**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 |
|   |   |
| Category 2 | CEREBRAL PALSY |
|   | Dystonic Cerebral Palsy |
|   | Hereditary Spastic Paralysis |
|   | Spastic Quadriplegia |
|   | Mixed cerebral palsy |
|   |   |
| 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 |
|   |   |
| Category 4 | PARAPLEGIA and QUADRIPLEGIA |
|   | Paraparesis |
|   | Spinal Cord Compression |
|   | Spinal Cord Infarction |
|   | Spinal Cord Damage |
|   | Tetraplegia |
|   | Transverse Myelitis |
|   |   |
| 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) |
|   |   |
| 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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