

Established Conditions (Not an Exhaustive List)

Genetic and Metabolic Disorders	
Albinism	
Albright's Hereditary Osteodystrophy	
Angelman Syndrome (Happy Puppet Syndrome)	
Adrenoleukodystrophy	
Antley-Bixler Syndrome (Multisynostotic Osteodysgenesis, Craniosynostosis, Choanal Atresia, Radial Humeral Synostosis, Trapezoidocephaly-Multiple Synostosis Syndrome, ABS, Multisynostotic Osteodysgenesis with Long Bone Fractures)	
Apert Syndrome (Acrocephalosyndactyly)	
Arthrogryposis Multiplex Congenita	
Ataxia-Telangiectasia Syndrome (Louis-Bar Syndrome)	
Canavan Disease	
Cardio-Facio-Cutaneo Syndrome	
Cerebral Lipidosis	
Cerebro-Oculo-Facio-Skeletal (COFS) Syndrome	
CHARGE Syndrome/Association	
Chromosome Syndromes 10p+, 13q+, 3q+, 4Q+	
Chromosome Syndromes, such as 11p- (Jacobsen syndrome), 12p-, 13q-, 18q-, 21q-, 22q-, , 4q-, (Wolf-Hirschhorn syndrome), 5p- (Cri-du-chat syndrome)	
Coffin-Lowry Syndrome	
Coffin-Siris Syndrome	
Cornelia de Lange Syndrome (Brachmann de Lange)	
Cystic Fibrosis	
Dandy Walker Syndrome	
Down Syndrome (Trisomy 21)	
Duchenne Muscular Dystrophy	
Dyggve-Melchior-Clausen Syndrome (DMC Disease, DMC Syndrome, Smith-McCort Dysplasia)	
Fragile X Syndrome	
Fraser Syndrome (Cryptophthalmos Syndrome, Meyer-Schwickerath's syndrome, Fraser-Francois syndrome, Ullrich-Feichtiger syndrome)	
Galactosemia	
Gaucher Syndrome (Glucosylceramide storage disease; GSDI)	
Glutaric Aciduria (Type I, Type II)	
Glycogen Storage Disease	
Jeune Syndrome	
Joubert Syndrome	
Krabbe's disease	
Lesch-Nyhan Syndrome	
Lissencephaly Syndrome (Miller-Dieker Syndrome, Agyria)	
Maple Syrup Urine	
Mucopolysaccharidosis I (MPS I)	
Mucopolysaccharidosis II, III	
Organic Acidemias	
Pelizaeus-Merzbacher disease	
Peroxisomal Disorders	
Phenylketonuria (PKU)	

Established Conditions
(Not an Exhaustive List)

Phelan-McDermid syndrome
Pompe
Prader-Willi Syndrome
Rubenstein-Taybi Syndrome
Schwartz-Jampel Syndrome
Spinal Muscular Atrophy (SMA)
Steinert Myotonic Dystrophy Syndrome (Curschmann-Batten-Steinert syndrome)
Tay-Sachs disease (Sandhoff)
Trisomy 8
Trisomy 9
Tetrasomy 12p
Trisomy 13 (Patau Syndrome)
Trisomy 18 (Edward's Syndrome)
Tuberous Sclerosis Complex
Urea Cycle Defect
Very Long Chain Fatty Acid Storage Diseases
Walker-Warburg Syndrome (XO)
Williams Syndrome
Zellweger Syndrome (Cerebro-Hepato-Renal Syndrome)
Neurological Disorders
Agyria (Miller-Dieker lissencephaly syndrome (MDLS), agyria syndrome, agyria-pachygyria syndrome, classical lissencephaly)
Aicardi Syndrome
Alpers Syndrome/Disease
Aphasia
Arachnoid cyst with Neuro-Developmental Delay
Arhinencephaly (Holoprosencephaly)
Arnold-Chiari Syndrome, type II (Malformation d'Arnold-Chiari)
Ataxia
Cerebral Palsy
Cerebral Aneurysm with Neuro-Developmental Delay
CNS Tumor with Neuro- Developmental Delay
Encephalopathy, Congenital Only
Encephalopathy, Static
Erb's Palsy (Brachial Plexus Injury, Perinatal Origin)
Extracorporeal Membrane Oxygenation (ECMO)
Holoprosencephaly
Hypertonia (persistent only)
Hypoxic Ischemic Encephalopathy (HIE)
Lennox-Gastaut Syndrome
Intracranial Calcifications
Intraventricular Hemorrhage (Grade 3, Grade 4)
Meningocele (cervical)
Microcephaly
Miller-Dieker Syndrome
Mitochondrial Disorder

Established Conditions (Not an Exhaustive List)

Multiple Anomalies of the Brain
Myopathy
Neonatal/Perinatal Asphyxia (5-minute Apgar score of 6 or less, Cord PH < 7, Evidence of Central Nervous System involvement, Organ failure, Resuscitation)
Periventricular Leukomalacia (PVL)
Spina Bifida
Spinocerebellar Disorders
Severe Attachment Disorders
Anxiety Disorders of Infancy and Early Childhood
Depression of Infancy and Early Childhood
Infantile Anorexia
Autism Spectrum Disorders
Asperger's Disorder
Autism Spectrum Disorder
Pervasive Developmental Disorder
Rett's Syndrome
Significant Sensory Impairment
Auditory Neuropathy/Auditory Dyssynchrony
Aural Atresia/Microtia and Atresia (bilateral or unilateral)
Blindness ("legal" blindness or 20/200 best acuity with correction)
Optic Nerve Hypoplasia (De Morsier's Syndrome), Septo Optic Dysplasia
Permanent Conductive/Sensorineural Hearing Loss (in excess of 25 dB HL)
Retinopathy of Prematurity Stage III and/or IV (ROP)
• Stage 3 unspecified (Bilateral, Left eye, Right eye) • Stage 4 unspecified (Bilateral, Left eye, Right eye) • Stage 5 unspecified (Bilateral, Left eye, Right eye)
Other
Fetal Alcohol Syndrome
Hydrocephalus (congenital or acquired)
Lead Poisoning
Low Birth Weight (<1,200 grams at birth)
Zika Probable or Confirmed (Congenital with Symptoms, Congenital with No Symptoms)