CHAPTER VI

Diseases of the nervous system

(G00-G99)

certain conditions originating in the perinatal period

certain infectious and parasitic diseases

complications of pregnancy, childbirth and the puerperium

congenital malformations, deformations and chromosomal abnormalities

endocrine, nutritional and metabolic diseases

injury, poisoning and certain other consequences of external causes

neoplasms

symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified

G00-G09 Inflammatory diseases of the central nervous system

G10-G14 Systemic atrophies primarily affecting the central nervous system

G20-G26 Extrapyramidal and movement disorders

G30-G32 Other degenerative diseases of the nervous system

G35-G37 Demyelinating diseases of the central nervous system

G40-G47 Episodic and paroxysmal disorders

G50-G59 Nerve, nerve root and plexus disorders

G60-G64 Polyneuropathies and other disorders of the peripheral nervous system

G70-G73 Diseases of myoneural junction and muscle

G80-G83 Cerebral palsy and other paralytic syndromes

G90-G99 Other disorders of the nervous system

Inflammatory diseases of the central nervous system

G00-G09

G00 Bacterial meningitis, not elsewhere classified

arachnoiditisbacterial

leptomeningitisbacterial

meningitisbacterial

pachymeningitisbacterial

bacterial:meningoencephalitis

bacterial:meningomyelitis

G00.0 Haemophilus meningitis

Meningitis due to Haemophilus influenzae

G00.1 Pneumococcal meningitis

G00.2 Streptococcal meningitis

G00.3 Staphylococcal meningitis

G00.8 Other bacterial meningitis

Meningitis due to:Escherichia coli

Meningitis due to:Friedl?nder bacillus

Meningitis due to:Klebsiella

G00.9 Bacterial meningitis, unspecified

Meningitis:purulent NOS

Meningitis:pyogenic NOS

Meningitis:suppurative NOS

G01 Meningitis in bacterial diseases classified elsewhere

Meningitis (in):anthrax

Meningitis (in):gonococcal

Meningitis (in):leptospirosis

Meningitis (in):listerial

Meningitis (in):Lyme disease

Meningitis (in):meningococcal

Meningitis (in):neurosyphilis

Meningitis (in):salmonella infection

Meningitis (in):syphilis:congenital

Meningitis (in):syphilis:secondary

Meningitis (in):tuberculous

Meningitis (in):typhoid fever

meningoencephalitis and meningomyelitis in bacterial diseases classified elsewhere

G02 Meningitis in other infectious and parasitic diseases classified elsewhere

meningoencephalitis and meningomyelitis in other infectious and parasitic diseases classified elsewhere

G02.0 Meningitis in viral diseases classified elsewhere

Meningitis (due to):adenoviral

Meningitis (due to):enteroviral

Meningitis (due to):herpesviral [herpes simplex]

Meningitis (due to):infectious mononucleosis

Meningitis (due to):measles

Meningitis (due to):mumps

Meningitis (due to):rubella

Meningitis (due to):varicella [chickenpox]

Meningitis (due to):zoster

G02.1 Meningitis in mycoses

Meningitis (in):candidal

Meningitis (in):coccidioidomycosis

Meningitis (in):cryptococcal

G02.8 Meningitis in other specified infectious and parasitic diseases classified elsewhere

Meningitis due to:African trypanosomiasis

Meningitis due to:Chagas disease (chronic)

G03 Meningitis due to other and unspecified causes

arachnoiditisdue to other and unspecified causes

leptomeningitisdue to other and unspecified causes

meningitisdue to other and unspecified causes

pachymeningitisdue to other and unspecified causes

meningoencephalitis

meningomyelitis

G03.0 Nonpyogenic meningitis

Nonbacterial meningitis

G03.1 Chronic meningitis

G03.2 Benign recurrent meningitis [Mollaret]

G03.8 Meningitis due to other specified causes

G03.9 Meningitis, unspecified

Arachnoiditis (spinal) NOS

G04 Encephalitis, myelitis and encephalomyelitis

acute ascending myelitis

meningoencephalitis

meningomyelitis

benign myalgic encephalomyelitis

encephalopathy:NOS

encephalopathy:alcoholic

encephalopathy:toxic

multiple sclerosis

myelitis:acute transverse

myelitis:subacute necrotizing

G04.0 Acute disseminated encephalitis

Encephalitispostimmunization

Encephalomyelitispostimmunization

Use additional external cause code (Chapter XX), if desired, to identify vaccine.

G04.1 Tropical spastic paraplegia

G04.2 Bacterial meningoencephalitis and meningomyelitis, not elsewhere classified

G04.8 Other encephalitis, myelitis and encephalomyelitis

Postinfectious encephalitis and encephalomyelitis NOS

G04.9 Encephalitis, myelitis and encephalomyelitis, unspecified

Ventriculitis (cerebral) NOS

G05 Encephalitis, myelitis and encephalomyelitis in diseases classified elsewhere

meningoencephalitis and meningomyelitis in diseases classified elsewhere

G05.0 Encephalitis, myelitis and encephalomyelitis in bacterial diseases classified elsewhere

Encephalitis, myelitis or encephalomyelitis (in):listerial

Encephalitis, myelitis or encephalomyelitis (in):meningococcal

Encephalitis, myelitis or encephalomyelitis (in):syphilis:congenital

Encephalitis, myelitis or encephalomyelitis (in):syphilis:late

Encephalitis, myelitis or encephalomyelitis (in):tuberculous

G05.1 Encephalitis, myelitis and encephalomyelitis in viral diseases classified elsewhere

