- D82.1 Di George’s Syndrome
- E75.0 GM2 gangliosidosis
- E75.00 GM2 gangliosidosis, unspecified
- E75.01 Sandhoff disease
- E75.02 Tay-Sachs disease
- E75.09 Other GM2 gangliosidosis
- E75.1 Other and unspecified gangliosidosis
- E75.10 Unspecified gangliosidosis
- E75.11 Mucolipidosis IV
- E75.19 Other gangliosidosis
- E75.2 Other sphingolipidosis
- E75.21 Fabry (-Anderson) disease
- E75.22 Gaucher disease
- E75.23 Krabbe disease
- E75.24 Niemann-Pick disease
- E75.240 Niemann-Pick disease type A
- E75.241 Niemann-Pick disease type B
- E75.242 Niemann-Pick disease type C
- E75.243 Niemann-Pick disease type D
- E75.248 Other Niemann-Pick disease
- E75.249 Niemann-Pick disease, unspecified
- E75.25 Metachromatic leukodystrophy
- E75.26 Sulfatase deficiency
- E75.29 Other sphingolipidosis
- E75.3 Sphingolipidosis, unspecified
- E75.4 Neuronal ceroid lipofuscinosis
- E75.5 Other lipid storage disorders
- E75.6 Lipid storage disorder, unspecified
- E76 Disorders of glycosaminoglycan metabolism
- E76.0 Mucopolysaccharidosis, type I
- E76.01 Hurler's syndrome
- E76.02 Hurler-Scheie syndrome
- E76.03 Scheie's syndrome
- E78.71 Barth syndrome
- E78.72 Smith-Lemli-Opitz syndrome
- G11.1 Early-onset cerebellar ataxia
- G12.0 Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]
- G12.1 Other inherited spinal muscular atrophy
- G31.84 Mild cognitive impairment, so stated
- G71.0 Muscular Dystrophy
- G71.00 Muscular dystrophy, unspecified
- G71.01 Duchenne or Becker muscular dystrophy
- G71.02 Facioscapulohumeral muscular dystrophy
- G71.09 Other specified muscular dystrophy
- G71.11 Myotonic muscular dystrophy
- G71.12 Myotonia congenita
- G71.13 Myotonic chondrodystrophy

- G71.14 Drug induced myotonia
- G71.19 Other specified myotonic disorders
- G71.2 Congenital myopathies
- G80.0 Spastic quadriplegic cerebral palsy
- G80.1 Spastic diplegic cerebral palsy
- G80.2 Spastic hemiplegic cerebral palsy
- G80.3 Athetoid cerebral palsy
- G80.4 Ataxic cerebral palsy
- G80.8 Other cerebral palsy
- G80.9 Cerebral palsy, unspecified
- G82.51 Quadriplegia, C1-C4 complete
- G91.0 Communicating hydrocephalus
- G91.1 Obstructive hydrocephalus
- G91.3 Post-traumatic hydrocephalus, unspecified
- G91.4 Hydrocephalus in diseases classified elsewhere
- G91.8 Other hydrocephalus
- G91.9 Hydrocephalus, unspecified
- G93.1 Anoxic brain damage, not elsewhere classified
- G93.40 Encephalopathy, unspecified
- G93.5 Compression of brain
- G93.6 Cerebral edema
- G93.7 Reye's syndrome
- G93.89 Other specified disorders of brain
- G93.9 Disorder of brain, unspecified
- G96.9 Disorder of central nervous system, unspecified
- G98.8 Other disorders of nervous system
- P07.3 Preterm newborn
- P07.30 Preterm newborn, unspecified weeks of gestation
- P07.31 Preterm newborn, gestational age 28 completed weeks
- P07.32 Preterm newborn, gestational age 29 completed weeks
- P07.33 Preterm newborn, gestational age 30 completed weeks
- P07.34 Preterm newborn, gestational age 31 completed weeks
- P07.35 Preterm newborn, gestational age 32 completed weeks
- P07.36 Preterm newborn, gestational age 33 completed weeks
- P07.37 Preterm newborn, gestational age 34 completed weeks
- P07.38 Preterm newborn, gestational age 35 completed weeks
- P07.39 Preterm newborn, gestational age 36 completed weeks
- P83.2 Hydrops fetalis not due to hemolytic disease
- P92 Feeding problems of newborn
- Q01.0 Frontal encephalocele
- Q01.1 Nasofrontal encephalocele
- Q01.2 Occipital encephalocele
- Q01.8 Encephalocele of other sites
- Q01.9 Encephalocele, unspecified
- Q02 Microcephaly
- Q03.0 Malformations of aqueduct of Sylvius
- Q03.1 Atresia of foramina of Magendie and Luschka

