2014 ICD-10-CM Diagnosis Codes
Related to Speech, Language, and Swallowing Disorders

The codes in ICD-10 are not valid for any purpose or use in the United States until October 1, 2015.
General Information

This ASHA document provides a listing of the 2014 International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) codes related to speech, language, and swallowing disorders. This document is not a comprehensive list and a number of codes are included for information purposes only. Entries with only three or four digits may require coding to a higher degree of specificity than indicated here. However, in general, speech-language pathology related diagnoses will be listed to their highest level of specificity.

The codes in ICD-10 are not valid for any purpose or use in the United States until October 1, 2015. For more information on the transition, see www.asha.org/Practice/reimbursement/coding/ICD-10/.

For the most up-to-date information on ICD coding, go to ASHA’s Billing and Reimbursement website at www.asha.org/practice/reimbursement/coding/.

For additional information, contact the health care economics and advocacy team by e-mail at reimbursement@asha.org.
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## ICD-10-CM Tabular List of Diseases and Injuries

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ICD-10-CM Diagnostic Codes

Note: On April 1, 2014, President Obama signed into law the Protecting Access to Medicare Act of 2014. This law included a provision that delays the implementation of ICD-10 from October 1, 2014 to October 1, 2015.

Overview
On October 1, 2015, the International Classification of Diseases, 10th Revision (ICD-10) will replace ICD-9 (9th Revision) as the official system of assigning codes to diagnoses and procedures associated with hospital utilization in the United States. The ICD is also used to code and classify mortality data from death certificates.

The new ICD-10 will include the ICD-10-CM (clinical modification) and ICD-10-PCS (procedure coding system). The ICD-10 is owned by the World Health Organization (WHO). The clinical modification was developed by the Centers for Disease Control and Prevention for use in all U.S. health care treatment settings. The procedure coding system (i.e., ICD-9-PCS and ICD-10-PCS) was developed by the Centers for Medicare and Medicaid Services for use in the U.S. for inpatient hospital settings only. This product only includes speech-language pathology related ICD-10-CM codes.

Scope
The intent of ICD-10-CM is to standardize disease and procedure classification throughout the United States and to gather data about basic health statistics.

Purpose
HIPAA legislation requires the ICD-10-CM to be used for health services billing and record keeping. As noted above, the effective implementation date for ICD-10-CM (and ICD-10-PCS) is October 1, 2015. Updates to this version of ICD-10-CM are anticipated prior to its implementation.

Relation to Professional Scope of Practice
The speech-language pathologist practicing in a health care setting, especially a hospital, may have to code delivery of services according to the ICD-10-CM.

Official ICD-10-CM Websites
- National Center for Health Statistics: www.cdc.gov/nchs/icd/icd10.htm
- Centers for Medicare and Medicaid Services: www.cms.gov/ICD10/

ASHA Resources
- ICD-9 to ICD-10 Mapping Tool for Audiologists and Speech-Language Pathologists: www.asha.org/icdmapping.aspx
- ICD-10-CM Diagnosis Codes for Audiology and Speech-Language Pathology: www.asha.org/Practice/reimbursement/coding/ICD-10/
- ICD-9-CM Diagnosis Codes for Speech-Language Pathology: www.asha.org/practice/reimbursement/coding/icd9SLP/
- Coding Normal Results: www.asha.org/practice/reimbursement/coding/normalresults/
- Coding to the Highest Degree of Specificity: www.asha.org/practice/reimbursement/coding/codespecificity/
ICD-10-CM Tabular List of Diseases and Injuries

Related to speech, language, and swallowing disorders

**Note:** This is *not* a comprehensive list and a number of codes are included for information purposes only. Some categories of codes (e.g., neoplasms) may be more extensive, contain additional instructional notes, and may also require coding to a higher degree of specificity than indicated here. However, in general, speech-language pathology related diagnoses will be listed to their highest level of specificity. For a full list of ICD-10-CM codes, descriptors, and instructions, see the official ICD-10-CM publication at ftp://ftp.cdc.gov/pub/Health_Statistics/NCHS/Publications/ICD10CM/2014/.

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### Ch. 2 Neoplasms (C00-D49)

*Malignant neoplasms of lip, oral cavity, and pharynx (C00-C14)*

- **C00** Malignant neoplasm of lip
  - C00.0 Malignant neoplasm of external upper lip
  - C00.1 Malignant neoplasm of external lower lip
  - C00.2 Malignant neoplasm of external lip, unspecified
  - C00.3 Malignant neoplasm of upper lip, inner aspect
  - C00.4 Malignant neoplasm of lower lip, inner aspect
  - C00.5 Malignant neoplasm of lip, unspecified, inner aspect
  - C00.6 Malignant neoplasm of commissure of lip, unspecified
  - C00.8 Malignant neoplasm of overlapping sites of lip
  - C00.9 Malignant neoplasm of lip, unspecified

- **C01** Malignant neoplasm of base of tongue

- **C02** Malignant neoplasm of other and unspecified parts of tongue
  - C02.0 Malignant neoplasm of dorsal surface of tongue
  - C02.1 Malignant neoplasm of border of tongue
  - C02.2 Malignant neoplasm of ventral surface of tongue
  - C02.3 Malignant neoplasm of anterior two-thirds of tongue, part unspecified
  - C02.4 Malignant neoplasm of lingual tonsil
  - C02.8 Malignant neoplasm of overlapping sites of tongue
  - C02.9 Malignant neoplasm of tongue, unspecified

- **◊ C03** Malignant neoplasm of gum

- **◊ C04** Malignant neoplasm of floor of mouth

- **C05** Malignant neoplasm of palate
  - C05.0 Malignant neoplasm of hard palate
  - C05.1 Malignant neoplasm of soft palate
  - C05.2 Malignant neoplasm of uvula

- **◊ C06** Malignant neoplasm of other and unspecified parts of mouth

- **◊ C08** Malignant neoplasm of other and unspecified major salivary glands

- **◊ C09** Malignant neoplasm of tonsil

- **C10** Malignant neoplasm of oropharynx
  - C10.1 Malignant neoplasm of anterior surface of epiglottis
  - C10.2 Malignant neoplasm of lateral wall of oropharynx
  - C10.3 Malignant neoplasm of posterior wall of oropharynx

- **C11** Malignant neoplasm of nasopharynx
  - C11.0 Malignant neoplasm of superior wall of nasopharynx

- **✓** Code typically used by SLPs
- **◊** Additional digits not listed here
C11.1 Malignant neoplasm of posterior wall of nasopharynx
C11.2 Malignant neoplasm of lateral wall of nasopharynx
C11.3 Malignant neoplasm of anterior wall of nasopharynx
C11.8 Malignant neoplasm of overlapping sites of nasopharynx
C14 Malignant neoplasm of other and ill-defined sites in the lip, oral cavity and pharynx
C14.0 Malignant neoplasm of pharynx, unspecified
◊ C15 Malignant neoplasm of esophagus

Malignant neoplasms of digestive organs (C15-C26)

Malignant neoplasms of respiratory and intrathoracic organs (C30-C39)
◊ C30 Malignant neoplasm of nasal cavity and middle ear
C32 Malignant neoplasm of larynx
  C32.0 Malignant neoplasm of glottis
  C32.1 Malignant neoplasm of supraglottis
  C32.2 Malignant neoplasm of subglottis
  C32.3 Malignant neoplasm of laryngeal cartilage
  C32.8 Malignant neoplasm of overlapping sites of larynx
  C32.9 Malignant neoplasm of larynx, unspecified
◊ C33 Malignant neoplasm of trachea

Malignant neoplasms of eye, brain and other parts of central nervous system (C69-C72)
◊ C71 Malignant neoplasm of brain

In situ neoplasms (D00-D09)
D00 Carcinoma in situ of oral cavity, esophagus and stomach
  D00.07 Carcinoma in situ of tongue
D02 Carcinoma in situ of middle ear and respiratory system
  D02.1 Carcinoma in situ of trachea

Benign neoplasms, except benign neuroendocrine tumors (D10-D36)
◊ D10 Benign neoplasm of mouth and pharynx
D14 Benign neoplasm of middle ear and respiratory system
  D14.1 Benign neoplasm of larynx
  D14.2 Benign neoplasm of trachea
◊ D33 Benign neoplasm of brain and other parts of central nervous system
◊ D38 Neoplasm of uncertain behavior of middle ear and respiratory and intrathoracic organs

Ch. 5 Mental, behavioral, and neurodevelopmental disorders (F01-F99)

Mental disorders due to known physiological conditions (F01-F09)
F01 Vascular dementia
  F01.5 Vascular dementia
    F01.50 Vascular dementia without behavioral disturbance
    F01.51 Vascular dementia with behavioral disturbance
F02 Dementia in other diseases classified elsewhere
  Code first the underlying physiological condition, such as:
    Alzheimer's (G30.-)
    cerebral lipidosis (E75.4)
    Creutzfeldt-Jakob disease (A81.0-)

✓ Code typically used by SLPs  ◊ Additional digits not listed here
dementia with Lewy bodies (G31.83)
epilepsy and recurrent seizures (G40.-)
frontotemporal dementia (G31.09)
hepatolenticular degeneration (E83.0)
human immunodeficiency virus [HIV] disease (B20)
hypercalcemia (E83.52)
hypothyroidism, acquired (E00-E03.-)
intoxications (T36-T65)
Jakob-Creutzfeldt disease (A81.0-)
multiple sclerosis (G35)
neurosyphilis (A52.17)
niacin deficiency [pellagra] (E52)
Parkinson's disease (G20)
Pick's disease (G31.01)
polyarteritis nodosa (M30.0)
systemic lupus erythematosus (M32.-)
trypanosomiasis (B56.-, B57.-)
vitamin B deficiency (E53.8)