Encephalitis, myelitis or encephalomyelitis (in):adenoviral

Encephalitis, myelitis or encephalomyelitis (in):cytomegaloviral

Encephalitis, myelitis or encephalomyelitis (in):enteroviral

Encephalitis, myelitis or encephalomyelitis (in):herpesviral [herpes simplex]

Encephalitis, myelitis or encephalomyelitis (in):influenza:seasonal virus identified

Encephalitis, myelitis or encephalomyelitis (in):influenza:virus not identified

Encephalitis, myelitis or encephalomyelitis (in):influenza:zoonotic or pandemic influenza virus identified

Encephalitis, myelitis or encephalomyelitis (in):measles

Encephalitis, myelitis or encephalomyelitis (in):mumps

Encephalitis, myelitis or encephalomyelitis (in):postchickenpox

Encephalitis, myelitis or encephalomyelitis (in):rubella

Encephalitis, myelitis or encephalomyelitis (in):zoster

G05.2 Encephalitis, myelitis and encephalomyelitis in other infectious and parasitic diseases classified elsewhere

Encephalitis, myelitis or encephalomyelitis in:African trypanosomiasis

Encephalitis, myelitis or encephalomyelitis in:Chagas disease (chronic)

Encephalitis, myelitis or encephalomyelitis in:naegleriasis

Encephalitis, myelitis or encephalomyelitis in:toxoplasmosis

Eosinophilic meningoencephalitis

G05.8 Encephalitis, myelitis and encephalomyelitis in other diseases classified elsewhere

Encephalitis in systemic lupus erythematosus

G06 Intracranial and intraspinal abscess and granuloma

Use additional code (B95-B98), if desired, to identify infectious agent.

G06.0 Intracranial abscess and granuloma

Abscess (embolic)(of):brain [any part]

Abscess (embolic)(of):cerebellar

Abscess (embolic)(of):cerebral

Abscess (embolic)(of):otogenic

Intracranial abscess or granuloma:epidural

Intracranial abscess or granuloma:extradural

Intracranial abscess or granuloma:subdural

G06.1 Intraspinal abscess and granuloma

Abscess (embolic) of spinal cord [any part]

Intraspinal abscess or granuloma:epidural

Intraspinal abscess or granuloma:extradural

Intraspinal abscess or granuloma:subdural

G06.2 Extradural and subdural abscess, unspecified

G07 Intracranial and intraspinal abscess and granuloma in diseases classified elsewhere

Abscess of brain:amoebic

Abscess of brain:gonococcal

Abscess of brain:tuberculous

Schistosomiasis granuloma of brain

Tuberculoma of:brain

Tuberculoma of:meninges

G08 Intracranial and intraspinal phlebitis and thrombophlebitis

Septic:embolismof intracranial or intraspinal venous sinuses and veins

Septic:endophlebitisof intracranial or intraspinal venous sinuses and veins

Septic:phlebitisof intracranial or intraspinal venous sinuses and veins

Septic:thrombophlebitisof intracranial or intraspinal venous sinuses and veins

Septic:thrombosisof intracranial or intraspinal venous sinuses and veins

intracranial phlebitis and thrombophlebitis:complicating:abortion or ectopic or molar pregnancy

intracranial phlebitis and thrombophlebitis:complicating:pregnancy, childbirth and the puerperium

intracranial phlebitis and thrombophlebitis:of nonpyogenic origin

nonpyogenic intraspinal phlebitis and thrombophlebitis

G09 Sequelae of inflammatory diseases of central nervous system

Category G09 is to be used to indicate conditions whose primary classification is to G00-G08 (i.e. excluding those marked with an asterisk (\*)) as the cause of sequelae, themselves classifiable elsewhere. The "sequelae"" include conditions specified as such or as late effects

Not to be used for chronic inflammatory diseases of the central nervous system. Code these to current inflammatory diseases of the central nervous system.

Systemic atrophies primarily affecting the central nervous system

G10-G14

G10 Huntington disease

Huntington chorea

G11 Hereditary ataxia

cerebral palsy

hereditary and idiopathic neuropathy

metabolic disorders

G11.0 Congenital nonprogressive ataxia

G11.1 Early-onset cerebellar ataxia

Early-onset cerebellar ataxia with:essential tremor

Early-onset cerebellar ataxia with:myoclonus [Hunt ataxia]

Early-onset cerebellar ataxia with:retained tendon reflexes

Friedreich ataxia (autosomal recessive)

X-linked recessive spinocerebellar ataxia

Onset usually before the age of 20

G11.2 Late-onset cerebellar ataxia

Onset usually after the age of 20

G11.3 Cerebellar ataxia with defective DNA repair

Ataxia telangiectasia [Louis-Bar]

Cockayne syndrome

xeroderma pigmentosum

G11.4 Hereditary spastic paraplegia

G11.8 Other hereditary ataxias

G11.9 Hereditary ataxia, unspecified

Hereditary cerebellar:ataxia NOS

Hereditary cerebellar:degeneration

Hereditary cerebellar:disease

Hereditary cerebellar:syndrome

G12 Spinal muscular atrophy and related syndromes

G12.0 Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]

G12.1 Other inherited spinal muscular atrophy

Progressive bulbar palsy of childhood [Fazio-Londe]

Spinal muscular atrophy:adult form

Spinal muscular atrophy:childhood form, type II

Spinal muscular atrophy:distal

Spinal muscular atrophy:juvenile form, type III [Kugelberg-Welander]