- Q03.8 Other congenital hydrocephalus
- Q03.9 Congenital hydrocephalus, unspecified
- Q04.0 Congenital malformations of corpus callosum
- Q04.1 Arhinencephaly
- Q04.2 Holoprosencephaly
- Q04.3 Other reduction deformities of brain
- Q04.4 Septo-optic dysplasia of brain
- Q04.5 Megalencephaly
- Q04.6 Congenital cerebral cysts
- Q04.8 Other specified congenital malformations of brain
- Q04.9 Congenital malformation of brain, unspecified
- Q05.0 Cervical spina bifida with hydrocephalus
- Q05.1 Thoracic spina bifida with hydrocephalus
- Q05.2 Lumbar spina bifida with hydrocephalus
- Q05.3 Sacral spina bifida with hydrocephalus
- Q05.4 Unspecified spina bifida with hydrocephalus
- Q05.5 Cervical spina bifida without hydrocephalus
- Q05.6 Thoracic spina bifida without hydrocephalus
- Q05.7 Lumbar spina bifida without hydrocephalus
- Q05.8 Sacral spina bifida without hydrocephalus
- Q05.9 Spina bifida, unspecified
- Q06 Other congenital malformations of spinal cord
- Q06.0 Amyelia
- Q06.1 Hypoplasia and dysplasia of spinal cord
- Q06.2 Diastematomyelia
- Q06.3 Other congenital cauda equina malformations
- Q06.4 Hydromyelia
- Q06.8 Other specified congenital malformations of spinal cord
- Q06.9 Congenital malformation of spinal cord, unspecified
- Q07.00 Arnold-Chiari syndrome without spina bifida or hydrocephalus
- Q07.01 Arnold-Chiari syndrome with spina bifida
- Q07.03 Arnold-Chiari syndrome with spina bifida and hydrocephalus
- Q07.8 Other specified congenital malformations of nervous system
- Q07.9 Congenital malformation of nervous system, unspecified
- Q90.0 Trisomy 21, nonmosaicism (meiotic nondisjunction)
- Q90.1 Trisomy 21, mosaicism (mitotic nondisjunction)
- Q90.2 Trisomy 21, translocation
- Q90.9 Down syndrome, unspecified
- Q91.0 Trisomy 18, nonmosaicism (meiotic nondisjunction)
- Q91.1 Trisomy 18, mosaicism (mitotic nondisjunction)
- Q91.2 Trisomy 18, translocation
- Q91.3 Trisomy 18, unspecified
- Q91.4 Trisomy 13, nonmosaicism (meiotic nondisjunction)
- Q91.5 Trisomy 13, mosaicism (mitotic nondisjunction)
- Q91.6 Trisomy 13, translocation
- Q91.7 Trisomy 13, unspecified
- Q92.0 Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)

- Q92.1 Whole chromosome trisomy, mosaicism (mitotic nondisjunction)
- Q92.2 Partial trisomy
- Q92.5 Duplications with other complex rearrangements
- Q92.6 Marker chromosomes
- Q92.61 Marker chromosomes in normal individual
- Q92.62 Marker chromosomes in abnormal individual
- Q92.7 Triploidy and polyploidy
- Q92.8 Other specified trisomies and partial trisomies of autosomes
- Q92.9 Trisomy and partial trisomy of autosomes, unspecified
- Q93 Monosomies and deletions from the autosomes, not elsewhere classified
- Q93.0 Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)
- Q93.1 Whole chromosome monosomy, mosaicism (mitotic nondisjunction)
- Q93.2 Chromosome replaced with ring, dicentric or isochromosome
- Q93.3 Deletion of short arm of chromosome 4
- Q93.4 Deletion of short arm of chromosome 5
- Q93.5 Other deletions of part of a chromosome
- Q93.51 Angelman syndrome
- Q93.59 Other deletions of part of a chromosome
- Q93.7 Deletions with other complex rearrangements
- Q93.8 Other deletions from the autosomes
- Q93.81 Velo-cardio-facial syndrome
- Q93.82 Williams syndrome
- Q93.88 Other microdeletions
- Q93.89 Other deletions from the autosomes
- Q93.9 Deletion from autosomes, unspecified
- Q95.2 Balanced autosomal rearrangement in abnormal individual
- Q95.3 Balanced sex/autosomal rearrangement in abnormal individual
- Q99.2 Fragile X chromosome
- Q99.8 Other specified chromosome abnormalities
- Q99.9 Chromosomal abnormality, unspecified
- R27.9 Unspecified lack of coordination
- R62.0 Delayed milestone in childhood
- R62.50 Unspecified lack of expected normal physiological development in childhood
- R62.51 Failure to thrive (child)
- R62.59 Other lack of expected normal physiological development in childhood
- R63.3 Feeding difficulties
- T74.4XXA Shaken infant syndrome, initial encounter
- T74.4XXD Shaken infant syndrome, subsequent encounter
- T74.4XXS Shaken infant syndrome, sequela
- F70 Mild intellectual disabilities
- F71 Moderate intellectual disabilities
- F72 Severe intellectual disabilities
- F73 Profound intellectual disabilities
- F78 Other intellectual disabilities
- F79 Unspecified intellectual disabilities
- F82 Specific developmental disorder of motor function
- F84 Pervasive developmental disorders

- F84.0 Autistic disorder
- F84.2 Rett's syndrome
- F84.3 Other childhood disintegrative disorder
- F84.5 Asperger's syndrome
- F84.8 Other pervasive developmental disorders
- F84.9 Pervasive developmental disorder, unspecified
- F88 Other disorders of psychological development
- F89 Disorder of psychological development, unspecified
- F90.0 Attention deficit hyperactivity disorder, predominantly inattentive type
- F90.1 Attention deficit hyperactivity disorder, primarily hyperactive type
- F90.2 Attention deficit hyperactivity disorder, combined type
- F90.8 Attention deficit hyperactivity disorder, other
- F90.9 Attention deficit hyperactivity disorder, unspecified type
- F98.2 Other feeding disorders of infancy and childhood
- F98.9 Unspecified behavioral and emotional disorders with onset usually occurring in childhood and adolescence