F02.8  Dementia in other diseases classified elsewhere
F02.80  Dementia in other diseases classified elsewhere, without behavioral disturbance
F02.81  Dementia in other diseases classified elsewhere, with behavioral disturbance

F03  Unspecified dementia
F03.9  Unspecified dementia
F03.90  Unspecified dementia without behavioral disturbance
Dementia NOS
F03.91  Unspecified dementia with behavioral disturbance
Unspecified dementia with aggressive behavior
Unspecified dementia with combative behavior
Unspecified dementia with violent behavior

Use additional code, if applicable, to identify wandering in unspecified dementia (Z91.83)

Schizophrenia, schizotypal, delusional, and other non-mood psychotic disorders (F20-F29)

◊ F20  Schizophrenia

Intellectual Disabilities (F70-F79)

Code first any associated physical or developmental disorders

Excludes1: borderline intellectual functioning, IQ above 70 to 84 (R41.83)

F70  Mild intellectual disabilities
IQ level 50-55 to approximately 70
Mild mental subnormality

F71  Moderate intellectual disabilities
IQ level 35-40 to 50-55
Moderate mental subnormality

F72  Severe intellectual disabilities
IQ 20-25 to 35-40
Severe mental subnormality
F73  Profound intellectual disabilities
   IQ level below 20-25
   Profound mental subnormality
F78  Other intellectual disabilities
F79  Unspecified intellectual disabilities
   Mental deficiency NOS
   Mental subnormality NOS

**Pervasive and specific developmental disorders (F80-F89)**

F80  Specific developmental disorders of speech and language

✓ F80.0  Phonological disorder
   Dyslalia
   Functional speech articulation disorder
   Lalling
   Lisping
   Phonological developmental disorder
   Speech articulation developmental disorder
   **Excludes1:** speech articulation impairment due to aphasia NOS (R47.01)
   speech articulation impairment due to apraxia (R48.2)
   **Excludes2:** speech articulation impairment due to hearing loss (F80.4)
   speech articulation impairment due to intellectual disabilities (F70-F79)
   speech articulation impairment with expressive language developmental disorder (F80.1)
   speech articulation impairment with mixed receptive expressive language developmental disorder (F80.2)

✓ F80.1  Expressive language disorder
   Developmental dysphasia or aphasia, expressive type
   **Excludes1:** mixed receptive-expressive language disorder (F80.2)
   dysphasia and aphasia NOS (R47.-)
   **Excludes2:** acquired aphasia with epilepsy [Landau-Kleffner] (G40.80-)
   selective mutism (F94.0)
   intellectual disabilities (F70-F79)
   pervasive developmental disorders (F84.-)

✓ F80.2  Mixed receptive-expressive language disorder
   Developmental dysphasia or aphasia, receptive type
   Developmental Wernicke's aphasia
   **Excludes1:** central auditory processing disorder (H93.25)
   dysphasia or aphasia NOS (R47.-)
   expressive language disorder (F80.1)
   expressive type dysphasia or aphasia (F80.1)
   word deafness (H93.25)
   **Excludes2:** acquired aphasia with epilepsy [Landau-Kleffner] (G40.80-)
   pervasive developmental disorders (F84.-)
   selective mutism (F94.0)
   intellectual disabilities (F70-F79)

✓ F80.4  Speech and language development delay due to hearing loss
   **Code also** type of hearing loss (H90.-, H91.-)

F80.8  Other developmental disorders of speech and language

✓ Code typically used by SLPs  ○ Additional digits not listed here
F80.81 Childhood onset fluency disorder
Cluttering NOS
Stuttering NOS
**Excludes1:** adult onset fluency disorder (F98.5)
fluency disorder in conditions classified elsewhere (R47.82)
fluency disorder (stuttering) following cerebrovascular disease (I69. with final characters-23)

F80.89 Other developmental disorders of speech and language

F80.9 Developmental disorder of speech and language, unspecified
Communication disorder NOS
Language disorder NOS

F81 Specific developmental disorders of scholastic skills
F81.0 Specific reading disorder
'Backward reading'
Developmental dyslexia
Specific reading retardation
**Excludes1:** alexia NOS (R48.0)
dyslexia NOS (R48.0)

F81.2 Mathematics disorder
Developmental acalculia
Developmental arithmetical disorder
Developmental Gerstmann's syndrome
**Excludes1:** acalculia NOS (R48.8)
**Excludes2:** arithmetical difficulties associated with a reading disorder (F81.0)
arithmetical difficulties associated with a spelling disorder (F81.81)
arithmetical difficulties due to inadequate teaching (Z55.8)

F81.8 Other developmental disorders of scholastic skills
F81.81 Disorder of written expression
Specific spelling disorder
F81.89 Other developmental disorders of scholastic skills

F81.9 Developmental disorder of scholastic skills, unspecified
Knowledge acquisition disability NOS
Learning disability NOS
Learning disorder NOS

F82 Specific developmental disorder of motor function
Clumsy child syndrome
Developmental coordination disorder
Developmental dyspraxia
**Excludes1:** abnormalities of gait and mobility (R26.-)
lack of coordination (R27.-)
**Excludes2:** lack of coordination secondary to intellectual disabilities (F70-F79)

F84 Pervasive developmental disorders
**Use additional** code to identify any associated medical condition and intellectual disabilities.

F84.0 Autistic disorder
Infantile autism
Infantile psychosis
Kanner's syndrome
**Excludes1:** Asperger's syndrome (F84.5)
F84.2  Rett's syndrome
   Excludes1:  Asperger's syndrome (F84.5)
               Autistic disorder (F84.0)
               Other childhood disintegrative disorder (F84.3)

F84.3  Other childhood disintegrative disorder
   Dementia infantilis
   Disintegrative psychosis
   Heller's syndrome
   Symbiotic psychosis
   Use additional code to identify any associated neurological condition.
   Excludes1:  Asperger's syndrome (F84.5)
               Autistic disorder (F84.0)
               Rett's syndrome (F84.2)

✓ F84.5  Asperger's syndrome
   Asperger's disorder
   Autistic psychopathy
   Schizoid disorder of childhood

✓ F84.8  Other pervasive developmental disorders
   Overactive disorder associated with intellectual disabilities and stereotyped movements

F84.9  Pervasive developmental disorder, unspecified
   Atypical autism

F88  Other disorders of psychological development
   Developmental agnosia

F89  Unspecified disorder of psychological development

Behavioral and emotional disorders with onset usually occurring in childhood and adolescence (F90-F98)

F90  Attention-deficit hyperactivity disorders
   Includes:  attention deficit disorder with hyperactivity
              attention deficit syndrome with hyperactivity
   Excludes2:  anxiety disorders (F40.-, F41.-)
               mood [affective] disorders (F30-F39)
               pervasive developmental disorders (F84.)
               schizophrenia (F20.)

F90.0  Attention-deficit hyperactivity disorder, predominantly inattentive type
F90.1  Attention-deficit hyperactivity disorder, predominantly hyperactive type
F90.2  Attention-deficit hyperactivity disorder, combined type
F90.8  Attention-deficit hyperactivity disorder, other type
F90.9  Attention-deficit hyperactivity disorder, unspecified type

F94  Disorders of social functioning with onset specific to childhood and adolescence
F94.0  Selective mutism
   Elective mutism
   Excludes2:  pervasive developmental disorders (F84.)
               schizophrenia (F20.)
               specific developmental disorders of speech and language (F80.)
               transient mutism as part of separation anxiety in young children (F93.0)

F98  Other behavioral and emotional disorders with onset usually occurring in childhood and adolescence

✓ Code typically used by SLPs  ◆ Additional digits not listed here
F98.5  Adult onset fluency disorder
**Excludes1:** childhood onset fluency disorder (F80.81)
  dysphasia (R47.02)
  fluency disorder in conditions classified elsewhere (R47.82)
  fluency disorder (stuttering) following cerebrovascular disease (I69. with
  final characters -23)
  tic disorders (F95.-)

**Ch. 6  Diseases of the nervous system (G00-G99)**

**Inflammatory diseases of the central nervous system (G00-G09)**

G00  Bacterial meningitis, not elsewhere classified
  G00.0  Hemophilus meningitis
  G00.1  Pneumococcal meningitis
  G00.2  Streptococcal meningitis
    **Use additional** code to further identify organism (B95.0-B95.5)
  G00.3  Staphylococcal meningitis
    **Use additional** code to further identify organism (B95.6-B95.8)
  G00.8  Other bacterial meningitis
  G00.9  Bacterial meningitis, unspecified

◊ G04  Encephalitis, myelitis and encephalomyelitis

**Systemic atrophies primarily affecting the central nervous system (G10-G14)**

G10  Huntington's disease
G12  Spinal muscular atrophy and related syndromes
  G12.2  Motor neuron disease
    G12.21  Amyotrophic lateral sclerosis

**Extrapyramidal and movement disorders (G20-G26)**

G20  Parkinson’s disease
  Hemiparkinsonism
  Idiopathic Parkinsonism or Parkinson’s disease
  Paralysis agitans
  Parkinsonism or Parkinson’s disease NOS
  Primary Parkinsonism or Parkinson’s disease
  **Excludes1:** dementia with Parkinsonism (G31.83)
G21  Secondary parkinsonism
  **Excludes1:** dementia with Parkinsonism (G31.83)
    Huntington's disease (G10)
    Shy-Drager syndrome (G90.3)
    syphilitic Parkinsonism (A52.19)
  G21.0  Malignant neuroleptic syndrome
  G21.1  Other drug-induced secondary parkinsonism
    G21.11  Neuroleptic induced parkinsonism
      **Use additional** code for adverse effect, if applicable, to identify drug (T43.3X5,
      T43.4X5, T43.505,T43.595)
      **Excludes1:** malignant neuroleptic syndrome (G21.0)
    G21.19  Other drug induced secondary parkinsonism
Use additional code for adverse effect, if applicable, to identify drug (T36-T50 with fifth or sixth character 5)

