EarlySteps Eligibility Criteria

ICD-9 to ICD-10 Crosswalk

Purpose: To assist early interventionists in identifying ICD-10 diagnosis codes using the EarlySteps eligibility criteria. The ICD-10 codes are required for documentation and for billing for Medicaid-paid services for dates of service beginning 10/1/2015. The list below replaces the eligibility criteria list found in earlier versions of the Practice Manual in Chapter 5 (July, 2010 version). The nature of ICD-10 allows for increased specificity in the use of diagnosis coding, therefore the most specific applicable code should always be used. The sequence of the criteria in the list generally follows the sequence of the "general categories" Practice Manual Chapter 5 with some additional categories added to assist in locating codes. If you have questions about specific criteria, please contact your regional coordinator.

Criteria Description	ICD 9	ICD 10	Criteria Description
I. Developmental Delay			
General Category—Specific delays in development	315		
Developmental speech/language disorder	315.3	F80.89 F80.9	Other developmental disorders of speech and language Developmental disorder of speech and language not otherwise specified
Expressive language delay	315.31	F80.1	Expressive language disorder
Mixed receptive and expressive language delay	315.32	F80.2 H93.25	Mixed receptive-expressive language disorder Central auditory processing disorder
Speech and language delay due to hearing loss	315.34	F80.4	Speech and language development delay due to hearing loss
Developmental Coordination Disorder	315.4	F82	Specific Developmental Disorder: Motor Function
Mixed Developmental Disorder	315.5	F82	Specific Developmental Disorder: Motor Function
Other specified delays in development	315.8	F88	Other disorders of psychological development
II. Use of Informed Clinical Opinion to Determine Eligibility			
Abnormal sensory-motor response Affective or social disorder/condition	783.42	R62.0	Delayed milestone in childhood
Oral-motor skills dysfunction, including feeding difficulties	783.3	R63.3	Feeding difficulties
III. Established Medical Conditions			
Genetic Disorders			
B. Chromosomal Abnormality Syndromes – General Category	758		
Down syndrome	758.0	Q90.9 Q90.2 Q90.0 Q90.1	Down syndrome, unspecified Trisomy 21, translocation Trisomy 21, nonmosaicism (meiotic nondisjunction) Trisomy 21, mosaicism (mitotic nondisjunction)
Trisomy 13	758.1	Q91.7	Trisomy 13, unspecified (Patau's syndrome)
Trisomy 18	758.2	Q91.3	Trisomy 18, unspecified
General Category Autosomal deletion syndromes	758.3_		
Cri-du-chat	758.31	Q93.4	Deletion of short arm of chromosome 5
Velo-cardio-facial syndrome (VCFS)	758.32	Q93.81	Velo-cardio-facial syndrome

Criteria Description	ICD 9	ICD 10	Criteria Description
Other micro-deletion syndromes: include Miller-Dieker and Smith-Magenis syndromes	758.33	Q93.88	Other microdeletions
DiGeorge Syndrome	279.11	D82.1	Di George's syndrome
Fragile X	759.83	Q99.2	Fragile x chromosome
Prader-Willi	759.81	Q87.1	Congenital malformation syndromes predominantly associated with short stature
Other conditions due to autosomal anomalies	758.5	Q92.8	Other specified trisomies and partial trisomies of autosomes
Other conditions due to chromosomal anomalies Conditions due to sex chromosome anomalies	758.8_ 758.81	 Q97.0 Q97.1 Q97.2 Q97.8 Q98.4 Q98.5 Q98.7 Q98.8 Q98.9 Q99.8	 Karyotype 47, xxx Female with more than three x chromosomes Mosaicism, lines with various numbers of x chromosomes Other specified sex chromosome abnormalities, female phenotype Klinefelter's Syndrome (XXY) Karyotype 47, xyy Male with sex chromosome mosaicism Other specified sex chromosome abnormalities, male phenotype Turner's Syndrome (XO) Other specified chromosome abnormalities
Conditions due to anomaly of unspecified chromosome (includes Williams Syndrome)	758.9	Q99.9	Chromosomal abnormality, unspecified
C. **Pre-natal exposures**			
Fetal alcohol syndrome	760.71	P04.3 Q86.0	Newborn affected by alcohol affecting fetus or newborn via placenta or breast by maternal use of alcohol Fetal alcohol syndrome (dysmorphic)
Fetal hydantoin syndrome/Other	760.79	P04.8	Newborn (suspected to be) affected by other maternal noxious substances
Narcotics exposure	760.72	P04.49	Newborn (suspected to be) affected by maternal use of other drugs of addiction
Hallucinogenic agent exposure	760.73	P04.49	Newborn (suspected to be) affected by maternal use of drugs of addiction
Cocaine exposure	760.75	P04.41	Newborn (suspected to be) affected by maternal use of cocaine
Anticonvulsant exposure	760.77	P04.1	Newborn (suspected to be) affected by oth maternal medication
Other Noxious influences affecting fetus or newborn via placenta or breast milk	760.79	P04.8 P04.40	Newborn (suspected to be) affected by other maternal noxious substances Newborn affected by maternal use of unspecified drugs of addiction
Drug Withdrawal Syndrome	779.5	P96.1 P96.2	Neonatal withdrawal symptoms of maternal use of drugs of addiction Withdrawal symptoms from therapeutic use of drugs of newborn
D. **Neurocutaneous Syndromes**			
Congenital pigmentary anomalies of the skin	757.33	Q82.1 Q82.2	Xeroderma pigmentosum Mastocytosis
Neurofibromatosis	237.70	Q85.00	Neurofibromatosis, unspecified

