ICD10	Description	Type	Eligibility	Secondary
E71.111	3-methylglutaconic aciduria	D	1	1
Q77.0	Achondrogenesis	D	1	1
Q77.4	Achondroplasia	D	1	1
I67.81	Acute cerebrovascular insufficiency	D	1	1
R48.1	Agnosia	D	1	1
G31.81	Alpers disease	D	1	1
Q87.81	Alport syndrome	D	1	1
Q11.1	anopthalmia, bilateral agenesis of the eye	D	1	1
Q87.0	Apert syndrome	D	1	1
R48.2	Apraxia	D	1	1
E72.21	Argininemia	D	1	1
E72.22	Arginosuccinic aciduria	D	1	1
Q04.1	Arhinencephaly	D	1	1
Q74.3	Arthrogryposis multiplex congenita	D	1	1
F84.5	Asperger's syndrome	D	1	1
G80.4	Ataxic cerebral palsy	D	1	1
Q03.1	Atresia of foramina of Magendie and Luschka	D	1	1
E78.71	Barth syndrome	D	1	1
Q87.3	Beckwith-Wiedeman syndrome	D	1	1
H54.4	Blindness & low vision- monocular	D	1	1
Q76.49	Caudal Regression Syndrome	D	1	1
I67.82	Cerebral ischemia	D	1	1
G80.9	cerebral palsy, unspecified	D	1	1
Q89.8	CHARGE syndrome	D	1	1
E70.330	Chediak-Higashi syndrome	D	1	1
F84.0	Childhood Autism	D	1	1
F80.81	Childhood onset fluency disorder	D	1	1
Q99.9	Chromosomal abnormality, unspecified	E	1	1
Q99.9	Chromosomal abnormality, unspecified	D	1	1
E72.23	Citrullinemia	D	1	1
Q37.4	Cleft hard and soft palate with bilateral cleft lip	D	1	1
Q37.5	Cleft hard and soft palate with unilateral cleft lip	D	1	1
Q37.0	Cleft hard palate with bilateral cleft lip	D	1	1

Q37.1	Cleft hard palate with unilateral cleft lip	D	1	1
Q35.9	Cleft palate, unspecified	D	1	1
Q37.2	Cleft soft palate with bilateral cleft lip	D	1	1
Q37.3	Cleft soft palate with unilateral cleft lip	D	1	1
Z13.4	Clinical Opinion of Dev Delay Only	O	1	0
H90.0	Conductive hearing loss, bilateral	D	1	1
H90.12	Conductive hearing loss, unilateral, left ear, with unrestri	D	1	1
H90.11	Conductive hearing loss, unilateral, right ear, with unrestr	D	1	1
H90.2	Conductive hearing loss, unspecified	D	1	1
Q04.6	Congenital cerebral cysts	D	1	1
P35.2	Congenital herpes viral [herpes simplex] infection	D	1	1
Q03.9	Congenital hydrocephalus, unspecified	D	1	1
P94.1	Congenital Hypertonia	E	0	1
Q04.9	Congenital malformation of brain, unspecified	D	1	1
Q06.9	Congenital malformation of spinal cord, unspecified	D	1	1
Q87.1	Congenital malformation syndromes predominantly assoc	D	1	1
Q04.0	Congenital malformations of corpus callosum	D	1	1
P37.1	Congenital toxoplasmosis	D	1	1
H47.612	Cortical blindness, left side of brain	D	1	1
H47.611	Cortical blindness, right side of brain	D	1	1
H47.619	Cortical blindness, unspecified side of brain	D	1	1
P35.1	cytomegalovirus, congenital	D	1	1
H91.3	Deaf nonspeaking, not elsewhere classified	D	1	1
R62.0	Delayed Milestones	E	1	1
F80.9	Developmental disorders of speech and language, unspecified	E	1	1
D82.1	Di George's syndrome	D	1	1
Q77.5	Diastrophic dysplasia	D	1	1
E70.9	Disorder of aromatic amino-acid metabolism, unspecified	D	1	1
E71.2	Disorder of branched-chain amino-acid metabolism, unspecifie	D	1	1
E72.20	Disorder of urea cycle metabolism, unspecified	D	1	1
E74.20	Disorders of galactose metabolism, unspecified	D	1	1
E71.32	Disorders of ketone metabolism	D	1	1
E72.4	Disorders of ornithine metabolism	D	1	1
E70.5	Disorders of tryptophan metabolism	D	1	1