G21.2 Secondary parkinsonism due to other external agents
Code first (T51-T65) to identify external agent
G21.3 Postencephalitic parkinsonism
G21.4 Vascular parkinsonism
G21.8 Other secondary parkinsonism
G21.9 Secondary parkinsonism, unspecified

Other degenerative diseases of the nervous system (G30-G32)

G30 Alzheimer's disease
Includes: Alzheimer's dementia senile and presenile forms
Use additional code to identify:
delirium, if applicable (F05)
dementia with behavioral disturbance (F02.81)
dementia without behavioral disturbance (F02.80)
Excludes1: senile degeneration of brain NEC (G31.1)
senile dementia NOS (F03)
senility NOS (R41.81)
G30.0 Alzheimer's disease with early onset
G30.1 Alzheimer's disease with late onset
G30.8 Other Alzheimer's disease
G30.9 Alzheimer's disease, unspecified

G31 Other degenerative diseases of nervous system, not elsewhere classified
Use additional code to identify: dementia with behavioral disturbance (F02.81)
dementia without behavioral disturbance (F02.80)
Excludes2: Reye's syndrome (G93.7)
G31.0 Frontotemporal dementia
G31.01 Pick's disease
Circumscribed brain atrophy
Progressive isolated aphasia
G31.09 Other frontotemporal dementia
Frontal dementia
G31.1 Senile degeneration of brain, not elsewhere classified
Excludes1: Alzheimer's disease (G30.-)
senility NOS (R41.81)
G31.8 Other specified degenerative diseases of nervous system
G31.84 Mild cognitive impairment, so stated
Excludes1: age related cognitive decline (R41.81)
altered mental status (R41.82)
cerebral degeneration (G31.9)
change in mental status (R41.82)
cognitive deficits following (sequelae of) cerebral hemorrhage or infarction (I69.01, I69.11, I69.21, I69.31, I69.81, I69.91)
cognitive impairment due to intracranial or head injury (S06.-)
dementia (F01.-, F02.-, F03)
mild memory disturbance (F06.8)
neurologic neglect syndrome (R41.4)
personality change, nonpsychotic (F68.8)

Demyelinating diseases of the central nervous system (G35-G37)

G35 Multiple sclerosis
  Disseminated multiple sclerosis
  Generalized multiple sclerosis
  Multiple sclerosis NOS
  Multiple sclerosis of brain stem
  Multiple sclerosis of cord

Episodic and paroxysmal disorders (G40-G47)

G40 Epilepsy and recurrent seizures
  G40.8 Other epilepsy and recurrent seizures
    Epilepsies and epileptic syndromes undetermined as to whether they are focal or generalized
    Landau-Kleffner syndrome
  G40.80 Other epilepsy
    G40.801 Other epilepsy, not intractable, with status epilepticus
    G40.802 Other epilepsy, not intractable, without status epilepticus
    G40.803 Other epilepsy, intractable, with status epilepticus
    G40.804 Other epilepsy, intractable, without status epilepticus

◊ G45 Transient cerebral ischemic attacks and related syndromes
◊ G46 Vascular syndromes of brain in cerebrovascular diseases

Nerve, nerve root and plexus disorders (G50-G59)

G51 Facial nerve disorders
  Includes: disorders of 7th cranial nerve
  G51.0 Bell's palsy
    Facial palsy
  G51.1 Geniculate ganglionitis
    Excludes1: postherpetic geniculate ganglionitis (B02.21)
  G51.2 Melkersson's syndrome
    Melkerson-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

Cerebral palsy and other paralytic syndromes (G80-G83)

G80 Cerebral palsy
  Excludes1: hereditary spastic paraplegia (G11.4)
  G80.0 Spastic quadriplegic cerebral palsy
    Congenital spastic paralysis (cerebral)
  G80.1 Spastic diplegic cerebral palsy
    Spastic cerebral palsy NOS
  G80.2 Spastic hemiplegic cerebral palsy
G80.3  Athetoid cerebral palsy
   Double athetosis (syndrome)
   Dyskinetic cerebral palsy
   Dystonic cerebral palsy
   Vogt disease
G80.4  Ataxic cerebral palsy
G80.8  Other cerebral palsy
   Mixed cerebral palsy syndromes
G80.9  Cerebral palsy, unspecified
   Cerebral palsy NOS
◊ G81  Hemiplegia and hemiparesis
   Note: 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.
Excludes1: congenital cerebral palsy (G80.-)
   hemiplegia and hemiparesis due to sequel of cerebrovascular disease (I69.05-, I69.15-, I69.25-, I69.35-, I69.45-, I69.85-, I69.95-)

Other disorders of the nervous system (G89-G99)
◊ G91  Hydrocephalus
◊ G93  Other disorders of brain
G96  Other disorders of central nervous system
   G96.0  Cerebrospinal fluid leak
   Excludes1: cerebrospinal fluid leak from spinal puncture (G97.0)

Ch. 9 Diseases of the circulatory system (I00-I99)

Pulmonary heart disease and diseases of pulmonary circulation (I26-I28)

Other forms of heart disease (I30-I52)
◊ I46  Cardiac arrest

Cerebrovascular diseases (I60-I69)
◊ I63  Cerebral infarction
I69  Sequelae of cerebrovascular disease
   Note: Category I69 is to be used to indicate conditions in I60-I67 as the cause of sequelae. The
   'sequelae' include conditions specified as such or as residuals which may occur at any time after
   the onset of the causal condition.
I69.0  Sequelae of nontraumatic subarachnoid hemorrhage
   I69.00  Unspecified sequelae of nontraumatic subarachnoid hemorrhage
   I69.01  Cognitive deficits following nontraumatic subarachnoid hemorrhage
   I69.02  Speech and language deficits following nontraumatic subarachnoid hemorrhage
      ✓ I69.020  Aphasia following nontraumatic subarachnoid hemorrhage
      ✓ I69.021  Dysphasia following nontraumatic subarachnoid hemorrhage
      ✓ I69.022  Dysarthria following nontraumatic subarachnoid hemorrhage
      ✓ I69.023  Fluency disorder following nontraumatic subarachnoid hemorrhage
      ✓ I69.028  Other speech and language deficits following nontraumatic subarachnoid hemorrhage

✓ Code typically used by SLPs  ◊ Additional digits not listed here
I69.09 Other sequelae of nontraumatic subarachnoid hemorrhage
  ✓ I69.090 Apraxia following nontraumatic subarachnoid hemorrhage
  ✓ I69.091 Dysphagia following nontraumatic subarachnoid hemorrhage
  **Use additional** code to identify the type of dysphagia, if known (R13.1-)
  I69.092 Facial weakness following nontraumatic subarachnoid hemorrhage
  Facial droop following nontraumatic subarachnoid hemorrhage
  I69.093 Ataxia following nontraumatic subarachnoid hemorrhage
  I69.098 Other sequelae following nontraumatic subarachnoid hemorrhage
  Alterations of sensation following nontraumatic subarachnoid hemorrhage
  Disturbance of vision following nontraumatic subarachnoid hemorrhage
  **Use additional** code to identify the sequelae

I69.1 Sequelae of nontraumatic intracerebral hemorrhage
  I69.10 Unspecified sequelae of nontraumatic intracerebral hemorrhage
  I69.11 Cognitive deficits following nontraumatic intracerebral hemorrhage
  I69.12 Speech and language deficits following nontraumatic intracerebral hemorrhage
    ✓ I69.120 Aphasia following nontraumatic intracerebral hemorrhage
    ✓ I69.121 Dysphasia following nontraumatic intracerebral hemorrhage
    ✓ I69.122 Dysarthria following nontraumatic intracerebral hemorrhage
    ✓ I69.123 Fluency disorder following nontraumatic intracerebral hemorrhage
    Stuttering following nontraumatic subarachnoid hemorrhage
    ✓ I69.128 Other speech and language deficits following nontraumatic intracerebral hemorrhage

I69.19 Other sequelae of nontraumatic intracerebral hemorrhage
  ✓ I69.190 Apraxia following nontraumatic intracerebral hemorrhage
  ✓ I69.191 Dysphagia following nontraumatic intracerebral hemorrhage
  **Use additional** code to identify the type of dysphagia, if known (R13.1-)
  I69.192 Facial weakness following nontraumatic intracerebral hemorrhage
  Facial droop following nontraumatic intracerebral hemorrhage
  I69.193 Ataxia following nontraumatic intracerebral hemorrhage
  I69.198 Other sequelae of nontraumatic intracerebral hemorrhage
  Alteration of sensations following nontraumatic intracerebral hemorrhage
  Disturbance of vision following nontraumatic intracerebral hemorrhage
  **Use additional** code to identify the sequelae

I69.2 Sequelae of other nontraumatic intracranial hemorrhage
  I69.20 Unspecified sequelae of other nontraumatic intracranial hemorrhage
  ✓ I69.21 Cognitive deficits following other nontraumatic intracranial hemorrhage
  ✓ I69.22 Speech and language deficits following other nontraumatic intracranial hemorrhage
    ✓ I69.220 Aphasia following other nontraumatic intracranial hemorrhage
    ✓ I69.221 Dysphasia following other nontraumatic intracranial hemorrhage
    ✓ I69.222 Dysarthria following other nontraumatic intracranial hemorrhage
    ✓ I69.223 Fluency disorder following other nontraumatic intracranial hemorrhage

✓ Code typically used by SLPs  ◊ Additional digits not listed here
hemorrhage
Stuttering following nontraumatic subarachnoid hemorrhage

I69.228 Other speech and language deficits following other nontraumatic intracranial hemorrhage

I69.29 Other sequelae of other nontraumatic intracranial hemorrhage

I69.290 Apraxia following other nontraumatic intracranial hemorrhage

I69.291 Dysphagia following other nontraumatic intracranial hemorrhage
Use additional code to identify the type of dysphagia, if known (R13.1-)