Criteria Description	ICD 9	ICD 10	Criteria Description
Other Neurofibromatosis	237.79	Q85.09	Other neurofibromatosis
Sturge-Weber syndrome	759.6	Q85.8	Other phakomatoses, not elsewhere classified
Tuberous sclerosis	759.5	Q85.1	Tuberous sclerosis
D. Inborn Error of Metabolism			
Disorders of amino-acid transport and metabolism	270.0	E72.00 E72.01 E72.04 E72.09	Disorders of amino-acid transport, unspecified Cystinuria Cystinosis Other disorders of amino-acid transport
Phenylketonuria (PKU)	270.1	E70.0	Classical phenylketonuria
Other Disturbances of aromatic amino-acid metabolism	270.2	E70.21 E70.29 E70.30 E70.5 E70.8	Tyrosinemia Other disorders of tyrosine metabolism Albinism, unspecified Disorders of tryptophan metabolism Other disorders of aromatic amino-acid metabolism
Maple Sugar Urine Disease	270.3	E71.0 E71.120 E71.19 E71.2	Maple-syrup-urine disease Methylmalonic acidemia Other disorders of branched-chain amino-acid metabolism Disorder of branched-chain amino-acid metabolism, unspecified
Disturbances of Sulpher-bearing amino acid metabolism	270.4	E72.10	Disorders of sulphur-bearing amino-acid metabolism Homocystinuria Other disorders of sulphur-bearing amino-acid metabolism
Disorder of Urea cycle metabolism	270.6	E72.20 E72.22 E72.23 E72.29	Disorder of urea cycle metabolism, unspecified Arginosuccinic aciduria Citrullinemia Other disorders of urea cycle metabolism
Other disturbances of straight-chain amino-acid metabolism	270.7	E72.3 E72.8	Disorders of lysine and hydroxylysine metabolism Other specified disorders of amino-acid metabolism
Other specified disorders of amino-acid metabolism	270.8	E72.03 E72.8	Lowe's syndrome Other specified disorders of amino-acid metabolism
Unspecified disorder of amino acid metabolism	270.9	E72.9	Disorder of amino-acid metabolism, unspecified
General Category Disorders of Carbohydrate Metabolism Glycogenosis	271.0	E74.00 E74.01 E74.04 E74.09	Glycogen storage disease, unspecified von Gierke disease McArdle disease Other glycogen storage disease
Galactosemia	271.1	E74.21	Galactosemia
General Category Disorders of Lipid Metabolism	272.0	E78.0	Pure hypercholesterolemia
Lipidoses Fabry's disease - Gaucher's disease - Niemann Pick - sphingolipidoses	272.7	 E75.21 E75.22 E75.249 E77.0 E77.1	 Fabry (or Anderson-Fabry) disease Gaucher disease Niemann-Pick disease, unspecified Defects in post-translational modification of lysosomal enzymes Defects in glycoprotein degradation
Other disorders of lipid metabolism	272.8	E78.81 E78.89 E88.89	Lipoid dermatoarthritis Other lipoprotein metabolism disorders Other specified metabolic disorders

