

CHILDREN'S SPECIAL HEALTH SERVICES

MEDICAL CONDITION LIST

Medical eligibility is based on a list of conditions which has been established with the advice of a Medical Advisory Council and is subject to change.

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| <p>ACQUIRED BRAIN INJURY
ADENOID HYPERTROPHY causing SLEEP APNEA
ALPHA 1-ANTITRYPSIN DEFICIENCY
AMPUTATION
AMYOTONIA CONGENITA requiring rehabilitative measures
ANAL STENOSIS & IMPERFORATE ANUS
ANEMIAS (excluding minor anemias), including sickle cell
APLASIA CUTIS CONGENITA, severe, requiring surgery & ECTODERMAL DYSPLASIA
ARNOLD-CHIARI DEFORMITY
ARTHROGRYPOSIS
ASTHMA, persistent, requiring controller medications
ATAXIAS, FAMILIAL DEGENERATIVE DISEASE requiring rehabilitative measures</p> <p>BILE DUCT ATRESIA
BIRTH INJURY (ERB's PALSY, etc.) requiring bracing or surgery
BONE CYST requiring surgery
BONE TUMORS, benign, requiring surgery, including OSTEOCHONDROMAS
BONY DEFORMITIES requiring bracing, casting or surgery & POST-TRAUMATIC DEFORMITY (orthopedic or severe soft tissue deformity due to injury; excluding acute fracture without an underlying condition)
BOWED LEGS, severe
BRAIN TUMORS requiring surgery and/or radiation
BRANCHIOGENIC CLEFT CYST requiring surgery
BREAST HYPOPLASIA causing considerable psychological problems requiring surgery
BURNS, severe, acute, including residuals</p> <p>CANCER, including CANCER OF EYE
CATARACTS
CELIAC DISEASE
CEREBRAL PALSY, congenital or acquired, requiring rehabilitative measures
CHOANAL ATRESIA
CLEFT LIP AND/OR PALATE, including SHORT PALATE
CONGENITAL ADRENAL HYPERPLASIA (CAH)
CORNEAL TRANSPLANTS
CRANIOSTENOSIS (premature synostosis)
CYSTIC FIBROSIS
CYSTIC HYGROMA
CYSTINOSIS</p> <p>DENTAL DISORDERS, congenital
DIABETES INSIPIDUS
DIABETES MELLITUS, TYPE I and TYPE II
DIAPHRAGMATIC HERNIA
DISLOCATION OF HIPS OR OTHER JOINTS</p> <p>EAR DEFORMITY
EHLERS-DANLOS DISEASE
ENCEPHALITIS, POLIOMYELITIS OR MENINGITIS, residuals of</p> | <p>ENUCLEATION (removal of eyeball)
EOSINOPHILIC GASTROENTERITIS
EPIDERMOLYSIS BULLOSA
ESOPHAGEAL VARICES
EYE WOUNDS, penetrating
EYELID DEFORMITY requiring surgery, congenital</p> <p>FACE DEFORMITY
FEMORAL CAPITAL EPIPHYSIS, slipped</p> <p>GASTROINTESTINAL TRACT ANOMALIES, congenital (including gastroschisis)
GENITO-URINARY TRACT ANOMALIES, congenital, severe and requiring surgery
GENU RECURVATUM, severe
GLAUCOMA, congenital
GROWTH HORMONE DEFICIENCY
GUILLAIN-BARRE DISEASE, severe, acute, requiring tracheotomy and/or ventilation, including residuals</p> <p>HALLERVORDEN-SPATZ DISEASE including infusion pump
HEARING LOSS
HEART CONDITIONS, congenital or acquired
HEMANGIOMA, medically significant
HEMOGLOBINOPATHIES, limited to:
Sickle cell anemia
Thalassemia
HEMOPHILIA including deformities
HISTIOCYTOSIS X (eosinophilic granuloma)
HYDROCEPHALUS requiring surgery
HYPERCHOLESTEROLEMIA, congenital, including familial combined hyperlipidemia
HYPERTHYROIDISM
HYPOPARATHYROIDISM, congenital or if suspected to last longer than two years
HYPOPHOSPHATEMIC RICKETS
HYPOTHALAMIC ADRENAL INSUFFICIENCY
HYPOTHYROIDISM and SUBMUCOUS CLEFT
ICHTHYOSIFORM ERYTHRODERMA, congenital, severe
IMMUNODEFICIENCY STATES including severe combined immunodeficiency (SCID)
INFLAMMATORY BOWEL DISEASE including Crohn's Disease and ulcerative colitis
INTERSEX DISORDERS, congenital</p> <p>JOINT DEFORMITY, CLUBFEET AND CLUBHANDS, severe, requiring bracing, casting, surgery or physical therapy
KNOCK-KNEES, severe
KYPHOSIS, adolescent, requiring bracing or surgery</p> <p>LARYNGEAL PAPILLOMA
LEUKEMIA</p> |
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LEUKODYSTROPHY