I69.292 Facial weakness following other nontraumatic intracranial hemorrhage
Facial droop following other nontraumatic intracranial hemorrhage

I69.293 Ataxia following other nontraumatic intracranial hemorrhage

I69.298 Other sequelae of other nontraumatic intracranial hemorrhage
Alteration of sensation following other nontraumatic intracranial hemorrhage
Disturbance of vision following other nontraumatic intracranial hemorrhage

Use additional code to identify the sequelae

I69.3 Sequelae of cerebral infarction
Sequelae of stroke NOS

I69.30 Unspecified sequelae of cerebral infarction

I69.31 Cognitive deficits following cerebral infarction

I69.32 Speech and language deficits following cerebral infarction

I69.320 Aphasia following cerebral infarction

I69.321 Dysphasia following cerebral infarction

I69.322 Dysarthria following cerebral infarction

I69.323 Fluency disorder following cerebral infarction
Stuttering following nontraumatic subarachnoid hemorrhage

I69.328 Other speech and language deficits following cerebral infarction

I69.39 Other sequelae of cerebral infarction

I69.390 Apraxia following cerebral infarction

I69.391 Dysphagia following cerebral infarction
Use additional code to identify the type of dysphagia, if known (R13.1-)

I69.392 Facial weakness following cerebral infarction
Facial droop following cerebral infarction

I69.393 Ataxia following cerebral infarction

I69.398 Other sequelae of cerebral infarction
Alteration of sensation following cerebral infarction
Disturbance of vision following cerebral infarction

Use additional code to identify the sequelae

I69.8 Sequelae of other cerebrovascular diseases

Excludes1: sequelae of traumatic intracranial injury (S06.-)

I69.80 Unspecified sequelae of other cerebrovascular disease

I69.81 Cognitive deficits following other cerebrovascular disease

I69.82 Speech and language deficits following other cerebrovascular disease

I69.820 Aphasia following other cerebrovascular disease

✓ Code typically used by SLPs  ◊ Additional digits not listed here
I69.821 Dysphasia following other cerebrovascular disease
I69.822 Dysarthria following other cerebrovascular disease
I69.823 Fluency disorder following other cerebrovascular disease
Stuttering following nontraumatic subarachnoid hemorrhage
I69.828 Other speech and language deficits following other cerebrovascular disease

I69.89 Other sequelae of other cerebrovascular disease
I69.890 Apraxia following other cerebrovascular disease
I69.891 Dysphagia following other cerebrovascular disease
Use additional code to identify the type of dysphagia, if known (R13.1-)
I69.892 Facial weakness following other cerebrovascular disease
Facial droop following other cerebrovascular disease
I69.893 Ataxia following other cerebrovascular disease
I69.898 Other sequelae of other cerebrovascular disease
Alteration of sensation following other cerebrovascular disease
Disturbance of vision following other cerebrovascular disease

Use additional code to identify the sequelae

I69.9 Sequelae of unspecified cerebrovascular diseases
Excludes1: sequelae of stroke (I63.3)
sequelae of traumatic intracranial injury (S06.-)
I69.90 Unspecified sequelae of unspecified cerebrovascular disease
I69.91 Cognitive deficits following unspecified cerebrovascular disease
I69.92 Speech and language deficits following unspecified cerebrovascular disease
I69.920 Aphasia following unspecified cerebrovascular disease
I69.921 Dysphasia following unspecified cerebrovascular disease
I69.922 Dysarthria following unspecified cerebrovascular disease
I69.923 Fluency disorder following unspecified cerebrovascular disease
Stuttering following nontraumatic subarachnoid hemorrhage
I69.928 Other speech and language deficits following unspecified cerebrovascular disease

I69.99 Other sequelae of unspecified cerebrovascular disease
I69.990 Apraxia following unspecified cerebrovascular disease
I69.991 Dysphagia following unspecified cerebrovascular disease
Use additional code to identify the type of dysphagia, if known (R13.1-)
I69.992 Facial weakness following unspecified cerebrovascular disease
Facial droop following unspecified cerebrovascular disease
I69.993 Ataxia following unspecified cerebrovascular disease
I69.998 Other sequelae following unspecified cerebrovascular disease
Alteration in sensation following unspecified cerebrovascular disease
Disturbance of vision following unspecified cerebrovascular disease
Use additional code to identify the sequelae

Ch. 10 Diseases of the respiratory system (J00-J99)

Acute upper respiratory infections (J00-J06)
J02 Acute pharyngitis
◊ J03 Acute tonsillitis
◊ J04 Acute laryngitis and tracheitis
◊ J05 Acute obstructive laryngitis [croup] and epiglottitis

**Other diseases of upper respiratory tract (J30-J39)**

◊ J31 Chronic rhinitis, nasopharyngitis and pharyngitis

**Use additional** code to identify:
- exposure to environmental tobacco smoke (Z77.22)
- exposure to tobacco smoke in the perinatal period (P96.81)
- history of tobacco use (Z87.891)
- occupational exposure to environmental tobacco smoke (Z57.31)
- tobacco dependence (F17.-)
- tobacco use (Z72.0)

J35.1 Hypertrophy of tonsils
- Enlargement of tonsils
  **Excludes1:** hypertrophy of tonsils with tonsillitis (J35.0-)

J35.2 Hypertrophy of adenoids
- Enlargement of adenoids
  **Excludes1:** hypertrophy of adenoids with adenoiditis (J35.0-)

J35.3 Hypertrophy of tonsils with hypertrophy of adenoids
  **Excludes1:** hypertrophy of tonsils and adenoids with tonsillitis and adenoiditis (J35.03)

J37 Chronic laryngitis and laryngotracheitis

J37.0 Chronic laryngitis
- Catarrhal laryngitis
- Hypertrophic laryngitis
- Sicca laryngitis
  **Excludes2:** acute laryngitis (J04.0)
  obstructive (acute) laryngitis (J05.0)

J38 Diseases of vocal cords and larynx, not elsewhere classified

J38.0 Paralysis of vocal cords and larynx
- Laryngoplegia
- Paralysis of glottis
  J38.00 Paralysis of vocal cords and larynx, unspecified
  J38.01 Paralysis of vocal cords and larynx, unilateral
  J38.02 Paralysis of vocal cords and larynx, bilateral

J38.1 Polyp of vocal cord and larynx
  **Excludes1:** adenomatous polyps (D14.1)

J38.2 Nodules of vocal cords
- Chorditis (fibrinous)(nodosa)(tuberosa)
- Singer's nodes
- Teacher's nodes

J38.3 Other diseases of vocal cords
- Abscess of vocal cords
- Cellulitis of vocal cords
- Granuloma of vocal cords
- Leukokeratosis of vocal cords
- Leukoplakia of vocal cords
J38.4 Edema of larynx  
   Edema (of) glottis  
   Subglottic edema  
   Supraglottic edema  
   **Excludes1:** acute obstructive laryngitis [croup] (J05.0)  
   edematous laryngitis (J04.0)

J38.5 Laryngeal spasm  
   Laryngismus (stridulus)

J38.6 Stenosis of larynx

J38.7 Other diseases of larynx  
   Abscess of larynx  
   Cellulitis of larynx  
   Disease of larynx NOS  
   Necrosis of larynx  
   Pachyderma of larynx  
   Perichondritis of larynx  
   Ulcer of larynx

J38.8 Upper respiratory tract hypersensitivity reaction, site unspecified  
   **Excludes1:** hypersensitivity reaction of upper respiratory tract, such as:  
   extrinsic allergic alveolitis (J67.9)  
   pneumoconiosis (J60-J67.9)

J38.9 Other specified diseases of upper respiratory tract

J39.0 Disease of upper respiratory tract, unspecified

**Lung diseases due to external agents (J60-J70)**

J69 Pneumonitis due to solids and liquids  
   **Excludes1:** neonatal aspiration syndromes (P24.-)  
   postprocedural pneumonitis (J95.4)

J69.0 Pneumonitis due to inhalation of food and vomit  
   Aspiration pneumonia NOS  
   Aspiration pneumonia (due to) food (regurgitated)  
   Aspiration pneumonia (due to) gastric secretions  
   Aspiration pneumonia (due to) milk  
   Aspiration pneumonia (due to) vomit  
   **Code also** any associated foreign body in respiratory tract (T17.-)  
   **Excludes1:** chemical pneumonitis due to anesthesia (J95.4)  
   obstetric aspiration pneumonitis (O74.0)

*Intraoperative and postprocedural complications and disorders of respiratory system, not elsewhere classified (J95)*

J95 Intraoperative and postprocedural complications and disorders of respiratory system, not elsewhere classified  
   **Excludes2:** aspiration pneumonia (J69.-)  
   emphysema (subcutaneous) resulting from a procedure (T81.82)  
   hypostatic pneumonia (J18.2)  
   pulmonary manifestations due to radiation (J70.0-J70.1)

J95.0 Tracheostomy complications  
   J95.00 Unspecified tracheostomy complication

- Code typically used by SLPs  
- Additional digits not listed here
J95.01  Hemorrhage from tracheostomy stoma
J95.02  Infection of tracheostomy stoma
Use additional code to identify type of infection, such as:
cellulitis of neck (L03.8)
sepsis (A40, A41.-)
J95.03  Malfunction of tracheostomy stoma
Mechanical complication of tracheostomy stoma
Obstruction of tracheostomy airway
Tracheal stenosis due to tracheostomy
J95.04  Tracheo-esophageal fistula following tracheostomy
J95.09  Other tracheostomy complication

Ch. 11 Diseases of the digestive system (K00-K95)

Diseases of oral cavity and salivary glands (K00-K14)

◊ K00  Disorders of tooth development and eruption
K08  Other disorders of teeth and supporting structures
    Excludes2: dentofacial anomalies [including malocclusion] (M26.-)
    disorders of jaw (M27.-)
◊ K08.2  Atrophy of edentulous alveolar ridge