Criteria Description	ICD 9	ICD 10	Criteria Description
Mucopolysaccaridoses	277.5	E76.01 E76.03 E76.1 E76.219 E76.22 E76.29 E76.3	Hurler's syndrome Scheie's syndrome Mucopolysaccharidosis, type II-Hunter's syndrome Morquio mucopolysaccharidoses, unspecified Sanfilippo mucopolysaccharidoses Other mucopolysaccharidoses Mucopolysaccharidosis, unspecified
E. General Category: Cerebral degenerations of the central nervous system—usually manifested in childhood	330.__		
Leukodystrophy	330.0	E75.23 E75.25 E75.29	Krabbe disease Metachromatic leukodystrophy Other sphingolipidosis
Cerebral lipidoses such as TaySach's	330.1	E75.02 E75.19 E75.4	Tay-Sachs disease Other gangliosidosis Neuronal ceroid lipofuscinosis
Cerebral degeneration in generalized lipidoses	330.2	G93.89	Other specified disorders of brain
Cerebral Degenerations of childhood in other diseases	330.3	G93.9	Disorder of brain, unspecified
Other specified degenerations in childhood	330.8	F84.2 G31.81 G31.82	Rett's syndrome Alpers disease Leigh's disease
Unspecified cerebral degenerations in childhood	330.9	G94	Other disorders of brain in diseases classified elsewhere
F. Prenatal Infections			
TORCH" infections, including: Congenital rubella	771.0	P35.0	Congenital rubella syndrome
Congenital cytomegalovirus infection (CMV)	771.1	P35.1	Congenital cytomegalovirus infection
Congenital herpes simplex Congenital toxoplasmosis	771.2	P35.2 P37.1 P37.2 P37.8	Congenital herpesviral [herpes simplex] infection Congenital toxoplasmosis Neonatal (disseminated) listeriosis Other specified congenital infectious and parasitic diseases
G. Other Syndromes			
Cerebral gigantism	253.0	E22.0	Acromegaly and pituitary gigantism
General Category: Other and unspecified congenital anomalies	759.__		
Prader-willi syndrome Marfan syndrome Fragile x syndrome	759.81 759.82 759.83	Q87.1 Q87.40 Q99.2	Congenital malformation syndromes predominantly associated with short stature Marfan's syndrome, unspecified Fragile X chromosome
Other specified chromosome abnormalities	759.89	Q99.8 E78.71 E78.72 Q87.1 Q87.2 Q87.3 Q87.5 Q87.81	Other specified chromosome abnormalities Barth syndrome Smith-Lemli-Opitz syndrome Cornelia de Lange Congenital malformation syndromes predominantly involving limbs Congenital malformation syndromes involving early overgrowth—Beckwith Wiedemann Other congenital malformation syndromes with other skeletal changes Alport syndrome

Criteria Description	ICD 9	ICD 10	Criteria Description
		Q87.89 Q89.8	Other specified congenital malformation syndromes, not elsewhere classified Other specified congenital malformations
General Category: Congenital anomaly, unspecified	759.9	Q89.9	Congenital anomaly, unspecified
H. Sensory Impairment - Vision			
Vision--Impairment can be congenital or acquired (369—general category—more specific diagnosis obtained from physician) Profound impairment, both eyes (369.0-)	369.00	H54.0	Blindness, both eyes
Moderate or severe impairment, better eye, profound impairment lesser eye Blindness one eye; low vision other eye	369.01- 369.18	H54.10	Blindness, one eye, low vision other eye, unspecified eyes
Moderate or severe impairment, both eyes Low vision both eyes not otherwise specified	369.2- 369.20	H54.2	Low vision, both eyes
Better eye: severe vision impairment; lesser eye; impairment not further specified	369.21- 359.24	H54.10	Blindness, one eye, low vision other eye, unspecified eyes
Better eye: moderate vision impairment; lesser eye: moderate vision impairment	369.25	H54.2	Low vision, both eyes
Unqualified vision loss, both eyes	369.3	H54.3	Unqualified vision loss, both eyes
Legal blindness, as defined in USA	369.4	H54.8	Legal blindness, as defined in USA
Retrolental fibroplasia or retinopathy of prematurity ROP Stage 4	362.26	H35.159	Retinopathy of prematurity, stage 4, unspecified eye
ROP State 5	362.27	H35.169	Retinopathy of prematurity, stage 5, unspecified eye
Bilateral retrolental fibroplasia	362.21	H35.179	Retrolental fibroplasia, unspecified eye
Cortical Blindness	377.75	H47.619	Cortical blindness, unspecified side of brain
I.Sensory Impairment - Hearing-- Hearing impairment (25dB loss or greater) unilateral or bilateral General Category	389		
Conductive hearing loss, unspecified—includes: Conductive hearing loss external ear Conductive hearing loss tympanic membrane Conductive hearing loss middle ear Conductive hearing loss inner ear Conductive hearing loss, unilateral Conductive hearing loss, bilateral Conductive hearing loss of combined types	389.00 389.01 389.02 389.03 389.04 389.05 389.06 389.08	H90.2 H90.11 H90.12 H90.0 H90.2	Conductive hearing loss, unspecified (Conductive hearing loss, unilateral, right ear with unrestricted hearing on contralateral side Conductive hearing loss, unilateral, left ear with unrestricted hearing on contralateral side Conductive hearing loss, bilateral Conductive hearing loss, unspecified
Sensorineural hearing loss	389.10	H90.5	Unspecified sensorineural hearing loss
Sensory Hearing loss, bilateral Neural Hearing loss, bilateral Sensorineural Hearing loss, bilateral Sensorineural Hearing loss, left or right	389.11 389.12 389.18 389.15	H903 H90.41 H90.42	Sensorineural Hearing loss, bilateral Sensorineural Hearing loss, right ear Sensorineural Hearing loss, left ear
Mixed conductive and sensorineural hearing loss	389.20	H90.8	Mixed conductive and sensorineural hearing loss, unspecified

