FREQUENTLY USED ICD10 CODES
NEUROLOGY

DEVELOPMENTAL				
Delay/ID	F70	Mild intellectual disabilities	R62.0	Delayed milestone in childhood
	F71	Moderate intellectual disabilities	F81.9	Developmental disorder of scholastic skills, unspecified
	F79	Unspecified intellectual disabilities	R62.51	Failure to thrive (child)
	G31.84	Mild cognitive impairment, so stated	F82	Specific developmental disorder of motor function
Disorders	F84.0	Autistic disorder	F88	Other disorders of psychological development
	F84.9	Pervasive developmental disorder, unspecified	R62.50	Unspecified lack of expected normal physiological development in childhood
	Q99.2	Fragile X syndrome	F89	Unspecified disorder of psychological development
	F90.1	Attention-deficit hyperactivity disorder, predominantly hyperactive type		
Speech & Language	F80.2	Mixed receptive-expressive language disorder		
	F80.9	Developmental disorder of speech and language, unspecified		
	F95.2	Tourette's disorder		
	R47.1	Dysarthria and anarthria		

CNS-BRAIN				
Size	Q02	Microcephaly	Q04.5	Megalencephaly
	Q75.3	Macrocephaly		
Leukodystrophy	R90.82	White matter disease, unspecified		
	E75.25	Metachromatic leukodystrophy		
	I67.850	Cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy		
	E75.29	Other sphingolipidosis		
Imaging	R90.89	Other abnormal findings on diagnostic imaging of central nervous system		
	R93.0	Abnormal findings on diagnostic imaging of skull and head, not elsewhere classified		
Hydrocephalus	Q03.9	Congenital hydrocephalus, unspecified		
	G91.9	Hydrocephalus, unspecified		
Brain malformations	Q28.3	Other malformations of cerebral vessels	Q04.2	Holoprosencephaly
	Q04.0	Congenital malformations of corpus callosum	Q04.9	Congenital malformation of brain, unspecified
	Q03.1	Atresia of foramina of Magendie and Luschka	Q04.8	Other specified congenital malformations of brain
	Q07.8	Other specified congenital malformations of the nervous system		
Other	G90.9	Disorder of the autonomic nervous system, unspecified	Q04.3	Other reduction deformities of brain
	I63.9	Cerebral infarction, unspecified	G93.89	Other specified disorders of the brain

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FREQUENTLY USED ICD10 CODES
NEUROLOGY

EPILEPSY				
Intractable	With Status Epilepticus		Without Status Epilepticus	
	G40.311	Generalized idiopathic epilepsy and epileptic syndromes, intractable, with status epilepticus	G40.319	Generalized idiopathic epilepsy and epileptic syndromes, intractable, without status epilepticus
	G40.411	Other generalized epilepsy and epileptic syndromes, intractable, with status epilepticus	G40.419	Other generalized epilepsy and epileptic syndromes, intractable, without status epilepticus
	G40.911	Epilepsy, unspecified, intractable, with status epilepticus	G40.919	Epilepsy, unspecified, intractable, without status epilepticus
	G40.823	Epileptic spasms, intractable, with status epilepticus	G40.824	Epileptic spasms, intractable, without status epilepticus
	G40.111	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, intractable, with status epilepticus	G40.119	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, intractable, without status epilepticus
	G40.211	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, intractable, with status epilepticus	G40.219	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, intractable, without status epilepticus
	G40.011	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, intractable, with status epilepticus	G40.019	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, intractable, without status epilepticus
	G40.813	Lennox-Gastaut syndrome, intractable, with status epilepticus	G40.814	Lennox-Gastaut syndrome, intractable, without status epilepticus
	G40.A11	Absence epileptic syndrome, intractable, with status epilepticus	G40.A19	Absence epileptic syndrome, intractable, without status epilepticus
	G40.B11	Juvenile myoclonic epilepsy, intractable, with status epilepticus	G40.B19	Juvenile myoclonic epilepsy, intractable, without status epilepticus
Not Intractable	With Status Epilepticus		Without Status Epilepticus	
	G40.301	Generlized idiopathic epilepsy and epileptic syndromes, not intractable, with status epilepticus	G40.309	Generalized idiopathic epilepsy and epileptic syndromes, not intractable, without status epilepticus
	G40.401	Other generalized epilepsy and epileptic syndromes, not intractable, with status epilepticus	G40.409	Other generalized epilepsy and epileptic syndromes, not intractable, without status epilepticus
	G40.901	Epilepsy, unspecified, not intractable, with status epilepticus	G40.909	Epilepsy, unspecified, not intractable, without status epilepticus
	G40.822	Epileptic spasms, not intractable, with status epilepticus	G40.821	Epileptic spasms, not intractable, without status epilepticus
	G40.101	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, not intractable, with status epilepticus	G40.109	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, not intractable, without status epilepticus
	G40.201	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, not intractable, with status epilepticus	G40.209	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, not intractable, without status epilepticus
	G40.001	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, not intractable, with status epilepticus	G40.009	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, not intractable, without status epilepticus
	G40.811	Lennox-Gastaut syndrome, not intractable, with status epilepticus	G40.812	Lennox-Gastaut syndrome, not intractable, without status epilepticus
	G40.A01	Absence epileptic syndrome, not intractable, with status epilepticus	G40.802	Other epilepsy, not intractable, without status epilepticus
	G40.B01	Juvenile myoclonic epilepsy, not intractable, with status epilepticus	G40.A09	Absence epileptic syndrome, not intractable, without status epilepticus
			G40.B09	Juvenile myoclonic epilepsy, not intractable, without status epilepticus
Other	P90	Convulsions of newborn	G93.49	Other encephalopathy
	R56.9	Unspecified convulsions	G40.89	Other seizures
	G93.40	Encephalopathy, unspecified		