K13  Other diseases of lip and oral mucosa
K13.7  Other and unspecified lesions of oral mucosa
K13.70  Unspecified lesions of oral mucosa
K13.79  Other lesions of oral mucosa
    Focal oral mucinosis

K14  Diseases of tongue
K14.0  Glossitis
    Abscess of tongue
    Ulceration (traumatic) of tongue
    Excludes1: atrophic glossitis (K14.4)
K14.4  Atrophy of tongue papillae
    Atrophic glossitis
K14.5  Plicated tongue
    Fissured tongue
    Furrowed tongue
    Scrotal tongue
    Excludes1: fissured tongue, congenital (Q38.3)
K14.8  Other diseases of tongue
    Atrophy of tongue
    Crenated tongue
    Enlargement of tongue
    Glossocele
    Glossoptosis
    Hypertrophy of tongue

K21  Gastro-esophageal reflux disease
    Excludes1: newborn esophageal reflux (P78.83)
K21.0  Gastro-esophageal reflux disease with esophagitis
    Reflux esophagitis
K21.9  Gastro-esophageal reflux disease without esophagitis
          Esophageal reflux NOS

**Ch. 13  Diseases of the musculoskeletal system and connective tissue (M00-M99)**

*Dentofacial anomalies [including malocclusion] and other disorders of jaw (M26-M27)*

M26  Dentofacial anomalies [including malocclusion]

M26.0  Major anomalies of jaw size

*Excludes1:* acromegaly (E22.0)

- Robin's syndrome (Q87.0)
- Unspecified anomaly of jaw size
- Maxillary hyperplasia
- Maxillary hypoplasia
- Mandibular hyperplasia
- Mandibular hypoplasia
- Macrogenia
- Microgenia
- Excessive tuberosity of jaw
  - Entire maxillary tuberosity

M26.09  Other specified anomalies of jaw size

M26.1  Anomalies of jaw-cranial base relationship

M26.10  Unspecified anomaly of jaw-cranial base relationship

- Maxillary asymmetry
- Other jaw asymmetry
- Other specified anomalies of jaw-cranial base relationship

M26.2  Anomalies of dental arch relationship

M26.20  Unspecified anomaly of dental arch relationship

M26.21  Malocclusion, Angle's class

- Malocclusion, Angle's class I
  - Neutro-occlusion
- Malocclusion, Angle's class II
  - Disto-occlusion Division I
  - Disto-occlusion Division II
- Malocclusion, Angle's class III
  - Mesio-occlusion
- Malocclusion, Angle's class, unspecified

M26.22  Open occlusal relationship

- Open anterior occlusal relationship
  - Anterior openbite
- Open posterior occlusal relationship
  - Posterior openbite

M26.23  Excessive horizontal overlap
          Excessive horizontal overjet

M26.24  Reverse articulation
          Crossbite (anterior) (posterior)

M26.25  Anomalies of interarch distance

M26.29  Other anomalies of dental arch relationship

Midline deviation of dental arch
Overbite (excessive) deep
Overbite (excessive) horizontal
Overbite (excessive) vertical
Posterior lingual occlusion of mandibular teeth

- M26.3 Anomalies of tooth position of fully erupted tooth or teeth

- M26.4 Malocclusion, unspecified

- M26.5 Dentofacial functional abnormalities
  - **Exclude1**: bruxism (F45.8)
  - teeth-grinding NOS (F45.8)
  - M26.50 Dentofacial functional abnormalities, unspecified
  - M26.51 Abnormal jaw closure
  - M26.52 Limited mandibular range of motion
  - M26.53 Deviation in opening and closing of the mandible
  - M26.54 Insufficient anterior guidance
    - Insufficient anterior occlusal guidance
  - M26.55 Centric occlusion maximum intercuspation discrepancy
    - **Exclude1**: centric occlusion NOS (M26.59)
  - M26.56 Non-working side interference
    - Balancing side interference
  - M26.57 Lack of posterior occlusal support
  - M26.59 Other dentofacial functional abnormalities
    - Centric occlusion (of teeth) NOS
    - Malocclusion due to abnormal swallowing
    - Malocclusion due to mouth breathing
    - Malocclusion due to tongue, lip or finger habits

- M26.6 Temporomandibular joint disorders
  - **Exclude2**: current temporomandibular joint dislocation (S03.0)
    - current temporomandibular joint sprain (S03.4)
  - M26.60 Temporomandibular joint disorder, unspecified
  - M26.61 Adhesions and ankylosis of temporomandibular joint
  - M26.62 Arthralgia of temporomandibular joint
  - M26.63 Articular disc disorder of temporomandibular joint
  - M26.69 Other specified disorders of temporomandibular joint

- M26.7 Dental alveolar anomalies
  - M26.70 Unspecified alveolar anomaly
  - M26.71 Alveolar maxillary hyperplasia
  - M26.72 Alveolar mandibular hyperplasia
  - M26.73 Alveolar maxillary hypoplasia
  - M26.74 Alveolar mandibular hypoplasia
  - M26.79 Other specified alveolar anomalies

- M26.8 Other dentofacial anomalies
  - M26.81 Anterior soft tissue impingement
    - Anterior soft tissue impingement on teeth
  - M26.82 Posterior soft tissue impingement
    - Posterior soft tissue impingement on teeth
  - M26.89 Other dentofacial anomalies

- M26.9 Dentofacial anomaly, unspecified

- Code typically used by SLPs
- Additional digits not listed here
Ch. 16  Certain conditions originating in the perinatal period (P00-P96)

*Other disorders originating in the perinatal period (P90-P96)*

- **P92** Feeding problems of newborn
  - **Excludes1:** feeding problems in child over 28 days old (R63.3)
- **P92.2** Slow feeding of newborn
- **P92.6** Failure to thrive in newborn
  - **Excludes1:** failure to thrive in child over 28 days old (R62.51)
- **P92.8** Other feeding problems of newborn
- **P92.9** Feeding problem of newborn, unspecified

Ch. 17  Congenital malformations, deformations and chromosomal abnormalities (Q00-Q99)

*Congenital malformations of the nervous system (Q00-Q07)*

- **Q02** Microcephaly
- **Q03** Congenital hydrocephalus
- **Q04** Other congenital malformations of brain
  - **Q04.3** Other reduction deformities of brain
    - Absence of part of brain
    - Agenesis of part of brain
    - Agyria
    - Aplasia of part of brain
    - Hydranencephaly
    - Hypoplasia of part of brain
    - Lissencephaly
    - Microgyria
    - Pachygyria
  - **Excludes1:** congenital malformations of corpus callosum (Q04.0)
- **Q05** Spina bifida

*Congenital malformations of eye, ear, face and neck (Q10-Q18)*

- **Q16** Congenital malformations of ear causing impairment of hearing
  - **Excludes1:** congenital deafness (H90.-)
  - **Q16.0** Congenital absence of (ear) auricle
  - **Q16.1** Congenital absence, atresia and stricture of auditory canal (external)
    - Congenital atresia or stricture of osseous meatus
  - **Q16.2** Absence of eustachian tube
  - **Q16.3** Congenital malformation of ear ossicles
    - Congenital fusion of ear ossicles
  - **Q16.4** Other congenital malformations of middle ear
    - Congenital malformation of middle ear NOS
  - **Q16.5** Congenital malformation of inner ear
    - Congenital anomaly of membranous labyrinth
    - Congenital anomaly of organ of Corti
  - **Q16.9** Congenital malformation of ear causing impairment of hearing, unspecified
    - Congenital absence of ear NOS
  - **Q17** Other congenital malformations of ear
    - **Excludes1:** congenital malformations of ear with impairment of hearing (Q16.0-Q16.9)
    - Preauricular sinus (Q18.1)

- ✔ Code typically used by SLPs
- ◊ Additional digits not listed here
Q17.0  Accessory auricle
        Accessory tragus
        Polyotia
        Preauricular appendage or tag
        Supernumerary ear
        Supernumerary lobule

Q17.1  Macrotia
Q17.2  Microtia
Q17.3  Other misshapen ear
        Pointed ear
Q17.4  Misplaced ear
        Low-set ears
        **Excludes1**: cervical auricle (Q18.2)
Q17.5  Prominent ear
        Bat ear
Q17.8  Other specified congenital malformations of ear
        Congenital absence of lobe of ear
Q17.9  Congenital malformation of ear, unspecified
        Congenital anomaly of ear NOS

**Congenital malformations of the respiratory system (Q30-Q34)**

Q31  Congenital malformations of larynx
        **Excludes1**: congenital laryngeal stridor NOS (P28.89)
Q31.0  Web of larynx
        Glottic web of larynx
        Subglottic web of larynx
        Web of larynx NOS
Q31.1  Congenital subglottic stenosis
Q31.2  Laryngeal hypoplasia
Q31.3  Laryngocele
Q31.5  Congenital laryngomalacia
Q31.8  Other congenital malformations of larynx
        Absence of larynx
        Agenesis of larynx
        Atresia of larynx
        Congenital cleft thyroid cartilage
        Congenital fissure of epiglottis
        Congenital stenosis of larynx NEC
        Posterior cleft of cricoid cartilage
Q31.9  Congenital malformation of larynx, unspecified

**Cleft lip and cleft palate (Q35-Q37)**

**Use additional** code to identify associated malformation of the nose (Q30.2)
**Excludes1**: Robin's syndrome (Q87.0)

Q35  Cleft palate
        **Includes**: fissure of palate
        palatoschisis
        **Excludes1**: cleft palate with cleft lip (Q37.-)
Q35.1 Cleft hard palate
Q35.3 Cleft soft palate
Q35.5 Cleft hard palate with cleft soft palate
Q35.7 Cleft uvula
Q35.9 Cleft palate, unspecified
  Cleft palate NOS