Criteria Description	ICD 9	ICD 10	Criteria Description
Hearing loss unspecified	389.9	H91.90	Unspecified hearing loss, unspecified ear
Central hearing loss	389.14	H90.5	Unspecified sensorineural hearing loss
J. **Orthopedic and Neurological Disorders**			
Anoxic brain damage	348.1	G93.1	Anoxic brain damage, not elsewhere classified
Anterior horn cell disease Werdnig-Hoffmann disease	335.— 335.0	G12.0	Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]
Spinal muscular atrophy unspecified	335.10	G12.9	Spinal muscular atrophy, unspecified
Kugelberg-welander disease	335.11	G12.1	Other inherited spinal muscular atrophy
Other spinal muscular atrophy	335.19	G12.8	Other spinal muscular atrophies and related syndromes
Amyotrophic lateral sclerosis	335.20	G12.21	Amyotrophic lateral sclerosis
Progressive muscular atrophy	335.21	G12.21	Amyotrophic lateral sclerosis
Progressive bulbar palsy	335.22	G12.22	Progressive bulbar palsy
Pseudobulbar palsy	335.23	G12.8	Other spinal muscular atrophies and related syndromes
Primary lateral sclerosis	335.24	G12.29	Other motor neuron disease
Other motor neuron diseases	335.29	G12.29	Other motor neuron disease
Other anterior horn cell diseases	335.8	G12.8	Other spinal muscular atrophies and related syndromes
Anterior horn cell disease unspecified	335.9	G12.9	Spinal muscular atrophy, unspecified
General Category: other specified muscle disorders			
Arthrogryposis	728.3	M62.3 M62.89	Immobility syndrome (paraplegic) Other specified disorders of muscle
Arthrogryposis multiplex, congenita	754.89	Q74.3	Arthrogryposis multiplex, congenita
Injury to the Brachial plexus—birth trauma	767.6	P14.0 P14.1 P14.3	Erb's paralysis due to birth injury Klumpke's paralysis due to birth injury Other brachial plexus birth injuries
Brachial plexus—post perinatal origin	953.4	S14.3XXA	Injury of brachial plexus, initial encounter
Cerebral cysts	348.0	G93.0	Cerebral cysts
Cerebral palsy (all types)- General Category	343.__		
Congenital diplegia	343.0	G80.1	Spastic diplegic cerebral palsy
Congenital Hemiplegia	343.1	G80.2	Spastic hemiplegic cerebral palsy
Congenital Quadriplegia	343.2	G80.0	Spastic quadriplegic cerebral palsy
Congenital Monoplegia	343.3	G80.8	Other cerebral palsy
Infantile hemiplegia	343.4	G80.2	Spastic hemiplegic cerebral palsy
Other specified infantile cerebral palsy	343.8	G80.8	Other cerebral palsy
Infantile cerebral palsy unspecified	343.9	G80.9	Cerebral palsy, unspecified
Cleft hand	755.58	Q71.60	Lobster-claw hand, unspecified hand
Congenital anomalies of the central nervous system – General Category	742.--		
Encephalocele	742.0	Q01.9	Encephalocele, unspecified
Microcephaly	742.1	Q02	Microcephaly
Congenital reduction deformities of brain	742.2	Q04.1 Q04.2 Q04.3	Arhinencephaly Holoprosencephaly Other reduction deformities of brain
Congenital hydrocephaly	742.3	Q03.0 Q03.1 Q03.8 Q03.9	Malformations of aqueduct of Sylvius Atresia of foramina of Magendie and Luschka Other congenital hydrocephalus Congenital hydrocephalus, unspecified
Other specified congenital anomalies of brain	742.4	Q04.5 Q04.6 Q04.8	Megalencephaly Congenital cerebral cysts Other specified congenital malformations of

Criteria Description	ICD 9	ICD 10	Criteria Description
			brain
Other specified congenital anomalies of spinal cord—general category	742.5		
Diastematomyelia	742.51	Q06.2	Diastematomyelia
Hydromyelia	742.53	Q06.4	Hydromyelia
Other specified congenital anomalies of spinal cord	742.59	Q06.0 Q06.1 Q06.3 Q06.8	Amyelia Hypoplasia and dysplasia of spinal cord Other congenital cauda equina malformations Other specified congenital malformations of spinal cord
Other specified congenital anomalies of nervous system	742.8	G90.1 Q07.8	Familial dysautonomia [Riley-Day] Other specified congenital malformations of nervous system
Unspecified congenital anomaly of brain spinal cord and nervous system	742.9	Q07.9	Congenital malformation of nervous system, unspecified
Other congenital musculoskeletal anomalies - General Category	755.__		
Reduction of deformities of upper limb Transverse deficiency of upper limb Longitudinal deficiency of upper limb	755.20 755.21- 755.22	Q71.899 Q71.90 Q71.00	Other reduction defects of unspecified upper limb Unspecified reduction defect of unspecified upper limb Congenital complete absence of unspecified upper limb
Longitudinal deficiency combined involving humerus	755.23- 755.24	Q71.10	Congenital absence of unspecified upper arm and forearm with hand present.
Longitudinal deficiency , radioulner, complete or partial	755.25	Q71.20	Congenital absence of both forearm and hand, unspecified upper limb.
Longitudinal deficiency radial, complete or partial	755.26	Q71.40	Longitudinal reduction defect of snspecified ratius
Longitudinal deficiency, radial, complete or partial	755.27	Q71.50	Longitudinal reduction defect of unspecified ulna
Longitudinal deficiency ulnar, carpals or metacarpals, phalanges, finger	755.28- 755.29	Q71.30	Congenital absence of unspecified hand/finger
Reduction of deformities of lower limbs Longitudinal deficiency of lower limb, not classified elsewhere	755.30 755.32	Q72.899	Other reduction defects of unspecified lower limb
Transverse deficiency of lower lime	755.31	Q72.00	Congenital complete absence of unspecified lower limb
Longitudinal deficiency combined involving tibia and fibula	755.33	Q72.10	Congenital absence of unspecified thigh and lower leg with foot present
Longitudinal deficiency femoral, complete/incomplete	755.34	Q72.40	Longitudinal reduction defect of unspecified femur
Longitudinal deficiency tibiofibular complete or partial	755.35	Q72.20	Congenital absence of both lower leg and foot, unspecified lower limb
Longitudinal deficiency, tibia, complete/partial	755.36	Q72.50	Longitudinal reduction defect of unspecified tibia
Longitudinal deficiency, fibular, complete/partial	755.37	Q72.60	
Longitudinal deficiency, tarsals or metatarsals complete/partial Longitudinal deficiency, phalanges, complete/partial	755.38 755.39	Q72.30 Q72.70	Congenital absence of unspecified foot and toes Split foot, unspecified lower limb
Reduction deformities, unspecified limb	755.4	Q73.0 Q73.1 Q73.8	Congenital absence of unspecified limb(s) Phocomelia, unspecified limb(s) Other reduction defects of unspecified limb(s)
Congenital cleft hand	755.58	Q71.60	Lobster-claw hand, unspecified hand

