Amyloidosis |
|  | Arachnoiditis |
|  | Ascending Polyneuropathy |
|  | Astrocytoma |
|  | Autonomic Neuropathy |
|  | Basal Ganglia Infarction |
|  | Benign Meningioma |
|  | Brown-Sequard Syndrome |
|  | Cauda Equina compression syndrome |
|  | Cerebral Abscess |
|  | Cerebral Aneurysm |
|  | Cerebral Anoxia |
|  | Cerebral Toxoplasmosis |
|  | Cerebral Tumour |
|  | Cerebrovascular Disease |
|  | Chronic Hypoxia |
|  | Chronic Inflammatory Demyelinating Polyneuropathy (CIDP) |
|  | Cortical-Basal Ganglionic Degeneration |
|  | Dementia (any cause) |
|  | Developmental/Motor Dyspraxia |
|  | Diabetic Autonomic Neuropathy |
|  | Diabetic Neuropathic Bladder |
|  | Dorsal Pontine Band Syndrome |
|  | Encephalitis |
|  | Ependymoma |
|  | Epilepsy |
|  | Focal Cerebral Degeneration |
|  | Glioblastoma Multiforme |
|  | Glioblastoma of Spine |
|  | Hepatic Encephalopathy |
|  | Hydrocephalus (communicating or non-communicating) |
|  | Hypoxic Brain Injury |
|  | Inoperable Neurogenic Incontinence |
|  | Intracerebral Haemorrhage (Subarachnoid Haemorrhage, Subdural Haematoma) |
|  | Korsakoff’s Syndrome |
|  | Lambert-Eaton Myasthenic syndrome |
|  | Lewi Body Disease |
|  | Macroencephaly |
|  | Malignant Meningioma |
|  | Meningoencephalitis |
|  | Metastatic Carcinoma with Neurological Syndrome |
|  | Multiple Systems Atrophy |
|  | Myopathy |
|  | Nemaline Myopathy |
|  | Oligodendroglioma |
|  | Pachymeningitis |
|  | Periventricular Leukomalacia |
|  | Picks Disease |
|  | Pilocytic Astrocytoma |
|  | Poliomyelitis |
|  | Polymyoneuropathy |
|  | Posterior Leuco Encephalopathy |
|  | Primary Dystonia (case by case) |
|  | Primary or secondary CNS B-cell neoplasm |
|  | Progressive supranuclear palsy |
|  | Progressive Systemic Sclerosis |
|  | Sacral Neuroplexy |
|  | Sacral Plexopathy |
|  | Schizophrenia (Catatonic) |
|  | Schwannoma |
|  | Spinal Canal Disease |
|  | Spinal Chordoma |
|  | Spinal Ependymoma |
|  | Spinal Tumour |
|  | Stroke/Cerebrovascular Accident (CVA) |
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| Category 6 | DEGENERATIVE NEUROLOGICAL DISEASES |
|  | Alexander Disease |
|  | Amyotrophic Lateral Sclerosis |
|  | Ataxia Telangiectasia |
|  | Canavan disease |
|  | Cauda Equina Syndrome |
|  | Cervical Myelopathy |
|  | Creutzfeldt-Jakob Disease (CJD) |
|  | Cytochrome C Oxidase Deficiency |
|  | Dejerine-Sottas Disease |
|  | Demyelinating Neuropathy |
|  | Demyelination of White Matter |
|  | Fahr’s Disease |
|  | Friedreich’s Ataxia |
|  | Guillain Barre Syndrome |
|  | Huntington Chorea |
|  | Huntington Disease |
|  | Hypoxic Ischaemic Encephalopathy |
|  | Idiopathic Axonal Neuropathy |
|  | Krabbe disease |
|  | Kugelberg-Welander Syndrome |
|  | Machado Joseph Disease |
|  | Metachromatic Leukodystrophy |
|  | Mitochondrial Myopathy with Encephalopathy |
|  | Morquio Syndrome |
|  | Motor Neurone Disease |
|  | Multiple Sclerosis |
|  | Muscular Dystrophy |
|  | Myotonic dystrophy |
|  | Myoneural Disorders |
|  | Neuroaxonal Dystrophy |
|  | Neurofibromatosis NF |
|  | Neurogenic Bowel |
|  | Neuromyelitis optica |
|  | Niemann-Pick Disease Type C |
|  | Pallister-Hall Syndrome |
|  | Parkinson Disease |
|  | Parkinsonism |
|  | PEHO Syndrome (Progressive encephalopathy with oedema, hypsarrhythmia and optic atrophy) |
|  | Pelizaeus Merzbacher Disease |
|  | Primary Lateral Sclerosis |
|  | Progressive Supranuclear Palsy/Steele Richardson Syndrome |
|  | Sanfilippo Syndrome |
|  | Sarcoidosis of the Brain |
|  | Shy-Drager Syndrome |
|  | Spinal Cord Syndrome |
|  | Spinal Muscular Atrophy Type 1 |
|  | Spinal Muscular Atrophy Type 2 |
|  | Spinocerebellar Degeneration |
|  | Stiff-Mans Syndrome |
|  | Striato-Nigral Degeneration |
|  | Subacute sclerosing pan-encephalitis |
|  | Thiamine deficiency |
|  | Vascular Myelopathy |
|  | Vertebral Canal Stenosis |
|  | Vertebral Degeneration |
|  | Wallerian Degeneration of White Matter |
|  | Wilson’s Disease |
|  |  |
| Category 7 | BLADDER OR BOWEL INNERVATION DISORDERS |
|  | Atonic Bladder/Hypotonic Bladder |
|  | Bladder Innervation Urgency |
|  | Cystocele (not suitable for surgery) |
|  | Dysfunctional Voiding |
|  | Dystonic Bladder |
|  | Ectopia Vesica |
|  | Linear Sebaceous Nevus Genetic |
|  | Myasthenia Gravis |
|  | Neurogenic Bladder |
|  | Neuronal Intestinal Dysplasia |
|  | Neuropathic Bladder |
|  | Post Bladder Surgery |
|  | Prostatectomy with nerve removal or damage |
|  | Pudendal Nerve Palsy |
|  | Radical Prostatectomy |
|  | Schmidli Autonomic Neuropathy |
|  | Slow Transit Constipation |
|  | Smooth Muscle MyopathySphincter Deficiency (anal or bladder) |
|  | Spinal Stenosis |

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