

## CAPS Eligible Neurological Conditions - NO PCC Required

Category 1	SPINA BIFIDA and SYRINGOMYELIA
	Arnold-Chiari Syndrome
	Arthrogryposis
	Caudal Regression Syndrome
	Developmental Cord Disorder
	Holoprosencephaly
	Malformation of Spinal Cord
	Sacral Agenesis
	Spinal Agenesis
	Spinal Cord Congenital Abnormality
	Spinal Dysraphism
	Spinal Hemangioma
	Vater Syndrome/Vacterl Syndrome
Category 2	CEREBRAL PALSY
	Dystonic Cerebral Palsy
	Hereditary Spastic Paralysis
	Spastic Quadriplegia
Category 3	INTELLECTUAL DISABILITY
	2-Hydroxyglutaric Aciduria
	Agenesis of Corpus Callosum
	Alpers Disease
	Angelman Syndrome
	Apert Syndrome
	ATRX Syndrome
	Bardot Biedl Syndrome
	Batten Disease
	Beare-Stevenson Syndrome
	Cats Cry Syndrome/Cri Du Chat Syndrome
	CDKL5 Gene Mutation
	Cerebral Migration Disorders
	Charge Syndrome
	Chime Syndrome
	Chromosomal Abnormalities/Disorders
	Chromosome 1 Deletion
	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 Abnormality
	Chromosome 9p Deletion Syndrome
	Chromosome 9q Deletion Syndrome
	Chromosome Xp Duplication
	Cockayne Syndrome
	Coffin-Lowry Syndrome
	Cognitive Impairment
	Cohen Syndrome

Congenital Hydrocephalus
Congenital Neurological Infections
Cornelia de Lange Syndrome
Costello Syndrome
Cowden Disease
Developmental Delay
Developmental Delay associated with Autism, Autism Spectrum Disorder and Aspergers Syndrome
Down Syndrome/Trisomy 21
Dravet Syndrome
Edwards Syndrome/Trisomy 18
Fragile X Syndrome
Fumarase Deficiency
GLUT1-DS Condition
Glutaric Aciduria Type 1
Goldenhar's Syndrome
Hunter Syndrome
Hurler-Scheie Syndrome
Hypomyelination
Incomplete Corpus Callosum/Aicardi Syndrome
Inversion Duplication of Chromosome 8
Jacobsen Syndrome/11q Chromosome Deletion
Joubert Syndrome
Kabuki Syndrome
Langer-Gideon Syndrome
Lawrence Moon Biedel Syndrome
Lennox-Gastaut Syndrome
Lesch-Nyhan Syndrome
Lissencephaly
Lowe Syndrome
Mannosidosis
Maple Syrup Urine Disease
Meningitis
Menkes Syndrome
Microcephaly
Mitochondrial Deficiency
Molybdenum Cofactor Deficiency
Mowat-Wilson Syndrome
Mucopolipidosis IV
Myotonic Dystrophy (Type 1)
Neonatal Hypoxia
Neonatal Onset Multisystem Inflammatory Disease
Normal Pressure Hydrocephalus
OHDO Syndrome
Opitz Trignocephaly Syndrome
Ohtahara Syndrome
Ouvrier Syndrome
Pallister-Killian Mosaic Syndrome
Periventricular Leukomalacia
Peroxisome Biogenesis Disorder
Phelan McDermid Syndrome/22q 13 Deletion Syndrome
Phenylketonuria
Polymicrogyria
Pontocerebellar Hypoplasia

	Prader-Willi Syndrome
	Pyruvate Dehydrogenase Deficiency/Leigh's Disease
	Rare Congenital Neurological Syndromes and Conditions
	Rasmussen's Disease
	Rett Syndrome
	Rubinstein-Taybi Syndrome
	Schizencephaly
	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
	Trisomy 10
	Trisomy 13
	Trisomy 20p
	Trisomy 47
	Trisomy 4p
	Trisomy 9
	Tuberous Sclerosis
	Turner Syndrome
	Urea Cycle Defect
	Valproate Embryopathy
	Velocardiofacial Syndrome
	West Syndrome
	Williams Syndrome
	Wolf-Hirschhorn Syndrome
	X-Linked Adrenoleukodystrophy
	Young-Simpson Syndrome
<b>Category 4</b>	<b>PARAPLEGIA and QUADRIPLÉGIA</b>
	Paraparesis
	Spinal Cord Compression
	Spinal Cord Infarction
	Spinal Damage
	Syringomyelia
	Tetraplegia
	Transverse Myelitis
<b>Category 5</b>	<b>ACQUIRED NEUROLOGICAL CONDITIONS</b>
	Acquired Brain Injury
	Adhesive Arachnoiditis
	Alcoholic Encephalopathy
	Alzheimer's Disease
	Arachnoiditis
	Ascending Polyneuropathy
	Astrocytoma
	Autonomic Neuropathy Disease
	Basal Ganglia Infarction
	Benign Meningioma
	Brown-Sequard Syndrome