Criteria Description	ICD 9	ICD 10	Criteria Description
Anomalies of skull and face bone Premature closure of cranial sutures	756.0	Q75.0 Q75.2 Q75.9	Craniosynostosis Hypertelorism Congenital malformation of skull and face bones, unspecified
Absence of vertebra, congenital	756.13	Q76.49	Other congenital malformations of spine, not associated with scoliosis
Chondrodystrophies	756.4	Q77.1 Q77.4 Q77.8 Q78.4	Thanatophoric short stature Achondroplasia Other osteochondrodysplasia with defects of growth of tubular bones and spine Enchondromatosis
Osteodystrophies, unspecified	756.50	Q78.9	Osteochondrodysplasia, unspecified
Osteogenesis imperfecta	756.51	Q78.0	Osteogenesis imperfecta
Other symbolic dystunction-general category Developmental apraxia of speech	784.6 784.69	 R48.2 R48.8	 Apraxia Other symbolic dysfunctions
Encephalopathy Not Otherwise Specified	348.30	G93.40	Encephalopathy, unspecified
Hypoxic Ischemic Encephalopathy	768.70 768.73	P91.60 P91.63	Hypoxic Ischemic Encephalopathy Hypoxic Ischemic Encephalopathy, severe/Grade 3
Fracture of vertebral column with spinal cord injury (806) General Category—include additional diagnosis from physician	806.00	S12.000A S12.001A S12.100A S12.101A S12.200A S12.201A S12.300A S12.301A S14.101A S14.102A S14.103A S14.104A	Unspecified displaced fracture of first cervical vertebra, initial encounter for closed fracture Unspecified nondisplaced fracture of first cervical vertebra, initial encounter for closed fracture Unspecified displaced fracture of second cervical vertebra, initial encounter for closed fracture Unspecified nondisplaced fracture of second cervical vertebra, initial encounter for closed fracture Unspecified displaced fracture of third cervical vertebra, initial encounter for closed fracture Unspecified nondisplaced fracture of third cervical vertebra, initial encounter for closed fracture Unspecified displaced fracture of fourth cervical vertebra, initial encounter for closed fracture Unspecified nondisplaced fracture of fourth cervical vertebra, initial encounter for closed fracture Unspecified injury at C1 level of cervical spinal cord, initial encounter Unspecified injury at C2 level of cervical spinal cord, initial encounter Unspecified injury at C3 level of cervical spinal cord, initial encounter Unspecified injury at C4 level of cervical spinal cord, initial encounter
General Category: Hemiplegia and hemiparesis	342.--		
Flaccid hemiplegia	342.00	G81.00	Flaccid hemiplegia affecting unspecified side
Flaccid hemiplegia and hemiparesis affecting dominant side	342.01	G81.01 G81.02	Flaccid hemiplegia affecting right dominant side Flaccid hemiplegia affecting left dominant side
Flaccid hemiplegia and hemiparesis affecting nondominant side	342.02	G81.03 G81.04	Flaccid hemiplegia affecting right nondominant side Flaccid hemiplegia affecting left