Spinal muscular atrophy:scapuloperoneal form

G12.2 Motor neuron disease

Familial motor neuron disease

Lateral sclerosis:amyotrophic

Lateral sclerosis:primary

Progressive:bulbar palsy

Progressive:spinal muscular atrophy

G12.8 Other spinal muscular atrophies and related syndromes

G12.9 Spinal muscular atrophy, unspecified

G13 Systemic atrophies primarily affecting central nervous system in diseases classified elsewhere

G13.0 Paraneoplastic neuromyopathy and neuropathy

Carcinomatous neuromyopathy

Sensorial paraneoplastic neuropathy [Denny Brown]

G13.1 Other systemic atrophy primarily affecting central nervous system in neoplastic disease

Paraneoplastic limbic encephalopathy

G13.2 Systemic atrophy primarily affecting central nervous system in myxoedema

G13.8 Systemic atrophy primarily affecting central nervous system in other diseases classified elsewhere

G14 Postpolio syndrome

Postpolio myelitic syndrome

sequelae of poliomyelitis

Extrapyramidal and movement disorders

G20-G26

G20 Parkinson disease

Hemiparkinsonism

Paralysis agitans

Parkinsonism or Parkinson disease:NOS

Parkinsonism or Parkinson disease:idiopathic

Parkinsonism or Parkinson disease:primary

G21 Secondary parkinsonism

G21.0 Malignant neuroleptic syndrome

Use additional external cause code (Chapter XX), if desired, to identify drug.

G21.1 Other drug-induced secondary parkinsonism

Use additional external cause code (Chapter XX), if desired, to identify drug.

G21.2 Secondary parkinsonism due to other external agents

Use additional external cause code (Chapter XX), if desired, to identify external agent.

G21.3 Postencephalitic parkinsonism

G21.8 Other secondary parkinsonism

G21.9 Secondary parkinsonism, unspecified

G22 Parkinsonism in diseases classified elsewhere

Syphilitic parkinsonism

G23 Other degenerative diseases of basal ganglia

G23.0 Hallervorden-Spatz disease

Pigmentary pallidal degeneration

G23.1 Progressive supranuclear ophthalmoplegia [Steele-Richardson-Olszewski]

G23.2 Multiple system atrophy, parkinsonian type [MSA-P]

G23.8 Other specified degenerative diseases of basal ganglia

Calcification of basal ganglia

Neurogenic orthostatic hypotension [Shy-Drager]

orthostatic hypotension NOS

G23.9 Degenerative disease of basal ganglia, unspecified

G24 Dystonia

dyskinesia

athetoid cerebral palsy

G24.0 Drug-induced dystonia

Use additional external cause code (Chapter XX), if desired, to identify drug.

G24.1 Idiopathic familial dystonia

Idiopathic dystonia NOS

G24.2 Idiopathic nonfamilial dystonia

G24.3 Spasmodic torticollis

torticollis NOS

G24.4 Idiopathic orofacial dystonia

Orofacial dyskinesia

G24.5 Blepharospasm

G24.8 Other dystonia

G24.9 Dystonia, unspecified

Dyskinesia NOS

G25 Other extrapyramidal and movement disorders

G25.0 Essential tremor

Familial tremor

tremor NOS

G25.1 Drug-induced tremor

Use additional external cause code (Chapter XX), if desired, to identify drug.

G25.2 Other specified forms of tremor

Intention tremor

G25.3 Myoclonus

Drug-induced myoclonus

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

facial myokymia

myoclonic epilepsy

G25.4 Drug-induced chorea

Use additional external cause code (Chapter XX), if desired, to identify drug.

G25.5 Other chorea

Chorea NOS

chorea NOS with heart involvement

Huntington chorea

rheumatic chorea

Sydenham chorea

G25.6 Drug-induced tics and other tics of organic origin

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

de la Tourette syndrome

tic NOS

G25.8 Other specified extrapyramidal and movement disorders

Akathisia (drug-induced) (treatment-induced)

Stiff-man syndrome

G25.9 Extrapyramidal and movement disorder, unspecified

G26 Extrapyramidal and movement disorders in diseases classified elsewhere

G23.3 Multiple system atrophy, cerebellar type [MSA-C]

G21.4 Vascular parkinsonism

Other degenerative diseases of the nervous system

G30-G32

G30 Alzheimer disease

senile and presenile forms

senile:degeneration of brain NEC

senile:dementia NOS

senility NOS

G30.0 Alzheimer disease with early onset

Onset usually before the age of 65

G30.1 Alzheimer disease with late onset

Onset usually after the age of 65

G30.8 Other Alzheimer disease

G30.9 Alzheimer disease, unspecified

G31 Other degenerative diseases of nervous system, not elsewhere classified

Reye syndrome

G31.0 Circumscribed brain atrophy

Pick disease

Progressive isolated aphasia

Frontotemporal dementia (FTD)

G31.1 Senile degeneration of brain, not elsewhere classified

Alzheimer disease

senility NOS

G31.2 Degeneration of nervous system due to alcohol

Alcoholic:cerebellar:ataxia

Alcoholic:cerebellar:degeneration

Alcoholic:cerebral degeneration

Alcoholic:encephalopathy

Dysfunction of autonomic nervous system due to alcohol

G31.8 Other specified degenerative diseases of nervous system

Grey-matter degeneration [Alpers]

Lewy body(ies)(dementia)(disease)

Subacute necrotizing encephalopathy [Leigh]