Cauda Equina Lesion
Cerebral Abscess (Cryptococcus)
Cerebral Aneurysm
Cerebral Anoxia
Cerebral Toxoplasmosis
Cerebral Tumour
Cerebrovascular Disease
Chronic Hypoxia
Chronic Inflammatory Demyelination Polyneuropathy (CIDP)
Cortical-Basal Ganglionic Degeneration
Dementia
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
Hypoxic Brain Damage
Inoperable Neurogenic Incontinence
Intracerebral Haemorrhage
Korsakoff's Syndrome
Leuco Posterior Encephalopathy
Lewi Body Disease
Macroencephaly
Malignant Meningioma
Metastatic Carcinoma with Neurological Syndrome
Multiple Systems Atrophy
Myopathy
Nemaline Myopathy
Oligodendroglioma
Pachymeningitis
Picks Disease
Pilocytic Astrocytoma
Poliomyelitis
Polymyoneuropathy
Primary Dystonia (case by case)
Progressive Systemic Sclerosis
Sacral Neuroplexy
Sacral Plexopathy
Schizophrenia (Catatonic)
Spinal Canal Disease
Spinal Chordoma
Spinal Ependymoma
Spinal Sacral Chordoma
Spinal Tumour
Stroke/Cerebrovascular Accident (CVA)
Subarachnoid Haemorrhage
Subdural Haematoma

	Vascular Dementia
<b>Category 6</b>	<b>DEGENERATIVE NEUROLOGICAL DISEASES</b>
	Alexander Disease
	Amyotrophic Lateral Sclerosis
	Ataxia Telangiectasia
	Cauda Equina Syndrome
	Cervical Canal Stenosis
	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/Disease
	Hypoxic Ischaemic Encephalopathy
	Idiopathic Axonal Neuropathy
	Kugelberg-Welander Syndrome
	Machado Joseph Disease
	Metachromatic Leukodystrophy
	Mitochondrial Myopathy with Encephalopathy
	Morquio Syndrome
	Motor Neurone Disease
	Multiple Sclerosis
	Muscular Dystrophy
	Myoneural Disorders
	Neuroaxonal Dystrophy
	Neurofibromatosis NF
	Neurogenic Bowel
	Niemann-Pick Disease Type C
	Pallister-Hall Syndrome
	Parkinson Disease
	Parkinsonism
	PEHO Syndrome (Progressive encephalopathy with oedema, hysarrhythmia 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 Degeneration
	Spinal Muscular Atrophy Type 1
	Spinal Muscular Atrophy Type 2
	Spinocerebellar Degeneration
	Stiff-Mans Syndrome
	Striato-Nigral Degeneration
	Vascular Myelopathy
	Wallerian Degeneration of White Matter
	Wilson's Disease

<b>Category 7</b>	<b>BLADDER (BOWEL) INNERVATION DISORDERS</b>
	Atonic Bladder/Hypotonic Bladder
	Bladder Exstrophy
	Bladder Innervation Urgency
	Cloacal Exstrophy
	Cystocele (not suitable for surgery)
	Dysfunctional Voiding
	Dystonic Bladder
	Ectopia Vesica
	Hirschsprung's Disease
	Linear Sebaceous Nevus Genetic
	Myasthenia Gravis
	Neurogenic Bladder
	Neuronal Intestinal Dysplasia
	Neuropathic Bladder
	Post Bladder Surgery
	Prostatectomy with nerve removal
	Prune Belly Syndrome
	Pudendal Nerve Palsy
	Radical Prostatectomy
	Schmidli Autonomic Neuropathy
	Slow Transit Constipation
	Smooth Muscle Myopathy
	Sphincter Deficiency
	Spinal Stenosis