Criteria Description	ICD 9	ICD 10	Criteria Description
			nondominant side
Spastic hemiplegia	342.10	G81.10	Spastic hemiplegia affecting unspecified side
Spastic hemiplegia and hemiparesis affecting dominant side	342.11	G81.11 G81.12	Spastic hemiplegia and hemiparesis affecting right dominant side Spastic hemiplegia and hemiparesis affecting left dominant side
Spastic hemiplegia and hemiparesis affecting nondominant side	342.10	G81.10	Spastic hemiplegia affecting unspecified side
Other specified hemiplegia	342.80	G81.90	Hemiplegia, unspecified affecting unspecified side
Hemiplegia, unspecified	342.90	G81.90	Hemiplegia, unspecified affecting unspecified side
General Category: Hereditary\degenerative diseases of the central nervous system	331.__		
Communicating hydrocephalus	331.3	G91.0	Communicating hydrocephalus
Obstructive hydrocephalus	331.4	G91.1	Obstructive hydrocephalus
Cerebral degeneration in diseases classified elsewhere	331.7	G94	Other disorders of brain in diseases classified elsewhere
Werdnig-Hoffman disease	335.0	G12.0	Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]
Other Cerebral Degeneration	331.8	G31.89	Other specified degenerative diseases of the nervous system
Infantile spasms with intractable epilepsy-not including febrile seizures (R56.00, R56.01)	345.60	G40.401 G40.409	Other generalized epilepsy and epileptic syndromes, not intractable, with status epilepticus Other generalized epilepsy and epileptic syndromes, not intractable, without status epilepticus
Infantile spasms with intractable epilepsy	345.61	G40.411 G40.419	Other generalized epilepsy and epileptic syndromes, intractable, with status epilepticus Other generalized epilepsy and epileptic syndromes, intractable, without status epilepticus
Intraventricular hemorrhage (IVH) – Grade 3 Grade 4	772.13 772.14	P52.21 P52.22	Intraventricular (nontraumatic) hemorrhage, grade 3, of newborn Intraventricular (nontraumatic) hemorrhage, grade 4, of newborn
General Category--Spina Bifida/Neural Tube Defect	741.00	Q05.9 Q05.4 Q07.01 Q07.02 Q07.03	Spina bifida, unspecified Unspecified spina bifida with hydrocephalus Arnold-Chiari syndrome with spina bifida Arnold-Chiari syndrome with hydrocephalus Arnold-Chiari syndrome with spina bifida and hydrocephalus
Spina Bifida, Cervical region with Hydrocephalus	741.01	Q05.0	Cervical Spina Bifida with hydrocephalus
Spina Bifida, dorsal (thoracic) region with hydrocephalus	741.2	Q05.1	Thoracic Spina Bifida with hydrocephalus
Spina Bifida, lumbar region with hydrocephalus	741.03	Q05.2 Q05.3	Lumbar spina bilfida with hydrocephalus Sacral spina bifida with hydrocephalus
Meningomyelocele	741.90	Q05.8	Sacral spina bifida without hydrocephalus
Myelomeningocele Spina Bifida with hydrocephalus	741.90 741.91 741.92 741.93	Q05.8 Q05.5 Q05.6 Q05.7	Sacral spina bifida without hydrocephalus Cervical spina bifida without hydrocephalus Thoracic spina bifida without hydrocephalus Lumbar spina bifida without hydrocephalus

Criteria Description	ICD 9	ICD 10	Criteria Description
General Category: Congenital hereditary muscular dystrophy	359.__		
Muscular dystrophies and other myopathies	359.0	G71.2	Congenital myopathies
Hereditary progressive muscular dystrophy	359.1	G71.0	Muscular dystrophy
Myotonic muscular dystrophy	359.21	G71.11	Myotonic muscular dystrophy
Myotonia, congenita	359.22	G71.12	Myotonia congenita
Myontic chondrodystrophy	359.23	G71.13	Myotonic chondrodystrophy
General Category—other paralytic syndromes	344.__		
Quadriplegia and quadriparesis -- unspecified	344.00	G82.50	Quadriplegia, unspecified
	344.01	G82.51	Quadriplegia, C1-C4 complete
Quadriplegia c1-c4 complete	344.02	G82.52	Quadriplegia C1-C4 incomplete
Quadriplegia c1-c4 incomplete	344.03	G82.53	Quadriplegia, C5-C7 complete
Quadriplegia c5-c7 complete	344.04	G82.54	Quadriplegia, C5-C7 incomplete
Quadriplegia c5-c7 incomplete	344.09	G82.50	Quadriplegia, unspecified
Other quadriplegia	3441	G82.20	Paraplegia, unspecified
Paraplegia	344.2	G83.0	Diplegia of upper limbs
Diplegia of upper limbs	344.30	G83.10	Monoplegia of lower limb affecting unspecified side
Monoplegia of lower limb affecting unspecified side	344.31	G83.11	Monoplegia of lower limb affecting right dominant side
Monoplegia of lower limb affecting dominant side		G83.12	Monoplegia of lower limb affecting left dominant side
	344.32	G83.13	Monoplegia of lower limb affecting right nondominant side
Monoplegia of lower limb affecting nondominant side		G83.14	Monoplegia of lower limb affecting left nondominant side
	344.40	G83.20	Monoplegia of upper limb affecting unspecified side
Monoplegia of upper limb affecting unspecified side	344.41	G83.21	Monoplegia of upper limb affecting right dominant side
Monoplegia of upper limb affecting dominant side		G83.22	Monoplegia of upper limb affecting left dominant side
	344.42	G83.23	Monoplegia of upper limb affecting right nondominant side
Monoplegia of upper limb affecting nondominant side		G83.24	Monoplegia of upper limb affecting left nondominant side
	344.5	G83.30	Monoplegia, unspecified affecting unspecified side
Unspecified monoplegia			
	344.60	G83.4	Cauda equina syndrome
Cauda equina syndrome without neurogenic bladder	344.61	G83.4	Cauda equina syndrome
Cauda equina syndrome with neurogenic bladder	344.81	G83.5	Locked-in state
Locked-in state	344.89	G83.81	Brown-Sequard syndrome
Other specified paralytic		G83.84	Todd's paralysis (postepileptic)
Syndrome		G83.89	Other specified paralytic syndromes
	344.9	G83.9	Paralytic syndrome, unspecified
Paralysis unspecified			
Paraplegia	344.1	G82.20	Paraplegia, unspecified
Diplegia of upper limbs	344.2	G83.0	Diplegia of upper limbs
Monoplegia of lower limb	344.30	G83.10	Monoplegia of lower limb affecting unspecified side
Monoplegia of upper limb	344.40	G83.20	Monoplegia of upper limb affecting unspecified side