Q36 Cleft lip
  Includes: cheilosis
  congenital fissure of lip
  harelip
  labium leporinum
  Excludes1: cleft lip with cleft palate (Q37.-)
Q36.0 Cleft lip, bilateral
Q36.1 Cleft lip, median
Q36.9 Cleft lip, unilateral
  Cleft lip NOS

Q37 Cleft palate with cleft lip
  Includes: cheilopalatoschisis
Q37.0 Cleft hard palate with bilateral cleft lip
Q37.1 Cleft hard palate with unilateral cleft lip
  Cleft hard palate with cleft lip NOS
Q37.2 Cleft soft palate with bilateral cleft lip
Q37.3 Cleft soft palate with unilateral cleft lip
  Cleft soft palate with cleft lip NOS
Q37.4 Cleft hard and soft palate with bilateral cleft lip
Q37.5 Cleft hard and soft palate with unilateral cleft lip
  Cleft hard and soft palate with cleft lip NOS
Q37.8 Unspecified cleft palate with bilateral cleft lip
Q37.9 Unspecified cleft palate with unilateral cleft lip
  Cleft palate with cleft lip NOS

Other congenital malformations of the digestive system (Q38-Q45)

Q38 Other congenital malformations of tongue, mouth and pharynx
  Excludes1: dentofacial anomalies (M26.-)
  macrostomia (Q18.4)
  microstomia (Q18.5)
Q38.0 Congenital malformations of lips, not elsewhere classified
  Congenital fistula of lip
  Congenital malformation of lip NOS
  Van der Woude's syndrome
  Excludes1: cleft lip (Q36.-)
    cleft lip with cleft palate (Q37.-)
    macrocheilia (Q18.6)
    microcheilia (Q18.7)
Q38.1 Ankyloglossia
  Tongue tie
Q38.2 Macroglossia
  Congenital hypertrophy of tongue
Q38.3 Other congenital malformations of tongue
- Aglossia
- Bifid tongue
- Congenital adhesion of tongue
- Congenital fissure of tongue
- Congenital malformation of tongue NOS
- Double tongue
- Hypoglossia
- Hypoplasia of tongue
- Microglossia

Q38.4 Congenital malformations of salivary glands and ducts
- Atresia of salivary glands and ducts
- Congenital absence of salivary glands and ducts
- Congenital accessory salivary glands and ducts
- Congenital fistula of salivary gland

Q38.5 Congenital malformations of palate, not elsewhere classified
- Congenital absence of uvula
- Congenital malformation of palate NOS
- Congenital high arched palate

Excludes1: cleft palate (Q35.-)
- cleft palate with cleft lip (Q37.-)

Q38.6 Other congenital malformations of mouth
- Congenital malformation of mouth NOS

Q38.7 Congenital pharyngeal pouch
- Congenital diverticulum of pharynx

Excludes1: pharyngeal pouch syndrome (D82.1)

Q38.8 Other congenital malformations of pharynx
- Congenital malformation of pharynx NOS
- Imperforate pharynx

Congenital malformations and deformations of the musculoskeletal system (Q65-Q79)

Q67 Congenital musculoskeletal deformities of head, face, spine and chest

Q67.0 Congenital facial asymmetry

Q67.4 Other congenital deformities of skull, face and jaw
- Congenital depressions in skull
- Congenital hemifacial atrophy or hypertrophy
- Deviation of nasal septum, congenital
- Squashed or bent nose, congenital

Excludes1: dentofacial anomalies [including malocclusion] (M26-)
- Syphilitic saddle nose (A50.5)

Chromosomal abnormalities, not elsewhere classified (Q90-Q99)

Q90 Down syndrome

Use additional code(s) to identify any associated physical conditions and degree of intellectual disabilities (F70-F79)

Q90.0 Trisomy 21, nonmosaicism (meiotic nondisjunction)

Q90.1 Trisomy 21, mosaicism (mitotic nondisjunction)

Q90.2 Trisomy 21, translocation

Q38.3 is an option for a diagnosis of tongue thrust.
Q90.9  Down syndrome, unspecified
       Trisomy 21 NOS

Q91  Trisomy 18 and Trisomy 13
   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

Q93  Monosomies and deletions from the autosomes, not elsewhere classified
   Q93.3  Deletion of short arm of chromosome 4
          Wolff-Hirschhorn syndrome
   Q93.4  Deletion of short arm of chromosome 5
          Cri-du-chat syndrome
   Q93.8  Other deletions from the autosomes
   Q93.81  Velo-cardio-facial syndrome
          Deletion 22q11.2

Q98  Other sex chromosome abnormalities, male phenotype, not elsewhere classified
   Q98.0  Klinefelter syndrome karyotype 47, XXY
   Q98.1  Klinefelter syndrome, male with more than two X chromosomes

Ch. 18  Symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified (R00-R99)

Symptoms and signs involving the digestive system and abdomen (R10-R19)

R12  Heartburn
   Excludes1: dyspepsia NOS (R10.13)
              functional dyspepsia (K30)

R13  Aphagia and dysphagia
   ✓ R13.0  Aphagia
          Inability to swallow
          Excludes1: psychogenic aphagia (F50.9)
   ✓ R13.1  Dysphagia
          Code first, if applicable, dysphagia following cerebrovascular disease (I69. with final characters -91)
          Excludes1: psychogenic dysphagia (F45.8)
       ✓ R13.10  Dysphagia, unspecified
               Difficulty in swallowing NOS
       ✓ R13.11  Dysphagia, oral phase
       ✓ R13.12  Dysphagia, oropharyngeal phase
       ✓ R13.13  Dysphagia, pharyngeal phase
       ✓ R13.14  Dysphagia, pharyngoesophageal phase
       ✓ R13.19  Other dysphagia
              Cervical dysphagia
              Neurogenic dysphagia

Symptoms and signs involving the nervous and musculoskeletal systems (R25-R29)
R27 Other lack of coordination
   **Excludes1**: ataxic gait (R26.0)
      hereditary ataxia (G11-)
      vertigo NOS (R42)
R27.0 Ataxia, unspecified
   **Excludes1**: ataxia following cerebrovascular disease (I69. with final characters -93)
R27.8 Other lack of coordination
R27.9 Unspecified lack of coordination

R29 Other symptoms and signs involving the nervous and musculoskeletal systems
R29.8 Other symptoms and signs involving the nervous and musculoskeletal systems
R29.81 Other symptoms and signs involving the nervous system
   R29.810 Facial weakness
      Facial droop
      **Excludes1**: Bell's palsy (G51.0)
      facial weakness following cerebrovascular disease (I69. with final characters-92)
   R29.818 Other symptoms and signs involving the nervous system

*Symptoms and signs involving cognition, perception, emotional state and behavior (R40-R46)*

**Excludes1**: symptoms and signs constituting part of a pattern of mental disorder (F01-F99)

R41 Other symptoms and signs involving cognitive functions and awareness
   **Excludes1**: dissociative [conversion] disorders (F44-)
      mild cognitive impairment, so stated (G31.84)
R41.0 Disorientation, unspecified
      Confusion NOS
      Delirium NOS
R41.1 Anterograde amnesia
R41.2 Retrograde amnesia
R41.3 Other amnesia
      Amnesia NOS
      Memory loss NOS
      **Excludes1**: amnestic disorder due to known physiologic condition (F04)
      amnestic syndrome due to psychoactive substance use (F10-F19 with 5th character .6)
      transient global amnesia (G45.4)
R41.4 Neurologic neglect syndrome
      Asomatognosia
      Hemi-akinesia
      Hemi-inattention
      Hemispatial neglect
      Left-sided neglect
      Sensory neglect
      Visuospatial neglect
      **Excludes1**: visuospatial deficit (R41.842)
R41.8 Other symptoms and signs involving cognitive functions and awareness
R41.81 Age-related cognitive decline
      Senility NOS
R41.82 Altered mental status, unspecified
Change in mental status NOS

**Excludes**: altered level of consciousness (R40.-)
altered mental status due to known condition - code to condition delirium NOS (R41.0)

R41.83 Borderline intellectual functioning
IQ level 71 to 84

**Excludes**: intellectual disabilities (F70-F79)

R41.84 Other specified cognitive deficit

- R41.840 Attention and concentration deficit
  **Excludes**: attention-deficit hyperactivity disorders (F90.-)

- R41.841 Cognitive communication deficit

- R41.842 Visuospatial deficit

- R41.843 Psychomotor deficit

- R41.844 Frontal lobe and executive function deficit

R41.89 Other symptoms and signs involving cognitive functions and awareness
Anosognosia

R41.9 Unspecified symptoms and signs involving cognitive functions and awareness

R44 Other symptoms and signs involving general sensations and perceptions

**Excludes**: alcoholic hallucinations (F1.5)

- hallucinations in drug psychosis (F11-F19 with .5)
- hallucinations in mood disorders with psychotic symptoms (F30.2, F31.5, F32.3, F33.3)
- hallucinations in schizophrenia, schizotypal and delusional disorders (F20-F29)

**Excludes**: disturbances of skin sensation (R20.-)

R44.0 Auditory hallucinations

**Symptoms and signs involving speech and voice (R47-R49)**

R47 Speech disturbances, not elsewhere classified

**Excludes**: autism (F84.0)

- cluttering (F80.81)
- specific developmental disorders of speech and language (F80.-)
- stuttering (F80.81)

R47.0 Dysphasia and aphasia

- R47.01 Aphasia
  **Excludes**: aphasia following cerebrovascular disease (l69. with final characters -20)
  progressive isolated aphasia (G31.01)

- R47.02 Dysphasia
  **Excludes**: dysphasia following cerebrovascular disease (l69. with final characters -21)

- R47.1 Dysarthria and anarthria
  **Excludes**: dysarthria following cerebrovascular disease (l69. with final characters -22)