G31.9 Degenerative disease of nervous system, unspecified

G32 Other degenerative disorders of nervous system in diseases classified elsewhere

G32.0 Subacute combined degeneration of spinal cord in diseases classified elsewhere

Subacute combined degeneration of spinal cord in vitamin B deficiency

G32.8 Other specified degenerative disorders of nervous system in diseases classified elsewhere

Demyelinating diseases of the central nervous system

G35-G37

G35 Multiple sclerosis

Multiple sclerosis (of):NOS

Multiple sclerosis (of):brain stem

Multiple sclerosis (of):cord

Multiple sclerosis (of):disseminated

Multiple sclerosis (of):generalized

G36 Other acute disseminated demyelination

postinfectious encephalitis and encephalomyelitis NOS

G36.0 Neuromyelitis optica [Devic]

Demyelination in optic neuritis

optic neuritis NOS

G36.1 Acute and subacute haemorrhagic leukoencephalitis [Hurst]

G36.8 Other specified acute disseminated demyelination

G36.9 Acute disseminated demyelination, unspecified

G37 Other demyelinating diseases of central nervous system

G37.0 Diffuse sclerosis

Periaxial encephalitis

Schilder disease

adrenoleukodystrophy [Addison-Schilder]

G37.1 Central demyelination of corpus callosum

G37.2 Central pontine myelinolysis

G37.3 Acute transverse myelitis in demyelinating disease of central nervous system

Acute transverse myelitis NOS

multiple sclerosis

neuromyelitis optica [Devic]

G37.4 Subacute necrotizing myelitis

G37.5 Concentric sclerosis [Bal?]

G37.8 Other specified demyelinating diseases of central nervous system

G37.9 Demyelinating disease of central nervous system, unspecified

Episodic and paroxysmal disorders

G40-G47

G40 Epilepsy

Landau-Kleffner syndrome

seizure (convulsive) NOS

status epilepticus

Todd paralysis

G40.0 Localization-related (focal)(partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset

Benign childhood epilepsy with centrotemporal EEG spikes

Childhood epilepsy with occipital EEG paroxysms

G40.1 Localization-related (focal)(partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures

Attacks without alteration of consciousness

Simple partial seizures developing into secondarily generalized seizures

G40.2 Localization-related (focal)(partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures

Attacks with alteration of consciousness, often with automatisms

Complex partial seizures developing into secondarily generalized seizures

G40.3 Generalized idiopathic epilepsy and epileptic syndromes

Benign:myoclonic epilepsy in infancy

Benign:neonatal convulsions (familial)

Childhood absence epilepsy [pyknolepsy]

Epilepsy with grand mal seizures on awakening

Juvenile:absence epilepsy

Juvenile:myoclonic epilepsy [impulsive petit mal]

Nonspecific epileptic seizures:atonic

Nonspecific epileptic seizures:clonic

Nonspecific epileptic seizures:myoclonic

Nonspecific epileptic seizures:tonic

Nonspecific epileptic seizures:tonic-clonic

G40.4 Other generalized epilepsy and epileptic syndromes

Epilepsy with:myoclonic absences

Epilepsy with:myoclonic-astatic seizures

Infantile spasms

Lennox-Gastaut syndrome

Salaam attacks

Symptomatic early myoclonic encephalopathy

West syndrome

G40.5 Special epileptic syndromes

Epilepsia partialis continua [Kozhevnikof]

Epileptic seizures related to:alcohol

Epileptic seizures related to:drugs

Epileptic seizures related to:hormonal changes

Epileptic seizures related to:sleep deprivation

Epileptic seizures related to:stress

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

G40.6 Grand mal seizures, unspecified (with or without petit mal)

G40.7 Petit mal, unspecified, without grand mal seizures

G40.8 Other epilepsy

Epilepsies and epileptic syndromes undetermined as to whether they are focal or generalized

G40.9 Epilepsy, unspecified

Epileptic:convulsions NOS

Epileptic:fits NOS

Epileptic:seizures NOS

G41 Status epilepticus

G41.0 Grand mal status epilepticus

Tonic-clonic status epilepticus

epilepsia partialis continua [Kozhevnikof]

G41.1 Petit mal status epilepticus

Epileptic absence status

G41.2 Complex partial status epilepticus

G41.8 Other status epilepticus

G41.9 Status epilepticus, unspecified

G43 Migraine

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

headache NOS

G43.0 Migraine without aura [common migraine]

G43.1 Migraine with aura [classical migraine]

Migraine:aura without headache

Migraine:basilar

Migraine:equivalents

Migraine:familial hemiplegic

Migraine:with:acute-onset aura

Migraine:with:prolonged aura

Migraine:with:typical aura

G43.2 Status migrainosus

G43.3 Complicated migraine

G43.8 Other migraine

Ophthalmoplegic migraine

Retinal migraine

G43.9 Migraine, unspecified

G44 Other headache syndromes

atypical facial pain

headache NOS

trigeminal neuralgia

G44.0 Cluster headache syndrome

Chronic paroxysmal hemicrania

Cluster headache:chronic

Cluster headache:episodic

G44.1 Vascular headache, not elsewhere classified

Vascular headache NOS

G44.2 Tension-type headache

Chronic tension-type headache

Episodic tension headache

Tension headache NOS

G44.3 Chronic post-traumatic headache

G44.4 Drug-induced headache, not elsewhere classified

Use additional external cause code (Chapter XX), if desired, to identify drug.

