Part 1—Eligible neurological conditions

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