CAPS Eligible Other Conditions - PCC Required (Bladder Bowel)

This Eligibility List is not exhaustive. However, it does state many of the eligible conditions for the purpose of the CAPS. For conditions not listed, specialist medical advice will be sought on receipt of an Application Form. Please send any enquiries to: continence@health.gov.au

Code 8	**OTHER**
	Anal Carcinoma
	Anal Fistula
	Anorectal Malformation
	Anterior Prolapse
	Bilateral Nephrostomy Tubes
	Bladder Cancer
	Bladder Instability
	Bladder Muscle Dysfunction
	Bladder Neck Dysfunction
	Bladder Neck Fibrosis
	Bladder Prolapse
	Bowel Cancer
	Bowel Prolapse
	Cervical Cancer
	Chronic Urinary Retention
	Congenital Epispadias
	Detrusor Instability
	Detrusor Overactivity
	Enterocutaneous Fistula
	Hypertonic Bladder
	Imperforate Anus
	Irradiated Rectum/Radiation Proctitis
	Posterior Urethral Valve Syndrome
	Prostate Cancer
	Prostate Disease
	Rectal Prolapse
	Rectal Ulcer Syndrome
	Severe Ulcerative Proctitis
	Spastic Bladder
	TURP
	Urethral Stenosis
	Urinary Fistula
	Uterine Cancer
	Uterine Prolapse
	Vaginal Prolapse
	Vesico-Vaginal Fistula
	Vulva Cancer

	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
	Mowat-Wilson Syndrome
	Mucolipidosis IV
	Myotonic Dystrophy (Type 1)
	Neonatal Hypoxia
	Neonatal Onset Multisystem Inflammatory Disease
	Normal Pressure Hydrocephalus

	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
Category 4	**PARAPLEGIA and QUADRIPLEGIA**
	Paraparesis
	Spinal Cord Compression
	Spinal Cord Infarction
	Spinal Damage
	Syringomyelia
	Tetraplegia
	Transverse Myelitis
Category 5	**ACQUIRED NEUROLOGICAL CONDITIONS**
	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
Category 6	**DEGENERATIVE NEUROLOGICAL DISEASES**
	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, 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 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
Category 7	**BLADDER (BOWEL) INNERVATION DISORDERS**
	Atonic Bladder/Hypotonic Bladder
	Bladder Exstrophy
	Bladder Innervation Urgency
	Cloacal Exstrophy
	Cystocele (not suitable for surgery)
	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
	Smooth Muscle Myopathy
	Sphincter Deficiency
	Spinal Stenosis

	Stiff-Mans Syndrome
	Striato-Nigral Degeneration
	Vascular Myelopathy
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	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
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	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, 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 Degeneration
	Spinal Muscular Atrophy Type 1
	Spinal Muscular Atrophy Type 2
	Spinocerebellar Degeneration

CAPS Eligible Neurological Conditions - NO PCC Required

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	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
	Angelman Syndrome
	Apert Syndrome
	ATRX Syndrome
	Bardot Biedl Syndrome
	Batten Disease
	Beare-Stevenson Syndrome
	Cats Cry Syndrome/Cri Du Chat Syndrome
	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 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
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	Lowe Syndrome
	Mannosidosis
	Maple Syrup Urine Disease
	Meningitis
	Menkes Syndrome
	Microcephaly
	Mitochondrial Deficiency
	Mowat-Wilson Syndrome
	Mucolipidosis IV
	Myotonic Dystrophy (Type 1)
	Neonatal Hypoxia
	Neonatal Onset Multisystem Inflammatory Disease
	Normal Pressure Hydrocephalus
	OHDO Syndrome
	Opitz Trigonocephaly 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
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	Primary Dystonia (case by case)
	Progressive Systemic Sclerosis
	Sacral Neuroplexy
	Sacral Plexopathy
	Schizophrenia (Catatonic)
	Spinal Canal Disease
	Spinal Chordoma
	Spinal Ependymoma
	Spinal Sacral Chordoma