G44.8 Other specified headache syndromes

G45 Transient cerebral ischaemic attacks and related syndromes

neonatal cerebral ischaemia

G45.0 Vertebro-basilar artery syndrome

G45.1 Carotid artery syndrome (hemispheric)

G45.2 Multiple and bilateral precerebral artery syndromes

G45.3 Amaurosis fugax

G45.4 Transient global amnesia

amnesia NOS

G45.8 Other transient cerebral ischaemic attacks and related syndromes

G45.9 Transient cerebral ischaemic attack, unspecified

Spasm of cerebral artery

Transient cerebral ischaemia NOS

G46 Vascular syndromes of brain in cerebrovascular diseases

G46.0 Middle cerebral artery syndrome

G46.1 Anterior cerebral artery syndrome

G46.2 Posterior cerebral artery syndrome

G46.3 Brain stem stroke syndrome

Syndrome:Benedikt

Syndrome:Claude

Syndrome:Foville

Syndrome:Millard-Gubler

Syndrome:Wallenberg

Syndrome:Weber

G46.4 Cerebellar stroke syndrome

G46.5 Pure motor lacunar syndrome

G46.6 Pure sensory lacunar syndrome

G46.7 Other lacunar syndromes

G46.8 Other vascular syndromes of brain in cerebrovascular diseases

G47 Sleep disorders

nightmares

nonorganic sleep disorders

sleep terrors

sleepwalking

G47.0 Disorders of initiating and maintaining sleep [insomnias]

G47.1 Disorders of excessive somnolence [hypersomnias]

G47.2 Disorders of the sleep-wake schedule

Delayed sleep phase syndrome

Irregular sleep-wake pattern

G47.3 Sleep apnoea

Sleep apnoea:central

Sleep apnoea:obstructive

pickwickian syndrome

sleep apnoea of newborn

G47.4 Narcolepsy and cataplexy

G47.8 Other sleep disorders

Kleine-Levin syndrome

G47.9 Sleep disorder, unspecified

Nerve, nerve root and plexus disorders

G50-G59

current traumatic nerve, nerve root and plexus disorders - see nerve injury by body region

neuralgiaNOS

neuritisNOS

peripheral neuritis in pregnancy

radiculitis NOS

G50 Disorders of trigeminal nerve

disorders of 5th cranial nerve

G50.0 Trigeminal neuralgia

Syndrome of paroxysmal facial pain

Tic douloureux

G50.1 Atypical facial pain

G50.8 Other disorders of trigeminal nerve

G50.9 Disorder of trigeminal nerve, unspecified

G51 Facial nerve disorders

disorders of 7th cranial nerve

G51.0 Bell palsy

Facial palsy

G51.1 Geniculate ganglionitis

postherpetic geniculate ganglionitis

G51.2 Melkersson syndrome

Melkersson-Rosenthal syndrome

G51.3 Clonic hemifacial spasm

G51.4 Facial myokymia

G51.8 Other disorders of facial nerve

G51.9 Disorder of facial nerve, unspecified

G52 Disorders of other cranial nerves

disorders of:acoustic [8th] nerve

disorders of:optic [2nd] nerve

paralytic strabismus due to nerve palsy

G52.0 Disorders of olfactory nerve

Disorder of 1st cranial nerve

G52.1 Disorders of glossopharyngeal nerve

Disorder of 9th cranial nerve

Glossopharyngeal neuralgia

G52.2 Disorders of vagus nerve

Disorder of pneumogastric [10th] nerve

G52.3 Disorders of hypoglossal nerve

Disorder of 12th cranial nerve

G52.7 Disorders of multiple cranial nerves

Polyneuritis cranialis

G52.8 Disorders of other specified cranial nerves

G52.9 Cranial nerve disorder, unspecified

G53 Cranial nerve disorders in diseases classified elsewhere

G53.0 Postzoster neuralgia

Postherpetic:geniculate ganglionitis

Postherpetic:trigeminal neuralgia

G53.1 Multiple cranial nerve palsies in infectious and parasitic diseases classified elsewhere

G53.2 Multiple cranial nerve palsies in sarcoidosis

G53.3 Multiple cranial nerve palsies in neoplastic disease

G53.8 Other cranial nerve disorders in other diseases classified elsewhere

G54 Nerve root and plexus disorders

current traumatic nerve root and plexus disorders - see nerve injury by body region