Q93.5	Duplication of Chromosome 15	D	1	1
Q78.4	Enchondromatosis	D	1	1
G40.911	Epilepsy, unspecified, intractable, with status epi	D	1	1
G40.919	Epilepsy, unspecified, intractable, without status	D	1	1
F80.1	Expressive language disorder	E	1	1
P07.22	Extreme immaturity of newborn, GA 23 completed wks	D	1	1
P07.23	Extreme immaturity of newborn, GA 24 completed wks	D	1	1
P07.24	Extreme immaturity of newborn, GA 25 completed wks	D	1	1
P07.25	Extreme immaturity of newborn, GA 26 completed wks	D	1	1
P07.26	Extreme immaturity of newborn, GA 27 completed wks	D	1	1
P07.21	Extreme immaturity of newborn, GA less than 23 completed wks	D	1	1
P07.02	Extremely low birth weight newborn, 500-749 grams	D	1	1
P07.03	Extremely low birth weight newborn, 750-999 grams	D	1	1
P07.01	Extremely low birth weight newborn, less than 500 grams	D	1	1
F84.8	F84.8 - Other pervasive developmental disorders	D	1	1
R63.3	Feeding difficulties	E	1	1
Q86.0	Fetal alcohol syndrome (dysmorphic)	D	1	1
Q99.2	Fragile X chromosome	D	1	1
E75.22	Gaucher's Disease	D	1	1
F88	Global developmental delay	E	1	1
E71.313	Glutaric aciduria type II	D	1	1
E75.00	GM2 gangliosidosis, unspecified	D	1	1
E70.331	Hermansky-Pudlak syndrome	D	1	1
Q04.2	Holoprosencephaly	D	1	1
G91.9	Hydrocephalus, unspecified	D	1	1
P91.61	hypoxic ischemic encephalopathy	D	1	1
G40.822	Infantile Spasms/West Syndrome	D	1	1
F81.9	Intellectual delay	E	1	1
Q93.88	interstitial microdeletion, incl 14q22-23	D	1	1
G40.909	intractable epilepsy	D	1	1
P52.21	Intraventricular (nontraumatic) hemorrhage, grade 3, of newb	D	1	1
P52.22	Intraventricular (nontraumatic) hemorrhage, grade 4, of newb	D	1	1
P52.21	intraventricular hemorrhage of newborn, grade 3	D	1	1
P52.22	intraventricular hemorrhage of newborn, grade 4	D	1	1

E71.110	Isovaleric acidemia	D	1	1
Q96.0	Karyotype 45, X	D	1	1
Q96.1	Karyotype 46, X iso (Xq)	D	1	1
Q96.2	Karyotype 46, X with abnormal sex chromosome, except	D	1	1
H54.8	Legal blindness, as defined in USA	D	1	1
G31.82	Leigh's disease	D	1	1
E88.1	Lipodystrophy, congenital, not otherwise classified	D	1	1
Q04.3	Lissencephaly	D	1	1
E71.310	Long chain/very long chain acyl CoA dehydrogenase deficiency	D	1	1
Q03.0	Malformations of aqueduct of Sylvius	D	1	1
E71.0	Maple-syrup-urine disease	D	1	1
E71.311	Medium chain acyl CoA dehydrogenase deficiency	D	1	1
Q04.5	Megalencephaly	D	1	1
E71.120	Methylmalonic acidemia	D	1	1
Q02	Microcephaly	D	1	1
H90.6	Mixed conductive and sensorineural hearing loss, bilateral	D	1	1
H90.71	Mixed conductive and sensorineural hearing loss, unilateral,	D	1	1
H90.72	Mixed conductive and sensorineural hearing loss, unilateral,	D	1	1
H90.8	Mixed conductive and sensorineural hearing loss, unspecified	D	1	1
F80.2	mixed receptive-expressive language disorder	E	1	1
Q96.3	Mosaicism, 45, X/46, XX or XY	D	1	1
Q96.4	Mosaicism, 45, X/other cell line(s) with abnormal sex	D	1	1
E75.11	Mucolipidosis IV	D	1	1
E71.314	Muscle carnitine palmitoyltransferase deficiency	D	1	1
P96.1	Neonatal Abstinence syndrome- first 3 months of life only	D	1	1
P91.2	Neonatal cerebral leukomalacia	D	1	1
Q85.09	Neurofibromatosis	D	1	1
E75.4	Neuronal ceroid lipofuscinosis	D	1	1
P04.3	Newborn (suspected to be) affected by maternal use of alcoho	D	1	1
Q87.1	Noonan syndrome	D	1	1
G91.1	Obstructive hydrocephalus	D	1	1
Q77.9	Osteochondrodysplasia with defects of growth of tubular bone	D	1	1
Q78.0	Osteogenesis imperfecta	D	1	1
E71.118	Other branched-chain organic acidurias	D	1	1

