

<b>Established Conditions (Not an Exhaustive List)</b>
<b><i>Genetic and Metabolic Disorders</i></b>
<b>Albinism</b>
<b>Albright's Hereditary Osteodystrophy</b>
<b>Angelman Syndrome (Happy Puppet Syndrome)</b>
<b>Adrenoleukodystrophy</b>
<b>Antley-Bixler Syndrome (Multisynostotic Osteodysgenesis, Craniosynostosis, Choanal Atresia, Radial Humeral Synostosis, Trapezoidocephaly-Multiple Synostosis Syndrome, ABS, Multisynostotic Osteodysgenesis with Long Bone Fractures)</b>
<b>Apert Syndrome (Acrocephalosyndactyly)</b>
<b>Arthrogyposis Multiplex Congenita</b>
<b>Ataxia-Telangiectasia Syndrome (Louis-Bar Syndrome)</b>
<b>Canavan Disease</b>
<b>Cardio-Facio-Cutaneo Syndrome</b>
<b>Cerebral Lipidosis</b>
<b>Cerebro-Oculo-Facio-Skeletal (COFS) Syndrome</b>
<b>CHARGE Syndrome/Association</b>
<b>Chromosome Syndromes 10p+, 13q+, 3q+, 4Q+</b>
<b>Chromosome Syndromes 11p- (this one also called Jacobsen syndrome), 12p-, 13q-, 18q-, 21q-, 22q-, , 4q-, (this is also Wolf-Hirschhorn syndrome) 5p- (already below as cri-du-chat syndrome)</b>
<b>Coffin-Lowry Syndrome</b>
<b>Coffin-Siris Syndrome</b>
<b>Cornelia de Lange Syndrome (Brachmann de Lange)</b>
<b>Cri-du-chat Syndrome (Deletion 5p Syndrome)</b>
<b>Cystic Fibrosis</b>
<b>Dandy Walker Syndrome</b>
<b>Down Syndrome (Trisomy 21)</b>
<b>Duchenne Muscular Dystrophy</b>
<b>Dyggve-Melchior-Clausen Syndrome (DMC Disease, DMC Syndrome, Smith-McCort Dysplasia)</b>
<b>Fragile X Syndrome</b>
<b>Fraser Syndrome (Cryptophthalmos Syndrome, Meyer-Schwickerath's syndrome, Fraser-Francois syndrome, Ullrich-Feichtiger syndrome)</b>
<b>Galactosemia</b>
<b>Gaucher Syndrome (Glucosylceramide storage disease; GSDI)</b>
<b>Glutaric Aciduria</b>
<b>Type I</b>
<b>Type II</b>
<b>Glycogen Storage Disease</b>
<b>Jeune Syndrome</b>
<b>Joubert Syndrome</b>
<b>Krabbe's disease</b>
<b>Lesch-Nyhan Syndrome</b>
<b>Lissencephaly Syndrome (Miller-Dieker Syndrome, Agyria)</b>
<b>Maple Syrup Urine</b>
<b>Mucopolysaccharidosis I (MPS I)</b>
<b>Mucopolipidosis II, III</b>
<b>Organic Acidemias</b>

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

<b>Microcephaly</b>
<b>Miller-Dieker Syndrome</b>
<b>Mitochondrial Disorder</b>
<b>Multiple Anomalies of the Brain</b>
<b>Myopathy</b>
<b>Neonatal/Perinatal Asphyxia (5 minute Apgar score of 6 or less, Cord PH &lt; 7, Evidence of Central Nervous System involvement, Organ failure, Resuscitation)</b>
<b>Periventricular Leukomalacia (PVL)</b>
<b>Spina Bifida</b>
<b>Spinocerebellar Disorders</b>
<b><i>Severe Attachment Disorders</i></b>
<b>Anxiety Disorders of Infancy and Early Childhood</b>
<b>Depression of Infancy and Early Childhood</b>
<b>Infantile Anorexia</b>
<b><i>Autism Spectrum Disorders</i></b>
<b>Asperger's Disorder</b>
<b>Autism Spectrum Disorder</b>
<b>Pervasive Developmental Disorder</b>
<b>Rett's Syndrome</b>
<b><i>Significant Sensory Impairment</i></b>
<b>Auditory Neuropathy</b>
<b>Aural Atresia (bilateral or unilateral)</b>
<b>Blindness ("legal" blindness or 20/200 best acuity with correction)</b>
<b>Optic Nerve Hypoplasia (De Morsier's Syndrome)</b>
<ul style="list-style-type: none"> <li>• <b>Septo Optic Dysplasia</b></li> </ul>
<b>Retinopathy of Prematurity Stage III and/or IV (ROP)</b>
<ul style="list-style-type: none"> <li>• <b>Stage 3 unspecified</b> <ul style="list-style-type: none"> <li>○ <b>Bilateral</b></li> <li>○ <b>Left eye</b></li> <li>○ <b>Right eye</b></li> </ul> </li> <li>• <b>Stage 4 unspecified</b> <ul style="list-style-type: none"> <li>○ <b>Bilateral</b></li> <li>○ <b>Left eye</b></li> <li>○ <b>Right eye</b></li> </ul> </li> <li>• <b>Stage 5 unspecified</b> <ul style="list-style-type: none"> <li>○ <b>Bilateral</b></li> <li>○ <b>Left eye</b></li> <li>○ <b>Right eye</b></li> </ul> </li> </ul>
<b>Sensorineural hearing loss in excess of 25 dB HL</b>
<b><i>Other</i></b>
<b>Fetal Alcohol Syndrome</b>
<b>Hydrocephalus (congenital or acquired)</b>
<b>Lead Poisoning</b>
<b>Low Birth Weight (&lt;1,200 grams at birth)</b>
<b>Zika Confirmed Congenital with Symptoms</b>
<b>Zika Confirmed Congenital No Symptoms</b>
<b>Zika Probable Congenital with Symptoms</b>
<b>Zika Probable Congenital No Symptoms</b>