intervertebral disc disorders

neuralgia or neuritis NOS

neuritis or radiculitis:brachial NOS

neuritis or radiculitis:lumbar NOS

neuritis or radiculitis:lumbosacral NOS

neuritis or radiculitis:thoracic NOS

radiculitis NOS

radiculopathy NOS

spondylosis

G54.0 Brachial plexus disorders

Thoracic outlet syndrome

G54.1 Lumbosacral plexus disorders

G54.2 Cervical root disorders, not elsewhere classified

G54.3 Thoracic root disorders, not elsewhere classified

G54.4 Lumbosacral root disorders, not elsewhere classified

G54.5 Neuralgic amyotrophy

Parsonage-Aldren-Turner syndrome

Shoulder-girdle neuritis

G54.6 Phantom limb syndrome with pain

G54.7 Phantom limb syndrome without pain

Phantom limb syndrome NOS

G54.8 Other nerve root and plexus disorders

G54.9 Nerve root and plexus disorder, unspecified

G55 Nerve root and plexus compressions in diseases classified elsewhere

G55.0 Nerve root and plexus compressions in neoplastic disease

G55.1 Nerve root and plexus compressions in intervertebral disc disorders

G55.2 Nerve root and plexus compressions in spondylosis

G55.3 Nerve root and plexus compressions in other dorsopathies

G55.8 Nerve root and plexus compressions in other diseases classified elsewhere

G56 Mononeuropathies of upper limb

current traumatic nerve disorder - see nerve injury by body region

G56.0 Carpal tunnel syndrome

G56.1 Other lesions of median nerve

G56.2 Lesion of ulnar nerve

Tardy ulnar nerve palsy

G56.3 Lesion of radial nerve

G56.4 Causalgia

G56.8 Other mononeuropathies of upper limb

Interdigital neuroma of upper limb

G56.9 Mononeuropathy of upper limb, unspecified

G57 Mononeuropathies of lower limb

current traumatic nerve disorder - see nerve injury by body region

G57.0 Lesion of sciatic nerve

sciatica:NOS

sciatica:attributed to intervertebral disc disorder

G57.1 Meralgia paraesthetica

Lateral cutaneous nerve of thigh syndrome

G57.2 Lesion of femoral nerve

G57.3 Lesion of lateral popliteal nerve

Peroneal nerve palsy

G57.4 Lesion of medial popliteal nerve

G57.5 Tarsal tunnel syndrome

G57.6 Lesion of plantar nerve

Morton metatarsalgia

G57.8 Other mononeuropathies of lower limb

Interdigital neuroma of lower limb

G57.9 Mononeuropathy of lower limb, unspecified

G58 Other mononeuropathies

G58.0 Intercostal neuropathy

G58.7 Mononeuritis multiplex

G58.8 Other specified mononeuropathies

G58.9 Mononeuropathy, unspecified

G59 Mononeuropathy in diseases classified elsewhere

G59.0 Diabetic mononeuropathy

G59.8 Other mononeuropathies in diseases classified elsewhere

Polyneuropathies and other disorders of the peripheral nervous system

G60-G64

neuralgia NOS

neuritis NOS

peripheral neuritis in pregnancy

radiculitis NOS

G60 Hereditary and idiopathic neuropathy

G60.0 Hereditary motor and sensory neuropathy

Disease:Charcot-Marie-Tooth

Disease:D?jerine-Sottas

Hereditary motor and sensory neuropathy, types I-IV

Hypertrophic neuropathy of infancy

Peroneal muscular atrophy (axonal type)(hypertrophic type)

Roussy-L?vy syndrome

G60.1 Refsum disease

G60.2 Neuropathy in association with hereditary ataxia

G60.3 Idiopathic progressive neuropathy

G60.8 Other hereditary and idiopathic neuropathies

Morvan disease

Nelaton syndrome

Sensory neuropathy:dominantly inherited

Sensory neuropathy:recessively inherited

G60.9 Hereditary and idiopathic neuropathy, unspecified

G61 Inflammatory polyneuropathy

G61.0 Guillain-Barr? syndrome

Acute (post-)infective polyneuritis

Miller Fisher Syndrome

G61.1 Serum neuropathy

Use additional external cause code (Chapter XX), if desired, to identify cause.

G61.8 Other inflammatory polyneuropathies

G61.9 Inflammatory polyneuropathy, unspecified

G62 Other polyneuropathies

G62.0 Drug-induced polyneuropathy

Use additional external cause code (Chapter XX), if desired, to identify drug.

G62.1 Alcoholic polyneuropathy

G62.2 Polyneuropathy due to other toxic agents

Use additional external cause code (Chapter XX), if desired, to identify toxic agent.

G62.8 Other specified polyneuropathies

Radiation-induced polyneuropathy

Use additional external cause code (Chapter XX), if desired, to identify cause.

G62.9 Polyneuropathy, unspecified

Neuropathy NOS

G63 Polyneuropathy in diseases classified elsewhere

G63.0 Polyneuropathy in infectious and parasitic diseases classified elsewhere

Polyneuropathy (in):diphtheria

Polyneuropathy (in):infectious mononucleosis

Polyneuropathy (in):leprosy

Polyneuropathy (in):Lyme disease

Polyneuropathy (in):mumps

Polyneuropathy (in):postherpetic

Polyneuropathy (in):syphilis, late

Polyneuropathy (in):congenital syphilis, late

Polyneuropathy (in):tuberculous

G63.1 Polyneuropathy in neoplastic disease

G63.2 Diabetic polyneuropathy

G63.3 Polyneuropathy in other endocrine and metabolic diseases

G63.4 Polyneuropathy in nutritional deficiency

G63.5 Polyneuropathy in systemic connective tissue disorders

G63.6 Polyneuropathy in other musculoskeletal disorders

G63.8 Polyneuropathy in other diseases classified elsewhere

Uraemic neuropathy

G64 Other disorders of peripheral nervous system

Disorder of peripheral nervous system NOS

Diseases of myoneural junction and muscle

G70-G73

G70 Myasthenia gravis and other myoneural disorders

botulism

transient neonatal myasthenia gravis

G70.0 Myasthenia gravis

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

G70.1 Toxic myoneural disorders

Use additional external cause code (Chapter XX), if desired, to identify toxic agent.

