

18 Q Deletion	Menkes Syndrome
49 XXXXY Syndrome	Methylmalonic acidemia
Acardi Syndrome	Microphthalmia Moebius Sequence
Albinism	Muscular Dystrophy, Becker Type
All unbalanced structural chromosome syndrome	Muscular Dystrophy, Duchenne Type
Amyoplasia Congenita	Myopathies Neurofibromatosis Type 1
Angelman Syndrome	NICU Grad., Difficulty pacing/coordinating suck-swallow-breathe
Aniridia	NICU Grad., Fluctuating tone
Anophthalmia	NICU Grad., Inconsolability
Apert Syndrome	NICU Grad., Long, difficult time to learn to eat
Anoxic Brain Injury	NICU Grad., Neurologically based significant irritability
Arthrogryposis	NICU Grad., NG, NJ, or G tube feedings required either full or partial to maintain adequate nutrition
Autistic Spectrum Disorders	NICU Grad., Severe sleep disorder
Bardet-Biedl Syndrome	NICU Grad., Significant tremors when at rest
Brachial Plexopathy	NICU Grad., Unable to come to a quiet-alert state
Cerebral Palsy	NICU Grad., Unable to take 100% nutrition by mouth
CHARGE Syndrome	NICU Grad., Unusually high tone
Chromosome 9 Deletion	NICU Grad., Unusually low tone
Cockayne Syndrome	Niemann-Pick Disease
Congenital Bilateral Perisylvian	Optic Atrophy
Congenital Brain Malformation	Optic Nerve Hypoplasia
Congenital Infection, Cytomegalovirus	Osteogenesis imperfecta
Congenital Infection, Herpes	Pentasomy X
Congenital Infection, HIV	Persistent Hyperplastic Primary Vitreous (PHPV)
Congenital Infection, Rubella	Pervasive Developmental Delay
Congenital Infection, Syphilis	Phthisis Bulbi
Congenital Infection, Toxoplasmosis	Prader-Willi Syndrome
Cornelia de Lange Syndrome	Pierre-Robin Sequence
Cortical Visual Impairment	Reduction Deformity
Cri-du-chat	Retinal Detachment
Deafblind Delayed	Retinopathy of Prematurity (ROP) Grades 4,5
Visual Development/Maturation/Impairment	Rubenstein-Taybi Syndrome
Deletion 2p21	Sanfilippo Syndrome
DiGeorge Syndrome or Velo-Cardio-Facial	Schinzel-Giedion Syndrome
Down Syndrome (Trisomy 21)	Semilobar holoprosencephaly
Encephalocele	Septo Optic Dysplasia
Failure to Thrive	Sly Syndrome
Familial Exudative Vitreoretinopathy (FEVR)	Spina Bifida
Fetal Alcohol Syndrome	Spinal Cord Injury with Cord Involvement
Fetal Hydantoin Syndrome	Spinal Muscular Atrophy (SMA)
Fetal Valproate Syndrome	Sturge-Weber
FG Syndrome	Tay-Sachs Disease
Fragile X Syndrome	Tethered Cord Syndrome (TCS)
Goldenhar Syndrome	Treacher Collins
Hearing Loss	Trisomy 13
Hunter Syndrome	Trisomy 18
Hurler-Scheie Syndrome	Tuberous Sclerosis
Hypoxic Ischemic Encephalopathy (HIE)	Untreated Hypothyroidism
Infantile Gaucher Disease	Untreated PKU
Inflicted Neurotrauma	VACTERL Association
Jacobsen Syndrome	Velo-cardio-facial or DiGeorge Syndrome
Kleefstra Syndrome	Ventilator Dependent
Klinefelter Syndrome	Walker-Warburg Syndrome
Kugelburg-Wehlander Syndrome	Waardenburg Syndrome, Types I and II
Leber's Congenital Amaurosis	Werdnig-Hoffman Syndrome
Lesch-Nyhan Syndrome	Williams Syndrome
Lowe Syndrome	Wolf-Hirschhorn Syndrome
Maple Syrup Urine Disease	
Marshall-Smith Syndrome	