NORTH Dakota | Health & Human Services
Be Legendary.

SPECIAL HEALTH SERVICES MEDICAL CONDITIONS **Revised: 6-30-2023**

****

ACQUIRED BRAIN INJURY
ACUTE FLACCID MYELITIS
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
ATTENTION DEFICIT/HYPERACTIVITY DISORDER (ADD/ADHD)
AUTO-IMMUNE DISORDERS, chronic, severe, and complex in nature
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)
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
CANCER, including CANCER OF EYE
CATARACTS
CELIAC DISEASE
CEREBRAL PALSY, congenital or acquired, requiring rehabilitative measures
CHOANAL ATRESIA
CHRONIC LUNG/LOWER AIRWAY CONDITIONS, including chronic lung disease, chemical pneumonitis, and subglottic stenosis
CLEFT LIP AND/OR PALATE, including SHORT PALATE and SUBMUCOUS CLEFT
COLLAGEN VASCULAR DISORDERS, including but not limited to lupus, dermatomyositis, scleroderma, Sjogren's syndrome, and rheumatoid arthritis
CONGENITAL ADRENAL HYPERPLASIA (CAH)
CORNEAL TRANSPLANTS
CRANIOSTENOSIS (premature synostosis)
CYSTIC FIBROSIS
CYSTIC HYGROMA
CYSTINOSIS
DENTAL DISORDERS, congenital
DERMATOMYOSITIS
DIABETES INSIPIDUS
DIABETES MELLITUS, TYPE I and TYPE II
DIAPHRAGMATIC HERNIA
DISLOCATION OF HIPS OR OTHER JOINTS
DOWN SYNDROME
EAR DEFORMITY
EHLERS-DANLOS DISEASE
ENCEPHALITIS, POLIOMYELITIS OR MENINGITIS, residuals of
ENUCLEATION (removal of eyeball)
EOSINOPHILIC GASTROENTERITIS
EPIDERMOLYSIS BULLOSA
ESOPHAGEAL VARICES
EYE WOUNDS, penetrating
EYELID DEFORMITY requiring surgery, congenital
FACE DEFORMITY
FEMORAL CAPITAL EPIPHYSIS, slipped
GASTROINTESTINAL TRACT ANOMALIES, congenital (Including gastroschisis)
GENITO-URINARY TRACT ANOMALIES, congenital, severe, and requiring surgery
GENU RECURVATUM, severe
GENU VALGUM (Knock-knees), severe
GENU VARUM (Bowed legs), severe
GLAUCOMA, congenital
GROWTH HORMONE DEFICIENCY
GUILLAIN-BARRE DISEASE, severe, acute, requiring tracheotomy and/or ventilation, including residuals
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
HYPOPITUITARISM
HYPOTHALAMIC ADRENAL INSUFFICIENCY
HYPOTHYROIDISM
ICHTHYOSIFORM ERYTHRODERMA, congenital, severe
IMMUNODEFICIENCY STATES including severe combined immunodeficiency (SCID)
INFLAMMATORY BOWEL DISEASE including Crohn's Disease and ulcerative colitis
INTERSEX DISORDERS, congenital
JOINT DEFORMITY, CLUBFEET AND CLUBHANDS, severe, requiring bracing, casting, surgery, or physical therapy
JUVENILE IDIOPATHIC ARTHRITIS, (Juvenile Rheumatoid Arthritis)
KYPHOSIS, adolescent, requiring bracing or surgery
LARYNGEAL PAPILLOMA

LEUKEMIA
LEUKODYSTROPHY, including adrenoleukodystrophy
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 (MSUD)
- N-Acetylglutamate synthetase deficiency
- Nonketotic hyperglycinemia
- Ornithine aminotransferase deficiency
- Ornithine transcarbamylase deficiency (OTC)
- Phenylketonuria (PKU), includes phenylalanine hydroxylase deficiency (PAH) and hyperphenylalaninemia
- 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 academia)
- 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
Lysosomal Storage Disease
Organic Acid Disorders, limited to:
- 2-methylbuyryl-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
MUCOPOLYSACCHARIDOSIS (MPS) (including variants)
NARCOLEPSY (with or without Cataplexy)
NEPHROSIS & CHRONIC NEPHRITIS
NERVE INJURIES, chronic
NEUROFIBROMATOSIS
NEUROMUSCULAR DISORDERS limited to those covered by MDA including muscular dystrophy
NEVI with malignant potential
OCULAR ALBINISM, congenital
OSTEOCHONDRITIS of various bones
OSTEOGENESIS IMPERFECTA
OSTEOMYELITIS, residuals of
PANTOTHENATE KINASE-ASSOCIATED NEURODEGENERATION (PKAN)
(Hallervorden-Spatz Disease, including infusion pump)
PARAPLEGIA, traumatic, and its direct complications
PECTUS CARINATUM/PECTUS EXCAVATUM requiring surgery
PERTHES DISEASE
POLYCYSTIC KIDNEY DISEASE
PRECOCIOUS PUBERTY
PSEUDOHYPOPARATHYROIDISM
PSORIASIS
PTOSIS (drooping eyelids)
PULMONARY LOBAR EMPHYSEMA
RETINAL DETACHMENT in Marfan's syndrome
RETROLENTAL FIBROPLASIA (retinopathy of prematurity)
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; includes septo-optic dysplasia
THROMBOCYTOPENIA, congenital
THROMBOEMBOLISM
THYROGLOSSAL DUCT CYST
T-LYMPHOCYTE IMMUNE DEFICIENCY STATE
TORTICOLLIS (not spasmodic, requiring casting or surgery)
TRACHEAL STENOSIS
TRACHEOESOPHAGEAL FISTULA
TRANSVERSE MYELITIS
TUBERCULOSIS OF BONES AND JOINTS
TUBEROUS SCLEROSIS
UNDESCENDED TESTES
VASCULAR ABNORMALITIES
WEGENER'S GRANULOMATOSIS