G70.2 Congenital and developmental myasthenia

G70.8 Other specified myoneural disorders

G70.9 Myoneural disorder, unspecified

G71 Primary disorders of muscles

arthrogryposis multiplex congenita

metabolic disorders

myositis

G71.0 Muscular dystrophy

Muscular dystrophy:autosomal recessive, childhood type, resembling Duchenne or Becker

Muscular dystrophy:benign [Becker]

Muscular dystrophy:benign scapuloperoneal with early contractures [Emery-Dreifuss]

Muscular dystrophy:distal

Muscular dystrophy:facioscapulohumeral

Muscular dystrophy:limb-girdle

Muscular dystrophy:ocular

Muscular dystrophy:oculopharyngeal

Muscular dystrophy:scapuloperoneal

Muscular dystrophy:severe [Duchenne]

congenital muscular dystrophy:NOS

congenital muscular dystrophy:with specific morphological abnormalities of the muscle fibre

G71.1 Myotonic disorders

Dystrophia myotonica [Steinert]

Myotonia:chondrodystrophic

Myotonia:drug-induced

Myotonia:symptomatic

Myotonia congenita:NOS

Myotonia congenita:dominant [Thomsen]

Myotonia congenita:recessive [Becker]

Neuromyotonia [Isaacs]

Paramyotonia congenita

Pseudomyotonia

Use additional external cause code (Chapter XX), if desired, to identify drug, if drug-induced.

G71.2 Congenital myopathies

Congenital muscular dystrophy:NOS

Congenital muscular dystrophy:with specific morphological abnormalities of the muscle fibre

Disease:central core

Disease:minicore

Disease:multicore

Fibre-type disproportion

Myopathy:myotubular (centronuclear)

Myopathy:nemaline

G71.3 Mitochondrial myopathy, not elsewhere classified

G71.8 Other primary disorders of muscles

G71.9 Primary disorder of muscle, unspecified

Hereditary myopathy NOS

G72 Other myopathies

arthrogryposis multiplex congenita

dermatopolymyositis

ischaemic infarction of muscle

myositis

polymyositis

G72.0 Drug-induced myopathy

Use additional external cause code (Chapter XX), if desired, to identify drug.

G72.1 Alcoholic myopathy

G72.2 Myopathy due to other toxic agents

Use additional external cause code (Chapter XX), if desired, to identify toxic agent.

G72.3 Periodic paralysis

Periodic paralysis (familial):hyperkalaemic

Periodic paralysis (familial):hypokalaemic

Periodic paralysis (familial):myotonic

Periodic paralysis (familial):normokalaemic

G72.4 Inflammatory myopathy, not elsewhere classified

G72.8 Other specified myopathies

G72.9 Myopathy, unspecified

G73 Disorders of myoneural junction and muscle in diseases classified elsewhere

G73.0 Myasthenic syndromes in endocrine diseases

Myasthenic syndromes in:diabetic amyotrophy

Myasthenic syndromes in:thyrotoxicosis [hyperthyroidism]

G73.1 Lambert-Eaton syndrome

G73.2 Other myasthenic syndromes in neoplastic disease

G73.3 Myasthenic syndromes in other diseases classified elsewhere

G73.4 Myopathy in infectious and parasitic diseases classified elsewhere

G73.5 Myopathy in endocrine diseases

Myopathy in:hyperparathyroidism

Myopathy in:hypoparathyroidism

Thyrotoxic myopathy

G73.6 Myopathy in metabolic diseases

Myopathy in:glycogen storage disease

Myopathy in:lipid storage disorders

G73.7 Myopathy in other diseases classified elsewhere

Myopathy in:rheumatoid arthritis

Myopathy in:scleroderma

Myopathy in:sicca syndrome [Sj?gren]

Myopathy in:systemic lupus erythematosus

Cerebral palsy and other paralytic syndromes

G80-G83

G80 Cerebral palsy

hereditary spastic paraplegia

G80.0 Spastic quadriplegic cerebral palsy

Spastic tetraplegic cerebral palsy

G80.1 Spastic diplegic cerebral palsy

Congenital spastic paralysis (cerebral)

Spastic cerebral palsy NOS

G80.2 Spastic hemiplegic cerebral palsy

G80.3 Dyskinetic cerebral palsy

Athetoid cerebral palsy

Dystonic cerebral palsy

G80.4 Ataxic cerebral palsy

G80.8 Other cerebral palsy

Mixed cerebral palsy syndromes

G80.9 Cerebral palsy, unspecified

Cerebral palsy NOS

G81 Hemiplegia

For primary coding, this category is to be used only when hemiplegia (complete) (incomplete) is reported without further specification, or is stated to be old or longstanding but of unspecified cause. The category is also for use in multiple coding to identify these types of hemiplegia resulting from any cause.

congenital cerebral palsy

G81.0 Flaccid hemiplegia

G81.1 Spastic hemiplegia

G81.9 Hemiplegia, unspecified

G82 Paraplegia and tetraplegia

For primary coding, this category is to be used only when the listed conditions are reported without further specification, or are stated to be old or longstanding but of unspecified cause. The category is also for use in multiple coding to identify these conditions resulting from any cause.

congenital cerebral palsy

G82.0 Flaccid paraplegia

G82.1 Spastic paraplegia

G82.2 Paraplegia, unspecified

Paralysis of both lower limbs NOS

Paraplegia (lower) NOS

G82.3 Flaccid tetraplegia

G82.4 Spastic tetraplegia

G82.5 Tetraplegia, unspecified

Quadriplegia NOS

G83 Other paralytic syndromes

paralysis (complete)(incomplete), except as in G80-G82

For primary coding, this category is to be used only when the listed conditions are reported without further specification, or are stated to be old or longstanding but of unspecified cause. The category is also for use in multiple coding to identify these conditions resulting from any cause.