Criteria Description	ICD 9	ICD 10	Criteria Description
Unspecified monoplegia	344.5	G83.30	Monoplegia, unspecified affecting unspecified side
General Category— Spinal cord injury without evidence of spinal bone injury	952		
Cervical, Dorsal Range of Codes based on location of injury	952.0- 952.1	S14.101A- S34.139A	Unspecified injury at C1 level of cervical spinal cord, initial encounter
Unspecified site of spinal cord injury without spinal bone injury	952.9	S14.109A S24.109A S34.109A S34.139A	Unspecified injury at unspecified level of cervical spinal cord, initial encounter Unspecified injury at unspecified level of thoracic spinal cord, initial encounter Unspecified injury at unspecified level of lumbar spinal cord, initial encounter Unspecified injury at unspecified level of sacral spinal cord, initial encounter
General Category--Occlusion of cerebral arteries or stroke	434		
cerebral thrombosis without cerebral infarction	434.00	I66.09 I66.19 I66.29	Occlusion and stenosis of unspecified middle cerebral artery Occlusion and stenosis of unspecified anterior cerebral artery Occlusion and stenosis of unspecified posterior cerebral artery
cerebral embolism with cerebral infarction	434.01	I63.40	Cerebral infarction due to embolism of unspecified cerebral artery
Cerebral embolism without cerebral infarction	434.10	I66.09 I66.19 I66.29 I66.9	Occlusion and stenosis of unspecified middle cerebral artery Occlusion and stenosis of unspecified anterior cerebral artery Occlusion and stenosis of unspecified posterior cerebral artery Occlusion and stenosis of unspecified cerebral artery
Cerebral artery occlusion unspecified without cerebral infarction	434.90	I66.9	Occlusion and stenosis of unspecified cerebral artery
General category--Cerebral laceration and contusion or traumatic brain injury Includes range of codes for intracranial injury	851.00- 854.00	S06.330A- S06.339A	Contusion and laceration of cerebrum, unspecified, without loss of consciousness, initial encounter —range of codes for specific diagnoses
Shaken Infant Syndrome	995.55	T74.4XXA	Shaken Infant Syndrome
K. Social Emotional Disorders			
Social Emotional Disorders Childhood Depressive disorders, not elsewhere classified	311	F32.9	Major depressive disorder, single episode, unspecified
Reactive attachment disorder	313.89	F93.8 F94.1 F98.8	Other childhood emotional disorders Reactive attachment disorder of childhood Other specified behavioral and emotional disorders with onset usually occurring in childhood and adolescence
L. Pervasive Developmental Disorders			