F84.3	other childhood disintegrative disorder	D	1	1
F94.8	Other childhood disorder of social functioning	E	1	1
Q03.8	Other congenital hydrocephalus	D	1	1
Q86.8	Other congenital malformation syndromes due to known	D	1	1
Q87.5	Other congenital malformation syndromes with other sk	D	1	1
E70.8	Other disorders of aromatic amino-acid metabolism	D	1	1
G94	Other disorders of brain in diseases classified elsewhere	D	1	1
E71.19	Other disorders of branched-chain amino-acid metabolism	D	1	1
E71.318	Other disorders of fatty-acid oxidation	D	1	1
E74.29	Other disorders of galactose metabolism	D	1	1
E71.128	Other disorders of propionate metabolism	D	1	1
E72.29	Other disorders of urea cycle metabolism	D	1	1
E75.19	Other gangliosidosis	D	1	1
E75.09	Other GM2 gangliosidosis	D	1	1
E70.318	Other ocular albinism	D	1	1
Q77.8	Other osteochondrodysplasia with defects of growth of tubula	D	1	1
Q04.3	Other reduction deformities of brain	D	1	1
Q87.89	Other specified congenital malformation syndromes, n	D	1	1
Q04.8	Other specified congenital malformations of brain	D	1	1
G93.89	Other specified disorders of brain	D	1	1
H91.8X3	Other specified hearing loss, bilateral	D	1	1
H91.8X2	Other specified hearing loss, left ear	D	1	1
H91.8X1	Other specified hearing loss, right ear	D	1	1
H91.8X9	Other specified hearing loss, unspecified ear	D	1	1
Q96.8	Other variants of Turner's syndrome	D	1	1
H91.03	Ototoxic hearing loss, bilateral	D	1	1
H91.02	Ototoxic hearing loss, left ear	D	1	1
H91.01	Ototoxic hearing loss, right ear	D	1	1
H91.09	Ototoxic hearing loss, unspecified ear	D	1	1
P14.9	Paralysis; birth injury to peripheral nervous system,	D	1	1
I63.9	Periventricular Leukomalacia	D	1	1
Z86.59	Personal history of other mental and behavioral disorders	D	1	1
Z87.820	Personal History of Traumatic Brain Injury	D	1	1
F84.9	Pervasive developmental disorder, unspecified	D	1	1

Q79.8	Poland syndrome	D	1	1
G91.3	Post-traumatic hydrocephalus, unspecified	D	1	1
Q87.1	Prader-Willi syndrome	D	1	1
P07.31	Preterm newborn, gestational age 28 completed weeks	D	1	1
E71.121	Propionic acidemia	D	1	1
Q79.4	Prune belly syndrome	D	1	1
C69.20	Retinoblastoma	D	1	1
H35.173	Retrolental fibroplasia, bilateral	D	1	1
H35.172	Retrolental fibroplasia, left eye	D	1	1
H35.171	Retrolental fibroplasia, right eye	D	1	1
H35.179	Retrolental fibroplasia, unspecified eye	D	1	1
F84.2	Rett's syndrome	D	1	1
E75.01	Sandhoff disease	D	1	1
H90.3	Sensorineural hearing loss, bilateral	D	1	1
H90.42	Sensorineural hearing loss, unilateral, left ear, with unres	D	1	1
H90.41	Sensorineural hearing loss, unilateral, right ear, with unre	D	1	1
Q04.4	Septo-optic dysplasia of brain	D	1	1
E71.312	Short chain acyl CoA dehydrogenase deficiency	D	1	1
E78.72	Smith-Lemli-Opitz syndrome	D	1	1
G80.2	Spastic hemiplegic cerebral palsy	D	1	1
G80.0	Spastic quadriplegic cerebral palsy	D	1	1
F82	Specific developmental disorder of motor function	E	1	1
F80.0	Speech Sound Disorder	D	1	1
Q05.4	Spina Bifida with hydrocephalus	D	1	1
Q05.2	Spina Bifida, myelomeningocele - with hydrocephalus	D	1	1
Q05.8	Spina Bifida, myelomeningocele - without hydrocephalus	D	1	1
Q06.9	Spinal Cord, tethered/congenital malformation	D	1	1
Q77.7	Spondyloepiphyseal dysplasia	D	1	1
I63.9	Stroke, cerebral infarction	D	1	1
E75.02	Tay-Sachs disease	D	1	1
T56.0X1A	Toxic effect of lead and its compounds, accidental (unintent	D	1	1
S09.90XA	Traumatic head injury	D	1	1
Q92.9	Trisomies and partial trisomies, unspecified	D	1	1
Q91.5	Trisomy 13, mosaicism (mitotic nondisjunction)	D	1	1

Q91.4	Trisomy 13, nonmosaicism (meiotic nondisjunction)	D	1	1
Q91.6	Trisomy 13, translocation	D	1	1
Q91.7	Trisomy 13, unspecified	D	1	1
Q91.1	Trisomy 18, mosaicism (mitotic nondisjunction)	D	1	1
Q91.0	Trisomy 18, nonmosaicism (meiotic nondisjunction)	D	1	1
Q91.2	Trisomy 18, translocation	D	1	1
Q91.3	Trisomy 18, unspecified	D	1	1
Q90.9	Trisomy 21 (Down Syndrome) unspecified	D	1	1
Q 92.8	Trisomy, chromosome specified	D	1	1
Q85.1	Tuberous sclerosis	D	1	1
Q96.9	Turner's syndrome, unspecified	D	1	1
E70.21	Tyrosinemia	D	1	1
Q37.8	Unspecified cleft palate with bilateral cleft lip	D	1	1
Q37.9	Unspecified cleft palate with unilateral cleft lip	D	1	1
R62.50	Unspecified Delay in Development	E	1	1
E75.10	Unspecified gangliosidosis	D	1	1
H90.5	Unspecified sensorineural hearing loss	D	1	1
A92.5	Zika syndrome	D	1	1