G83.0 Diplegia of upper limbs

Diplegia (upper)

Paralysis of both upper limbs

G83.1 Monoplegia of lower limb

Paralysis of lower limb

G83.2 Monoplegia of upper limb

Paralysis of upper limb

G83.3 Monoplegia, unspecified

G83.4 Cauda equina syndrome

Neurogenic bladder due to cauda equina syndrome

cord bladder NOS

G83.8 Other specified paralytic syndromes

Todd paralysis (postepileptic)

G83.9 Paralytic syndrome, unspecified

G83.5 Locked-in syndrome

Other disorders of the nervous system

G90-G99

G90 Disorders of autonomic nervous system

dysfunction of autonomic nervous system due to alcohol

G90.0 Idiopathic peripheral autonomic neuropathy

Carotid sinus syncope

G90.1 Familial dysautonomia [Riley-Day]

G90.2 Horner syndrome

Bernard(-Horner) syndrome

G90.4 Autonomic dysreflexia

G90.8 Other disorders of autonomic nervous system

G90.9 Disorder of autonomic nervous system, unspecified

G91 Hydrocephalus

acquired hydrocephalus

hydrocephalus:congenital

hydrocephalus:acquired, of newborn

hydrocephalus:due to congenital toxoplasmosis

G91.0 Communicating hydrocephalus

G91.1 Obstructive hydrocephalus

G91.2 Normal-pressure hydrocephalus

G91.3 Post-traumatic hydrocephalus, unspecified

G91.8 Other hydrocephalus

G91.9 Hydrocephalus, unspecified

G92 Toxic encephalopathy

Use additional external cause code (Chapter XX), if desired, to identify toxic agent.

G93 Other disorders of brain

G93.0 Cerebral cysts

Arachnoid cyst

Porencephalic cyst, acquired

acquired periventricular cysts of newborn

congenital cerebral cysts

G93.1 Anoxic brain damage, not elsewhere classified

complicating:abortion or ectopic or molar pregnancy

complicating:pregnancy, labour or delivery

complicating:surgical and medical care

neonatal anoxia

G93.2 Benign intracranial hypertension

hypertensive encephalopathy

G93.3 Postviral fatigue syndrome

Benign myalgic encephalomyelitis

G93.4 Encephalopathy, unspecified

encephalopathy:alcoholic

encephalopathy:toxic

G93.5 Compression of brain

Compressionof brain (stem)

Herniationof brain (stem)

traumatic compression of brain (diffuse)

traumatic compression of brain (diffuse)focal

G93.6 Cerebral oedema

cerebral oedema:due to birth injury

cerebral oedema:traumatic

G93.7 Reye syndrome

Use additional external cause code (Chapter XX), if desired, to identify cause.

G93.8 Other specified disorders of brain

Postradiation encephalopathy

Use additional external cause code (Chapter XX), if desired, to identify cause.

G93.9 Disorder of brain, unspecified

G94 Other disorders of brain in diseases classified elsewhere

G94.0 Hydrocephalus in infectious and parasitic diseases classified elsewhere

G94.1 Hydrocephalus in neoplastic disease

G94.2 Hydrocephalus in other diseases classified elsewhere

G94.8 Other specified disorders of brain in diseases classified elsewhere

G95 Other diseases of spinal cord

myelitis

G95.0 Syringomyelia and syringobulbia

G95.1 Vascular myelopathies

Acute infarction of spinal cord (embolic)(nonembolic)

Arterial thrombosis of spinal cord

Haematomyelia

Nonpyogenic intraspinal phlebitis and thrombophlebitis

Oedema of spinal cord

Subacute necrotic myelopathy

intraspinal phlebitis and thrombophlebitis, except non-pyogenic

G95.2 Cord compression, unspecified

G95.8 Other specified diseases of spinal cord

Cord bladder NOS

Myelopathy:drug-induced

Myelopathy:radiation-induced

Use additional external cause code (Chapter XX), if desired, to identify external agent.

neurogenic bladder:NOS

neurogenic bladder:due to cauda equina syndrome

neuromuscular dysfunction of bladder without mention of spinal cord lesion

G95.9 Disease of spinal cord, unspecified

Myelopathy NOS

G96 Other disorders of central nervous system

G96.0 Cerebrospinal fluid leak

from spinal puncture

G96.1 Disorders of meninges, not elsewhere classified

Meningeal adhesions (cerebral)(spinal)

G96.8 Other specified disorders of central nervous system

G96.9 Disorder of central nervous system, unspecified

G97 Postprocedural disorders of nervous system, not elsewhere classified

G97.0 Cerebrospinal fluid leak from spinal puncture

G97.1 Other reaction to spinal and lumbar puncture

G97.2 Intracranial hypotension following ventricular shunting

G97.8 Other postprocedural disorders of nervous system

G97.9 Postprocedural disorder of nervous system, unspecified

G98 Other disorders of nervous system, not elsewhere classified

Nervous system disorder NOS

G99 Other disorders of nervous system in diseases classified elsewhere

G99.0 Autonomic neuropathy in endocrine and metabolic diseases

Amyloid autonomic neuropathy

Diabetic autonomic neuropathy

G99.1 Other disorders of autonomic nervous system in other diseases classified elsewhere

G99.2 Myelopathy in diseases classified elsewhere

Anterior spinal and vertebral artery compression syndromes

Myelopathy in:intervertebral disc disorders

Myelopathy in:neoplastic disease

Myelopathy in:spondylosis

G99.8 Other specified disorders of nervous system in diseases classified elsewhere

Uraemic paralysis