R47.8 Other speech disturbances

**Excludes**: dysarthria following cerebrovascular disease (l69. with final characters -28)

- R47.81 Slurred speech

- R47.82 Fluency disorder in conditions classified elsewhere
  Stuttering in conditions classified elsewhere

**Code first**: underlying disease or condition, such as:

- Code typically used by SLPs
- Additional digits not listed here
Parkinson’s disease (G20)

**Excludes1:** adult onset fluency disorder (F98.5)
childhood onset fluency disorder (F80.81)
fluency disorder (stuttering) following cerebrovascular disease (I69. with final characters-23)

✔ R47.89 Other speech disturbances
R47.9 Unspecified speech disturbances

R48 Dyslexia and other symbolic dysfunctions, not elsewhere classified

**Excludes1:** specific developmental disorders of scholastic skills (F81.-)

✔ R48.0 Dyslexia and alexia
✔ R48.1 Agnosia
  Astereognosia (astereognosis)
  Autotopagnosia
  **Excludes1:** visual object agnosia H53.16

✔ R48.2 Apraxia
  **Excludes1:** apraxia following cerebrovascular disease (I69. with final characters -90)

R48.3 Visual agnosia
  Prosopagnosia
  Simultanagnosia (asimultagnosia)

✔ R48.8 Other symbolic dysfunctions
  Acalculia
  Agraphia

✔ R48.9 Unspecified symbolic dysfunctions

R49 Voice and resonance disorders

**Excludes1:** psychogenic voice and resonance disorders (F44.4)

✔ R49.0 Dysphonia
  Hoarseness

✔ R49.1 Aphonias
  Loss of voice

R49.2 Hypertautophonia and hyponautophonia
  ✔ R49.21 Hypertautophonia
  ✔ R49.22 Hyponautophonia

✔ R49.8 Other voice and resonance disorders

R49.9 Unspecified voice and resonance disorder
  Change in voice NOS
  Resonance disorder NOS

**General symptoms and signs (R50-R69)**

R62 Lack of expected normal physiological development in childhood and adults

**Excludes1:** delayed puberty (E30.0)
  gonadal dysgenesis (Q99.1)
  hypopituitarism (E23.0)

✔ R62.0 Delayed milestone in childhood
  Delayed attainment of expected physiological developmental stage
  Late talker
  Late walker

R62.5 Other and unspecified lack of expected normal physiological development in childhood

**Excludes1:** HIV disease resulting in failure to thrive (B20)
physical retardation due to malnutrition (E45)
R62.50 Unspecified lack of expected normal physiological development in childhood
Infantilism NOS
R62.51 Failure to thrive (child)
Failure to gain weight
Excludes1: failure to thrive in child under 28 days old (P92.6)
R63.3 Feeding difficulties
Feeding problem (elderly) (infant) NOS
Excludes1: feeding problems of newborn (P92.-)
infant feeding disorder of nonorganic origin (F98.2-)
R63.4 Abnormal weight loss

Abnormal findings on diagnostic imaging and in function studies, without diagnosis (R90-R94)
R94 Abnormal results of function studies
R94.0 Abnormal results of function studies of central nervous system
R94.01 Abnormal electroencephalogram [EEG]
R94.02 Abnormal brain scan
R94.09 Abnormal results of other function studies of central nervous system
R94.1 Abnormal results of function studies of peripheral nervous system and special senses
R94.12 Abnormal results of function studies of ear and other special senses
R94.120 Abnormal auditory function study
R94.121 Abnormal vestibular function study
R94.128 Abnormal results of other function studies of ear and other special senses

Ch. 19 Injury, poisoning and certain other consequences of external causes (S00-T88)

Note: Use secondary code(s) from Chapter 20, External causes of morbidity, to indicate cause of injury. Codes within the T section that include the external cause do not require an additional external cause code
Use additional code to identify any retained foreign body, if applicable (Z18.-)
Excludes1: birth trauma (P10-P15)
obstetric trauma (O70-O71)

Note: The chapter uses the S-section for coding different types of injuries related to single body regions and the T-section to cover injuries to unspecified body regions as well as poisoning and certain other consequences of external causes.

Injuries to the head (S00-S09)

Includes: injuries of ear
injuries of eye
injuries of face [any part]
injuries of gum
injuries of jaw
injuries of oral cavity
injuries of palate
injuries of periocular area
injuries of scalp
injuries of temporomandibular joint area
injuries of tongue
injuries of tooth

✓ Code typically used by SLPs ○ Additional digits not listed here
**Excludes2:** burns and corrosions (T20-T32)  
effects of foreign body in ear (T16)  
effects of foreign body in larynx (T17.3)  
effects of foreign body in mouth NOS (T18.0)  
effects of foreign body in nose (T17.0-T17.1)  
effects of foreign body in pharynx (T17.2)  
effects of foreign body on external eye (T15.-)  
frostbite (T33-T34)

S00  Superficial injury of head  
◊ S00.5  Superficial injury of lip and oral cavity
S01  Open wound of head  
◊ S01.5  Open wound of lip and oral cavity
◊ S02  Fracture of skull and facial bones
S06  Intracranial injury  
**Includes:** traumatic brain injury  
**Excludes1:** head injury NOS (S09.90)
◊ S06.0  Concussion
◊ S06.2  Diffuse traumatic brain injury
◊ S06.3  Focal traumatic brain injury
S12  Fracture of cervical vertebra and other parts of neck
S12.8  Fracture of other parts of neck  
The appropriate 7th character is to be added to code S12.8  
A - initial encounter  
D - subsequent encounter  
S - sequela
Hyoid bone  
Larynx  
Thyroid cartilage  
Trachea

**Injury, poisoning and certain other consequences of external causes (T07-T88)**

T17  Foreign body in respiratory tract  
The appropriate 7th character is to be added to each code from category T17  
A - initial encounter  
D - subsequent encounter  
S - sequela
T17.2  Foreign body in pharynx  
Foreign body in nasopharynx  
Foreign body in throat NOS  
T17.22  Food in pharynx  
Bones in pharynx  
Seeds in pharynx  
T17.220  Food in pharynx causing asphyxiation
T17.3  Foreign body in larynx  
T17.32  Food in larynx  
Bones in larynx  
Seeds in larynx

✔ Code typically used by SLPs  ✿ Additional digits not listed here
T17.320  Food in larynx causing asphyxiation
T17.4  Foreign body in trachea
T17.42  Food in trachea
Bones in trachea
Seeds in trachea
T17.420  Food in trachea causing asphyxiation
T18  Foreign body in alimentary tract
\textbf{Excludes2:} foreign body in pharynx (T17.2-)
\textbullet{} T18.1  Foreign body in esophagus

\textbf{Ch. 20  External causes of morbidity (V00-Y99)}

\textbf{Note:} This chapter permits the classification of environmental events and circumstances as the cause of injury, and other adverse effects. Where a code from this section is applicable, it is intended that it shall be used secondary to a code from another chapter of the Classification indicating the nature of the condition. Most often, the condition will be classifiable to Chapter 19, Injury, poisoning and certain other consequences of external causes (S00-T88). Other conditions that may be stated to be due to external causes are classified in Chapters 1 to 18. For these conditions, codes from Chapter 20 should be used to provide additional information as to the cause of the condition.

\textbf{Ch. 21  Factors Influencing Health Status and Contact with Health Services (Z00-Z99)}

\textbf{Note:} Z codes (formerly “V codes” in ICD-9-CM) represent reasons for encounters. A corresponding procedure code must accompany a Z code if a procedure is performed. Categories Z00-Z99 are provided for occasions when circumstances other than a disease, injury, or external cause classifiable to categories A00-Y89 are recorded as 'diagnoses' or 'problems'. This can arise in two main ways:

a. When a person who may or may not be sick encounters the health services for some specific purpose, such as to receive limited care or service for a current condition, to donate an organ or tissue, to receive prophylactic vaccination (immunization), or to discuss a problem which is in itself not a disease or injury.

b. When some circumstance or problem is present which influences the person's health status but is not in itself a current illness or injury.

\textit{Persons encountering health services for examinations (Z00-Z13)}

\textbf{Note:} Nonspecific abnormal findings disclosed at the time of these examinations are classified to categories R70-R94.

\textbf{Excludes1:} examinations related to pregnancy and reproduction (Z30-Z36, Z39.-)

Z01  Encounter for other special examination without complaint, suspected or reported diagnosis
\textbf{Includes:} routine examination of specific system
\textbf{Note:} Codes from category Z01 represent the reason for the encounter. A separate procedure code is required to identify any examinations or procedures performed

\textbf{Excludes1:} encounter for examination for administrative purposes (Z02.-)

encounter for examination for suspected conditions, proven not to exist (Z03.-)
encounter for laboratory and radiologic examinations as a component of general medical examinations(Z00.-)
encounter for laboratory, radiologic and imaging examinations for sign(s) and symptom(s) - code to the sign(s) or symptom(s)
**Excludes2:** screening examinations (Z11-Z13)

Z01.1  
Encounter for examination of ears and hearing  
Z01.10  
Encounter for examination of ears and hearing without abnormal findings  
Encounter for examination of ears and hearing NOS  
Z01.11  
Encounter for examination of ears and hearing with abnormal findings  
Z01.110  
Encounter for hearing examination following failed hearing screening  
Z01.118  
Encounter for examination of ears and hearing with other abnormal findings  
**Use additional** code to identify abnormal findings  
Z01.12  
Encounter for hearing conservation and treatment

**Z02**  
Encounter for administrative examination  
Z02.7  
Encounter for issue of medical certificate  
**Excludes1:** encounter for general medical examination (Z00-Z01, Z02.0-Z02.6, Z02.8-Z02.9)  
Z02.71  
Encounter for disability determination  
Encounter for issue of medical certificate of incapacity  
Encounter for issue of medical certificate of invalidity  
Z02.79  
Encounter for issue of other medical certificate