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FREQUENTLY USED ICD10 CODES

NEUROLOGY

NEURODEGENERATIVE				
	G30.0	Alzheimer's disease with early onset	G31.09	Other frontotemporal dementia
	G30.9	Alzheimer's disease, unspecified	G20	Parkinson's disease
	G12.21	Amyotrophic lateral sclerosis	F03.91	Unspecified dementia with behavioral disturbance
	R47.01	Aphasia	F03.90	Unspecified dementia without behavioral disturbance
	G31.9	Degenerative disease of nervous system, unspecified		

MUSCULOSKELETAL				
Neuropathy	G60.0	Hereditary motor and sensory neuropathy	G60.9	Hereditary and idiopathic neuropathy, unspecified
	G60.3	Idiopathic progressive neuropathy	G61.81	Chronic inflammatory demyelinating polyneuritis
	G60.8	Other hereditary and idiopathic neuropathies	G62.9	Polyneuropathy, unspecified
Myopathy	G12.20	Motor neuron disease, unspecified		
	G12.0	Infantile spinal muscular atrophy, type 1		
	G71.01	Duchenne or Becker muscular dystrophy		
	G71.11	Myotonic muscular dystrophy		
	G71.2	Congenital myopathies		
	G71.3	Mitochondrial myopathy, not elsewhere classified (Leber's disease, Leigh's encephalopathy, Kearns-Sayre syndrome, mitochondrial metabolism disorders)		
	G71.09	Other specified muscular dystrophies (Limb-Girdle, Ocular, Oculopharyngeal, scapuloperoneal, congenital)		
	M62.89	Other specified disorders of muscle		
	G72.9	Myopathy, unspecified		
Other neuromuscular	R25.2	Cramp and spasm	M62.81	Muscle weakness (generalized)
	R26.89	Other abnormalities of gait and mobility	R94.131	Abnormal electromyogram (EMG)
	R26.9	Unspecified abnormalities of gait and mobility	P94.2	Congenital hypotonia
	R29.898	Other symptoms and signs involving the musculoskeletal system	G70.2	Congenital and developmental myasthenia
	R53.1	Weakness	G70.9	Myoneural disorder, unspecified
Sensory	M79.673	Pain in unspecified foot	M79.643	Pain in unspecified hand
	M79.641	Pain in right hand	R20.2	Paresthesia of skin
	M79.642	Pain in left hand	M79.671	Pain in right foot
	M79.606	Pain in leg, unspecified	M79.672	Pain in left foot
Skeletal	Q87.1	Congenital malformation syndromes predominately associated with short stature		
	Q66.7	Congenital pes cavus		
	R62.52	Short stature (child)		
	Q67.5	Congenital deformity of the spine (scolisis/kyphoscholsis)		
	M41.9	Scoliosis, unspecified		
	G95.9	Disease of spinal cord, unspecified		

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NEUROLOGY

MOVEMENT				
Cerebral Palsy	G80.8	Other cerebral palsy	G80.2	Spastic hemiplegic cerebral palsy
	G80.9	Cerebral palsy, unspecified	G82.50	Quadriplegia, unspecified
	G80.1	Spastic diplegic cerebral palsy	G82.20	Paraplegia, unspecified
	G80.0	Spastic quadriplegic cerebral palsy		
Ataxia	G11.1	Early-onset cerebellar ataxia	G11.9	Hereditary ataxia, unspecified
	G11.2	Late-onset cerebellar ataxia	G32.81	Cerebellar ataxia in diseases classified elsewhere
	G11.8	Other hereditary ataxias	R27.0	Ataxia, unspecified
Dystonia	G24.9	Dystonia, unspecified		
	G24.8	Other dystonia		
Other	R26.1	Paralytic gait	G83.9	Paralytic syndrome, unspecified
	G11.4	Hereditary spastic paraplegia	F98.4	Stereotyped movement disorders
	R25.1	Tremor, unspecified		

OTHER				
	R74.8	Abnormal levels of other serum enzymes	Q38.2	Macroglossia
	D17.71	Benign lipomatous neoplasm of kidney	Q89.7	Multiple congenital malformations, not elsewhere classified
	D21.9	Benign neoplasm of connective and other soft tissue, unspecified	P05.10	Newborn small for gestational age, unspecified weight
	Q18.9	Congenital malformation of face and neck, unspecified	E28.39	Other primary ovarian failure
	Q87.3	Congenital malformation syndromes involving early overgrowth	Q85.1	Tuberous sclerosis
	L81.9	Disorder of pigmentation, unspecified	Z13.71	Encounter for nonprocreative screening for genetic disease carrier status
	P08.0	Exceptionally large newborn baby	O28.3	Abnormal ultrasonic finding on antenatal screening of mother

FAMILY HISTORY		
	Z82.0	Family history of epilepsy and other diseases of the nervous system
	Z81.8	Family history of other mental and behavioral disorders
	Z84.81	Family history of carrier of genetic disease

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