

DIAGNOSIS	NEW ICD-10 CODES	OLD ICD-9 CODES	DIAGNOSIS	NEW ICD-10 CODES	OLD ICD-9 CODES
Agenesis Corpus Callosum	Q04.3	742.2	Failure to thrive	R62.51	783.41
Anoxic Brain Damage/Cerebral Hypoxia	G93.1	348.1	Fetal Alcohol Syndrome	Q86.0	760.71
Arthrogryposis	Q74.3	728.3	Fragile X Syndrome	Q99.2	759.83
Asthma	J45.998	493.90	Gastrostomy- Tube Feed	293.1	V44.1
Attention Deficit Disorder (ADD)	F90.9	314.00	Hydrocephalus	Q03.8	742.3
Attention Deficit Disorder (ADHD)	F90.2	314.01	Lack of expected normal physiological development in childhood	R62.50	783.4
Autism	F84.0	299.00	Manic Depression	F31.9	296.1
Bipolar Depression	F31.30	296.50	Meningitis	G03.9	322.9
Cerebral Palsy (athetoid)	G80.3	333.7	Mentally Handicapped (moderate)	F71	318.0
Cerebral Palsy (quadriplegic)	G80.0	343.2	Mentally handicapped (profoundly) (PMH)	F73	318.2
Cerebral Palsy (spastic)	G80.9	343.9	Mentally handicapped (severe)	F72	318.1
Charcot-Marie-Tooth	G60.0	356.1	Microcephaly	Q02	742.1
Colostomy	243.3	V55.3	Myotonic Dystrophy	G71.2	359.0
Congenital Cytomegalic Inclusion Disease (CMV)	P35.1	771.1	Myelodysplasia Spinal Abnormality	Q06.8	742.59
Congenital Hydrocephalus	Q03.8	742.3	Obsessive Compulsive Disorder	F42	300.3
Congenital Nystagmus	H55.01	379.51	Oppositional Defiant Disorder	F91.3	313.81
Convulsions Febrile	R56.00	780.31	Paralysis (complete) (incompleat)	G83.9	344.9
Convulsions Seizures NOS	R56.9	780.39	Physical Retardation	R62.50	783.4
Contracture of Muscle	M62.40	728.85	Prader Willi Syndrome	Q87.1	759.81
Cystic Fibrosis	E84.9	277.00	Renal Disease	I12.9	403.9
Deaf (HI)	491.90	389.9	Tourettes Syndrome	F95.2	307.23
Delayed milestones (late talker, late talker)	R62.0	783.43	Severely Emotionally Disabled (SED)	F60.30	301.3
Developmentally Delayed (DD) physiological, unspecified	R62.50	783.40	Short stature (growth failure, growth retardation)	R62.52	783.43
Developmentally Delayed Unspecified mental	F89	315.9	Specific Learning Disability (SLD)	F81.89	315.2
Diabetes Type I	E10.9	250.01	Speech/ Language Impaired (S/L) dysphasia	R47.89	784.5
Diabetes Type II	E11.9		Spina Bifida	Q05.8	741.90
Down Syndrome	Q90.9	758.0	Spinal Muscular Atrophy	G12.8	335.1
Drowning	T25.1XXA	994.1	Sturge Weber	Q85.8	759.6
Duchenne Muscular Dystrophy	G71.0	359.1	Tar Syndrome (Radial Aplasia)	Q87.2	755.26
Educable Mentally Handicapped (EMH)	F70	317	Tramatic Brain Injury	509.8XXA	310.9
Emotionally Handicapped (EH)	F60.4	301.59	Tremor	R25.9	781.0
Encephalitis	G04.90	323.9	Unspecified Anomaly of Brian, Spinal Cord, Nervous	Q07.9	742.9
Epilepsy (Seizure Disorder) Convulsive	780.39	345.1	Unspecified Visual Loss Visually Impaired (VI)	H54.7	369.9
Epilepsy (Seizure Disorder) Non-Convulsive	G40.A01	345.00			