MALOCCLUSION, handicapping

MASTOIDITIS, chronic

MEGACOLON requiring surgery

METABOLIC DISORDERS/INBORN ERRORS OF METABOLISM

Amino Acid Disorders, limited to:

- Arginase deficiency/Hyperargininaemia
- Argininemia
- Argininosuccinic acidemia (ASA lyase deficiency)
- Carbamoyl phosphate synthetase deficiency
- Citrullinemia (ASA synthetase deficiency)
- Glutaric acidemia/aciduria
- Glutathione synthetase deficiency (5-oxoprolinuria)
- Homocystinuria (cystathione synthase deficiency)
- Hypermethioninemia
- Hyperornithinemia, hyperammonemia, Homocitrullinemia (HHH syndrome)
- Hyperornithinemia or ornithine oxo-acid aminotransferase deficiency
- Maple syrup urine disease
- N-Acetylglutamate synthetase deficiency
- Nonketotic hyperglycinemia
- Ornithine aminotransferase deficiency
- Ornithine transcarbamylase deficiency (OTC)
- Phenylketonuria;
- Tyrosinemia (I, II, III);

Biotinidase Deficiency

Fatty Acid Oxidation Disorders, limited to:

- 2,4 dienoyl-CoA reductase deficiency
- Long chain acyl-CoA dehydrogenase deficiency (LCADD)
- Long chain 3-OH acyl-CoA dehydrogenase deficiency (LCHAD)
- Carnitine/acylcarnitine translocase deficiency (CACT)
- Carnitine palmitoyltransferase deficiency-type I (CPTI)
- Carnitine palmitoyltransferase deficiency-type II (CPTII)
- Carnitine transport defect (CTD)
- Glutaric acidemia/aciduria
- Medium chain acyl-CoA dehydrogenase deficiency (MCAD)
- Multiple acyl-CoA dehydrogenase deficiency (MADD) or glutaric acidemia-type II (GAII)
- Short chain acyl-CoA dehydrogenase deficiency (SCAD) (ethylmalonic acidemia)
- Trifunctional protein deficiency (TFP Deficiency)
- Very long chain acyl-CoA dehydrogenase deficiency (VLCAD)

Galactosemia

GLUT 1 Deficiency (glucose 1 transporter deficiency)

Glycogen Storage Disease

Hereditary Fructose Intolerance

Organic Acid Disorders, limited to:

- 2-methylbutyryl-CoA dehydrogenase deficiency
- 3-methylcrotonyl-CoA carboxylase deficiency
- 3-methylglutaconic-CoA hydratase deficiency
- 3-hydroxy-3-methylglutaryl-CoA lyase deficiency
- Glutaric acidemia/aciduria

Isobutyryl-CoA dehydrogenase deficiency

Isovaleric acidemia (IVA)

Methylmalonic acidemia (MMA)

Propionic Acidemia

Mitochondrial acetoacetyl-CoA thiolase deficiency (BKT, 3-ketothiolase deficiency)

Multiple CoA carboxylase deficiency

Refsum's Disease (Phytanic acid restriction)

MICROCEPHALY, diagnosis only

MUCOPOLYSACCHARIDOSIS (MPS) (including variants)

MUSCULAR DYSTROPHY

NEPHROSIS & CHRONIC NEPHRITIS

NERVE INJURIES, chronic

NEUROFIBROMATOSIS

NEVI with malignant potential

OCULAR ALBINISM, congenital

OSTEOCHONDRITIS of various bones

OSTEOGENESIS IMPERFECTA

OSTEOMYELITIS, residuals of

PARAPLEGIA, traumatic, and its direct complications

PECTUS CARINATUM/PECTUS EXCAVATUM requiring surgery

PERTHES DISEASE

POLYCYSTIC KIDNEY DISEASE

PRECOCIOUS PUBERTY

PSEUDOHYPOPARATHYROIDISM

PTOSIS (drooping eyelids)

PULMONARY LOBAR EMPHYSEMA

RETINAL DETACHMENT in Marfan's syndrome

RETROLENTAL FIBROPLASIA (retinopathy of prematurity)

RHEUMATOID ARTHRITIS

SCLERODERMA

SCOLIOSIS requiring bracing or surgery

SEIZURE DISORDERS, excluding febrile seizures

SHORT BOWEL SYNDROME

SPINA BIFIDA, MENINGOCELE, MYELOCELE

STRABISMUS through age 10

SUBLUXATED EYE LENS in Marfan's syndrome

SUPERNUMERARY PARTS, severe

SYNDACTYLY

SYNDROMES, limited, requiring ongoing medical treatment

THROMBOCYTOPENIA, congenital

THYROGLOSSAL DUCT CYST

T-LYMPHOCYTE IMMUNE DEFICIENCY STATE

TORTICOLLIS (wryneck, not spasmodic, requiring casting or surgery)

TRACHEAL STENOSIS

TRACHEOESOPHAGEAL FISTULA

TUBERCULOSIS OF BONES AND JOINTS

TUBEROUS SCLEROSIS

UNDESCENDED TESTES

WEGENER'S GRANULOMATOSIS