

<b>Established Conditions</b> (Not an exhaustive list)	
<i>Genetic and Metabolic Disorders</i>	<i>ICD10 Code</i>
<b>Albinism</b>	E70.30
<b>Albright's Hereditary Osteodystrophy</b>	E20.1
<b>Angelman Syndrome (Happy Puppet Syndrome)</b>	Q93.5
<b>Adrenoleukodystrophy</b>	E71.529
<b>Antley-Bixler Syndrome (Multisynostotic Osteodysgenesis, Craniosynostosis, Choanal Atresia, Radial Humeral Synostosis, Trapezoidocephaly-Multiple Synostosis Syndrome, ABS, Multisynostotic Osteodysgenesis with Long Bone Fractures)</b>	Q87.5
<b>Apert Syndrome (Acrocephalosyndactyly)</b>	Q87.0
<b>Arthrogryposis Multiplex Congenita</b>	Q74.3
<b>Ataxia-Telangiectasia Syndrome (Louis-Bar Syndrome)</b>	G11.3
<b>Canavan Disease</b>	E75.29
<b>Cardio-Facio-Cutaneo Syndrome</b>	Q87.89
<b>Cerebral Lipidosis</b>	E75.6
<b>Cerebro-Oculo-Facio-Skeletal (COFS) Syndrome</b>	Q87.8
<b>CHARGE Syndrome/Association</b>	Q89.8
<b>Chromosome Syndromes 10p+, 13q+, 3q+, 4Q+</b>	Q92.5
<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>	Q93.89
<b>Coffin-Lowry Syndrome</b>	Q89.8
<b>Coffin-Siris Syndrome</b>	Q03.1
<b>Cornelia de Lange Syndrome (Brachmann de Lange)</b>	Q87.1
<b>Cri-du-chat Syndrome (Deletion 5p Syndrome)</b>	Q93.4
<b>Cystic Fibrosis</b>	E84.0
<b>Dandy Walker Syndrome</b>	Q03.1
<b>Down Syndrome (Trisomy 21)</b>	Q90.9
<b>Duchenne Muscular Dystrophy</b>	G71.0
<b>Dyggve-Melchior-Clausen Syndrome (DMC Disease, DMC Syndrome, Smith-McCort Dysplasia)</b>	Q77.7
<b>Fragile X Syndrome</b>	Q99.2
<b>Fraser Syndrome (Cryptophthalmos Syndrome, Meyer-Schwickerath's syndrome, Fraser-Francois syndrome, Ullrich-Feichtiger syndrome)</b>	Q87.0
<b>Galactosemia</b>	E74.21
<b>Gaucher Syndrome (Glucosylceramide storage disease; GSDI)</b>	E75.22
<b>Glutaric Aciduria</b>	
Type I	E72.3
Type II	E71.313
<b>Glycogen Storage Disease</b>	E74.00
<b>Jeune Syndrome</b>	Q77.2
<b>Joubert Syndrome</b>	Q03.1
<b>Krabbe's disease</b>	E75.23
<b>Lesch-Nyhan Syndrome</b>	E79.1
<b>Lissencephaly Syndrome (Miller-Dieker Syndrome, Agyria)</b>	Q93.88
<b>Maple Syrup Urine</b>	E71.0