**Z02**  
Encounter for screening for other diseases and disorders  
Screening is the testing for disease or disease precursors in asymptomatic individuals so that early detection and treatment can be provided for those who test positive for the disease.  
**Excludes1:** encounter for diagnostic examination-code to sign or symptom

Z13.4  
Encounter for screening for certain developmental disorders in childhood  
Encounter for screening for developmental handicaps in early childhood  
**Excludes1:** routine development testing of infant or child (Z00.1-)

Z13.5  
Encounter for screening for eye and ear disorders  
**Excludes2:** encounter for general hearing examination (Z01.1-)  
encounter for general vision examination (Z01.0-)  
Z13.8  
Encounter for screening for other specified diseases and disorders  
**Excludes2:** screening for malignant neoplasms (Z12.-)  
Z13.85  
Encounter for screening for nervous system disorders  
Z13.850  
Encounter for screening for traumatic brain injury

*Encounters for other specific health care (Z40-Z53)*

Categories Z40-Z53 are intended for use to indicate a reason for care. They may be used for patients who have already been treated for a disease or injury, but who are receiving aftercare or prophylactic care, or care to consolidate the treatment, or to deal with a residual state  
**Excludes2:** follow-up examination for medical surveillance after treatment (Z08-Z09)

Z43  
Encounter for attention to artificial openings  
**Includes:** closure of artificial openings  
passage of sounds or bougies through artificial openings  
reforming artificial openings  
removal of catheter from artificial openings  
toilet or cleansing of artificial openings  
**Excludes1:** artificial opening status only, without need for care (Z93.-)  
complications of external stoma (J95.0-, K94.-, N99.5-)  
**Excludes2:** fitting and adjustment of prosthetic and other devices (Z44-Z46)
Z43.0  Encounter for attention to tracheostomy
Z44  Encounter for fitting and adjustment of external prosthetic device
   Includes: removal or replacement of external prosthetic device
   Excludes1: malfunction or other complications of device - see Alphabetical Index
   presence of prosthetic device (Z97.)
Z44.8  Encounter for fitting and adjustment of other external prosthetic devices
Z44.9  Encounter for fitting and adjustment of unspecified external prosthetic device
Z45  Encounter for adjustment and management of implanted device
   Includes: removal or replacement of implanted device
   Excludes1: malfunction or other complications of device
   presence of prosthetic and other devices (Z95-Z97)
   Excludes2: encounter for fitting and adjustment of non-implanted device (Z46.-)
Z45.3  Encounter for adjustment and management of implanted devices of the special senses
   Z45.32  Encounter for adjustment and management of implanted hearing device
   Excludes1: Encounter for fitting and adjustment of hearing aide (Z46.1)
   Z45.320  Encounter for adjustment and management of bone conduction device
   Z45.321  Encounter for adjustment and management of cochlear device
   Z45.328  Encounter for adjustment and management of other implanted hearing device
Z46  Encounter for fitting and adjustment of other devices
   Includes: removal or replacement of other device
   Excludes1: malfunction or other complications of device - see Alphabetical Index
   Excludes2: encounter for fitting and management of implanted devices (Z45.-)
   issue of repeat prescription only (Z76.0)
   presence of prosthetic and other devices (Z95-Z97)
   Z46.1  Encounter for fitting and adjustment of hearing aid
   Excludes1: encounter for adjustment and management of implanted hearing device
   (Z45.32-)
Z51  Encounter for other aftercare
   Z51.8  Encounter for other specified aftercare
   Excludes1: holiday relief care (Z75.5)
   Z51.89  Encounter for other specified aftercare

Persons with potential health hazards related to socioeconomic and psychosocial circumstances (Z55-Z65)
Z57  Occupational exposure to risk factors
   Z57.0  Occupational exposure to noise

Persons encountering health services in other circumstances (Z69-Z76)
Z73  Problems related to life management difficulty
   Excludes2: problems related to socioeconomic and psychosocial circumstances (Z55-Z65)
   Z73.8  Other problems related to life management difficulty
   Z73.82  Dual sensory impairment

Persons with potential health hazards related to family and personal history and certain conditions influencing health status (Z77-Z99)

Code also any follow-up examination (Z08-Z09)
Z77  Other contact with and (suspected) exposures hazardous to health
  Z77.1  Contact with and (suspected) exposure to environmental pollution and hazards in the physical environment
  Z77.12  Contact with and (suspected) exposure to hazards in the physical environment
  Z77.122  Contact with and (suspected) exposure to noise
◊ Z81  Family history of mental and behavioral disorders
Z82  Family history of certain disabilities and chronic diseases (leading to disablement)
  Z82.2  Family history of deafness and hearing loss
  Conditions classifiable to H90-H91
Z83  Family history of other specific disorders
  Excludes2: contact with and (suspected) exposure to communicable disease in the family (Z20.-)
  Z83.5  Family history of eye and ear disorders
  Conditions classifiable to H00-H53, H55-H83, H92-H95
  Excludes2: family history of blindness and visual loss (Z82.1)
  family history of deafness and hearing loss (Z82.2)
  Z83.52  Family history of ear disorders
  Conditions classifiable to H60-H83, H92-H95
  Excludes2: family history of deafness and hearing loss (Z82.2)
Z86  Personal history of certain other diseases
  Code first any follow-up examination after treatment (Z09)
  Z86.5  Personal history of mental and behavioral disorders
  Conditions classifiable to F40-F59
  Z86.59  Personal history of other mental and behavioral disorders
Z87  Personal history of other diseases and conditions
  Code first any follow-up examination after treatment (Z09)
  Z87.7  Personal history of (corrected) congenital malformations
  Conditions classifiable to Q00-Q89 that have been repaired or corrected
  Z87.72  Personal history of (corrected) congenital malformations of nervous system and sense organs
  Z87.721  Personal history of (corrected) congenital malformations of ear
  Z87.73  Personal history of (corrected) congenital malformations of digestive system
  Z87.730  Personal history of (corrected) cleft lip and palate
  Z87.79  Personal history of other (corrected) congenital malformations
  Z87.790  Personal history of (corrected) congenital malformations of face and neck
Z87.8  Personal history of other specified conditions
  Excludes2: personal history of self harm (Z91.5)
  Z87.82  Personal history of other (healed) physical injury and trauma
  Conditions classifiable to S00-T88, except traumatic fractures
  Z87.820  Personal history of traumatic brain injury
  Excludes1: personal history of transient ischemic attack (TIA), and cerebral infarction without residual deficits (Z86.73)
Z90  Acquired absence of organs, not elsewhere classified
  Includes: postprocedural or post-traumatic loss of body part NEC
  Excludes1: congenital absence
  Z90.0  Acquired absence of part of head and neck
  Z90.02  Acquired absence of larynx
  Z90.09  Acquired absence of other part of head and neck
Acquired absence of nose

**Excludes2**: teeth (K08.1)

Z93  Artificial opening status

**Excludes1**: artificial openings requiring attention or management (Z43.-)
- complications of external stoma (J95.0-, K94.-, N99.5-)

Z93.0  Tracheostomy status

Z96  Presence of other functional implants

**Excludes2**: complications of internal prosthetic devices, implants and grafts (T82-T85)
- fitting and adjustment of prosthetic and other devices (Z44-Z46)

Z96.2  Presence of otological and audiological implants

Z96.20  Presence of otological and audiological implant, unspecified

Z96.21  Cochlear implant status

Z96.22  Myringotomy tube(s) status

Z96.29  Presence of other otological and audiological implants
- Presence of bone-conduction hearing device
- Presence of eustachian tube stent
- Stapes replacement

Z96.3  Presence of artificial larynx

Z97  Presence of other devices

**Excludes1**: complications of internal prosthetic devices, implants and grafts (T82-T85)
- fitting and adjustment of prosthetic and other devices (Z44-Z46)

Z97.4  Presence of external hearing-aid
Instructional Notations


Includes

The word 'Includes' appears immediately under certain categories to further define, or give examples of, the content of the category.

Excludes Notes

The ICD-10-CM has two types of excludes notes. Each note has a different definition for use but they are both similar in that they indicate that codes excluded from each other are independent of each other.

Excludes1

A type 1 Excludes note is a pure excludes. It means 'NOT CODED HERE!' An Excludes1 note indicates that the code excluded should never be used at the same time as the code above the Excludes1 note. An Excludes1 is used when two conditions cannot occur together, such as a congenital form versus an acquired form of the same condition.

Excludes2

A type 2 excludes note represents 'Not included here'. An excludes2 note indicates that the condition excluded is not part of the condition it is excluded from but a patient may have both conditions at the same time. When an Excludes2 note appears under a code it is acceptable to use both the code and the excluded code together.

Code First/Use Additional Code notes (etiology/manifestation paired codes)

Certain conditions have both an underlying etiology and multiple body system manifestations due to the underlying etiology. For such conditions the ICD-10-CM has a coding convention that requires the underlying condition be sequenced first followed by the manifestation. Wherever such a combination exists there is a 'use additional code' note at the etiology code, and a 'code first' note at the manifestation code. These instructional notes indicate the proper sequencing order of the codes, etiology followed by manifestation.

In most cases the manifestation codes will have in the code title, 'in diseases classified elsewhere.' Codes with this title area component of the etiology/manifestation convention. The code title indicates that it is a manifestation code. 'In diseases classified elsewhere' codes are never permitted to be used as first listed or principal diagnosis codes. They must be used in conjunction with an underlying condition code and they must be listed following the underlying condition.

Code Also

A code also note instructs that 2 codes may be required to fully describe a condition but the sequencing of the two codes is discretionary, depending on the severity of the conditions and the reason for the encounter.

7th characters and placeholder X

For codes less than 6 characters that require a 7th character a placeholder X should be assigned for all characters less than 6. The 7th character must always be the 7th character of a code