Criteria Description	ICD 9	ICD 10	Criteria Description
Pervasive Developmental Disorders General Category including: Autistic disorder current or active state	299.00	F84.0	Autistic disorder
Autistic disorder residual state	299.01	F84.0	Autistic disorder
Childhood disintegrative disorder current or active state	299.10	F84.3	Other childhood disintegrative disorder
Childhood disintegrative disorder residual state	299.11	F84.3	Other childhood disintegrative disorder
Other specified pervasive developmental disorders current or active state	299.80	F84.5 F84.8	Asperger's syndrome Other pervasive developmental disorders
Other specified pervasive developmental disorders residual state	299.81	F84.5 F84.8	Asperger's syndrome Other pervasive developmental disorders
Unspecified pervasive developmental disorder current or active state	299.90	F84.9	Pervasive developmental disorder, unspecified
Unspecified pervasive developmental disorder residual state	299.91	F84.9	Pervasive developmental disorder, unspecified
Asperger syndrome / disorder	299.80	F84.5 F84.8	Asperger's syndrome Other pervasive developmental disorders
M. Medically Related Disorders			
Congenital or infancy-onset hypothyroidism	243	E00.9	Congenital iodine-deficiency syndrome, unspecified
Cleft Palate- eligibility may continue post-operative repair	V136.4	Z87.730	Personal history (corrected) cleft lip and palate
Cleft palate —unspecified	749.00	Q35.9	Cleft palate, unspecified
unilateral, complete	749.01	Q35.9	Cleft palate, unspecified
unilateral, incomplete	749.02	Q35.7 Q35.9	Cleft uvula Cleft palate, unspecified
bilateral, complete	749.13	Q36.0	Cleft lip, bilateral
bilateral, incomplete	749.14	Q36.0	Cleft lip, bilateral
Cleft palate with cleft lip —unspecified	749.20	Q37.9	Unspecified cleft palate with unilateral cleft lip
unilateral, complete	749.21	Q37.9	Unspecified cleft palate with unilateral cleft lip
unilateral, incomplete	749.22	Q37.9	Unspecified cleft palate with unilateral cleft lip
bilateral, complete	749.23	Q37.8	Unspecified cleft palate with bilateral cleft lip
bilateral, incomplete	749.24	Q37.8	Unspecified cleft palate with bilateral cleft lip
Toxic effects of lead and its compounds (including fumes) General category	984.0	T56.0X1A T56.0X2A T56.0X3A T56.0X4A	Toxic effect of lead and its compounds, accidental (unintentional), initial encounter Toxic effect of lead and its compounds, intentional self-harm, initial encounter Toxic effect of lead and its compounds, assault, initial encounter Toxic effect of lead and its compounds, undetermined, initial encounter
unspecified lead compound effects	984.9	M1A.10X1 T56.0X1A T56.0X2A T56.0X3A T56.0X4A	Lead-induced chronic gout, unspecified site, with tophus (tophi) Toxic effect of lead and its compounds, accidental (unintentional), initial encounter Toxic effect of lead and its compounds, intentional self-harm, initial encounter Toxic effect of lead and its compounds, assault, initial encounter Toxic effect of lead and its compounds, undetermined, initial encounter
Non-organic failure to thrive	783.41	R62.51	Failure to thrive (child)
Chronic respiratory failure or ventilator dependence	518.83	J96.10	Chronic respiratory failure, unspecified whether with hypoxia or hypercapnia

Criteria Description	ICD 9	ICD 10	Criteria Description
N. Prematurity			
Bronchopulmonary Dysplasia (BPD)	770.7	P27.0 P27.1 P27.8	Wilson-Mikity syndrome Bronchopulmonary dysplasia originating in the perinatal period
Disorders relating to short gestation and low birth weight General Category—include 5 digit code	765.00	P07.00 P07.10	Extremely low birth weight newborn, unspecified weight Other low birth weight newborn, unspecified weight
Other preterm infant's birth weight of 1000-2499 grams —unspecified weight—an EarlySteps eligibility criterion is <1500 grams at birth:	765.10	P07.00 P07.10	Extremely low birth weight newborn, unspecified weight Other low birth weight newborn, unspecified weight
less than 500 grams	765.11	P07.01	Extremely low birth weight newborn, less than 500 grams
500 to 749 grams	765.12	P07.02	Extremely low birth weight newborn, 500-749 g
750 to 999 grams	765.13	P07.03	Extremely low birth weight newborn, 750-999 grams
1000 to 1249 grams	765.14	P07.14	Other low birth weight newborn, 1000-1249 grams
1250 to 1499 grams	765.15	P07.15	Other low birth weight newborn, 1250-1499 grams
Please refer to weight conversion table at the end of the chapter. --Weeks of gestation –unspecified gestation--General Category—	765.20	P07.20 P07.30	Extreme immaturity of newborn, unspecified weeks of gestation Preterm newborn, unspecified weeks of gestation
EarlySteps eligibility criteria is 32 weeks gestation or less Less than 24 weeks of gestation	765.21	P07.21 P07.22	Extreme immaturity of newborn, gestational age less than 23 completed weeks Extreme immaturity of newborn, gestational age 23 completed weeks
24 complete weeks of gestation	765.22	P07.23	Extreme immaturity of newborn, gestational age 24 completed weeks
25-26 weeks of gestation	765.23	P07.24 P07.25	Extreme immaturity of newborn, gestational age 25 completed weeks Extreme immaturity of newborn, gestational age 26 completed weeks
27-28 weeks of gestation	765.24	P07.26 P07.31	Extreme immaturity of newborn, gestational age 27 completed weeks Preterm newborn, gestational age 28 completed weeks
29-30 weeks of gestation	765.25	P07.32 P07.33	Preterm newborn, gestational age 29 completed weeks Preterm newborn, gestational age 30 completed weeks
31-32 weeks of gestation	765.26	P07.34 P07.35	Preterm newborn, gestational age 31 completed weeks Preterm newborn, gestational age 32 completed weeks

Additional information about Prematurity eligibility criteria, please see page 10.