<b>Mucopolysaccharidosis II, III</b>	E77.0
<b>Organic Acidemias</b>	E71.121
<b>Pelizaeus-Merzbacher disease</b>	E75.29
<b>Peroxisomal Disorders</b>	E71.5
<b>Phenylketonuria (PKU)</b>	E70.0
<b>Phelan-McDermid syndrome</b>	Q93.0
<b>Rubenstein-Taybi Syndrome</b>	Q87.2
<b>Schwartz-Jampel Syndrome</b>	G71.13
<b>Prader-Willi Syndrome</b>	Q87.1
<b>Steinert Myotonic Dystrophy Syndrome (Curschmann-Batten-Steinert syndrome)</b>	G71.11
<b>Tay-Sachs disease (Sandhoff)</b>	E75.02
<b>Trisomy 8</b>	Q92.9
<b>Trisomy 9</b>	Q92.9
<b>Tetrasomy 12p</b>	Q99.8
<b>Trisomy 13 (Patau Syndrome)</b>	Q91.7
<b>Trisomy 18 (Edward's Syndrome)</b>	Q91.3
<b>Tuberous Sclerosis Complex</b>	Q85.1
<b>Urea Cycle Defect</b>	E72.20
<b>Very long chain fatty acid storage diseases</b>	E71.310
<b>Walker-Warburg Syndrome (XO)</b>	G71.2
<b>Williams Syndrome</b>	Q99.8
<b>Zellweger Syndrome (Cerebro-Hepato-Renal Syndrome)</b>	Q87.89
<b>Neurological Disorders</b>	
<b>Agyria (Miller-Dieker lissencephaly syndrome (MDLS), agyria syndrome, agyria-pachygyria syndrome, classical lissencephaly)</b>	Q04.3
<b>Aicardi Syndrome</b>	Q04.0
<b>Alpers Syndrome/Disease</b>	G31.81
<b>Aphasia</b>	R47.01
<b>Arachnoid cyst with Neuro-Developmental Delay</b>	G93.0
<b>Arhinencephaly (Holoprosencephaly)</b>	Q04.1 Q04.2
<b>Arnold-Chiari Syndrome, type II (Malformation d'Arnold-Chiari)</b>	Q07.00
<b>Ataxia</b>	R27.0
<b>Cerebral Palsy</b>	G80.9
<b>Cerebral Aneurysm with Neuro-Developmental Delay</b>	I67.1
<b>CNS Tumor with Neuro- Developmental Delay</b>	D49.7
<b>Encephalopathy, Congenital Only</b>	G93.40
<b>Encephalopathy, Static</b>	G93.40
<b>Erb's Palsy (Brachial Plexus Injury, Perinatal Origin)</b>	P14.0
<b>Extracorporeal Membrane Oxygenation (ECMO)</b>	Z92.81
<b>Holoprosencephaly</b>	Q04.2
<b>Hypertonia (persistent only)</b>	P94.1
<b>Hypoxic Ischemic Encephalopathy (HIE)</b>	P91.60
<b>Lennox-Gastaut Syndrome</b>	G40.812
<b>Intracranial Calcifications</b>	93.89
<b>Intraventricular Hemorrhage</b>	
<b>Grade 3</b>	P52.21
<b>Grade 4</b>	P52.22
<b>Meningocele (cervical)</b>	Q05.5

Microcephaly	Q02
Miller-Dieker Syndrome	Q93.88
Mitochondrial Disorder	E88.40
Multiple Anomalies of the Brain	Q04.9
Myopathy	G72.89
Neonatal/Perinatal Asphyxia (5 minute Apgar score of 6 or less, Cord PH < 7, Evidence of Central Nervous System involvement, Organ failure, Resuscitation)	P84
Periventricular Leukomalacia (PVL)	91.2
Spina Bifida	Q05.9
Spinocerebellar Disorders	G11.8
<b>Severe Attachment Disorders</b>	
Anxiety Disorders of Infancy and Early Childhood	F41.1
Depression of Infancy and Early Childhood	F33.40
Infantile Anorexia	R63.0
<b>Autism Spectrum Disorders</b>	
Asperger's Disorder	F84.5
Autism Spectrum Disorder	F84.0
Pervasive Developmental Disorder	F84.9
Rett's Syndrome	F84.2
<b>Significant Sensory Impairment</b>	
Auditory Neuropathy	H93.299
Aural Atresia (bilateral or unilateral)	Q16.1
Blindness ("legal" blindness or 20/200 best acuity with correction)	H54.8
Optic Nerve Hypoplasia (De Morsier's Syndrome)	H47.039
• Septo Optic Dysplasia	Q04.4
Retinopathy of Prematurity Stage III and/or IV (ROP)	
• Stage 3 unspecified	H35.149
○ Bilateral	H35.143
○ Left eye	H35.142
○ Right eye	H35.141
• Stage 4 unspecified	H35.159
○ Bilateral	H35.153
○ Left eye	H35.152
○ Right eye	H35.151
• Stage 5 unspecified	H35.169
○ Bilateral	H35.163
○ Left eye	H35.162
○ Right eye	H35.161
Sensorineural hearing loss in excess of 25 dB HL	H90.5
<b>Other</b>	
Fetal Alcohol Syndrome	Q86.0
Hydrocephalus (congenital or acquired)	G91.9
Lead Poisoning	R78.71
Low Birth Weight (<1,200 grams at birth)	P07.00
Zika Confirmed Congenital with Symptoms	A92.5
Zika Confirmed Congenital No Symptoms	A92.5
Zika Probable Congenital with Symptoms	A92.5
Zika Probable Congenital No Symptoms	A92.5