2015 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 2015 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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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*

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

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

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>C00</td>
<td>Malignant neoplasm of lip</td>
</tr>
<tr>
<td>C00.0</td>
<td>Malignant neoplasm of external upper lip</td>
</tr>
<tr>
<td>C00.1</td>
<td>Malignant neoplasm of external lower lip</td>
</tr>
<tr>
<td>C00.2</td>
<td>Malignant neoplasm of external lip, unspecified</td>
</tr>
<tr>
<td>C00.3</td>
<td>Malignant neoplasm of upper lip, inner aspect</td>
</tr>
<tr>
<td>C00.4</td>
<td>Malignant neoplasm of lower lip, inner aspect</td>
</tr>
<tr>
<td>C00.5</td>
<td>Malignant neoplasm of lip, unspecified, inner aspect</td>
</tr>
<tr>
<td>C00.6</td>
<td>Malignant neoplasm of commissure of lip, unspecified</td>
</tr>
<tr>
<td>C00.8</td>
<td>Malignant neoplasm of overlapping sites of lip</td>
</tr>
<tr>
<td>C00.9</td>
<td>Malignant neoplasm of lip, unspecified</td>
</tr>
<tr>
<td>C01</td>
<td>Malignant neoplasm of base of tongue</td>
</tr>
<tr>
<td>C02</td>
<td>Malignant neoplasm of other and unspecified parts of tongue</td>
</tr>
<tr>
<td>C02.0</td>
<td>Malignant neoplasm of dorsal surface of tongue</td>
</tr>
<tr>
<td>C02.1</td>
<td>Malignant neoplasm of border of tongue</td>
</tr>
<tr>
<td>C02.2</td>
<td>Malignant neoplasm of ventral surface of tongue</td>
</tr>
<tr>
<td>C02.3</td>
<td>Malignant neoplasm of anterior two-thirds of tongue, part unspecified</td>
</tr>
<tr>
<td>C02.4</td>
<td>Malignant neoplasm of lingual tonsil</td>
</tr>
<tr>
<td>C02.8</td>
<td>Malignant neoplasm of overlapping sites of tongue</td>
</tr>
<tr>
<td>C02.9</td>
<td>Malignant neoplasm of tongue, unspecified</td>
</tr>
<tr>
<td>C03</td>
<td>Malignant neoplasm of gum</td>
</tr>
<tr>
<td>C04</td>
<td>Malignant neoplasm of floor of mouth</td>
</tr>
<tr>
<td>C05</td>
<td>Malignant neoplasm of palate</td>
</tr>
<tr>
<td>C05.0</td>
<td>Malignant neoplasm of hard palate</td>
</tr>
<tr>
<td>C05.1</td>
<td>Malignant neoplasm of soft palate</td>
</tr>
<tr>
<td>C05.2</td>
<td>Malignant neoplasm of uvula</td>
</tr>
<tr>
<td>C06</td>
<td>Malignant neoplasm of other and unspecified parts of mouth</td>
</tr>
<tr>
<td>C08</td>
<td>Malignant neoplasm of other and unspecified major salivary glands</td>
</tr>
<tr>
<td>C09</td>
<td>Malignant neoplasm of tonsil</td>
</tr>
<tr>
<td>C10</td>
<td>Malignant neoplasm of oropharynx</td>
</tr>
<tr>
<td>C10.1</td>
<td>Malignant neoplasm of anterior surface of epiglottis</td>
</tr>
<tr>
<td>C10.2</td>
<td>Malignant neoplasm of lateral wall of oropharynx</td>
</tr>
<tr>
<td>C10.3</td>
<td>Malignant neoplasm of posterior wall of oropharynx</td>
</tr>
<tr>
<td>C11</td>
<td>Malignant neoplasm of nasopharynx</td>
</tr>
<tr>
<td>C11.0</td>
<td>Malignant neoplasm of superior wall of nasopharynx</td>
</tr>
</tbody>
</table>

- ✓ 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 without behavioral disturbance
  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)
            Excludes2: speech articulation impairment due to apraxia (R48.2)
            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

☑  Code typically used by SLPs  ☥ Additional digits not listed here
**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:** Alcohol'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)

✔ Code typically used by SLPs  ◊ Additional digits not listed here
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
    Other epilepsy NOS
    Other epilepsy without intractability 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
  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

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

✓ Code typically used by SLPs  ◊ Additional digits not listed here
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 sequela 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
        Stuttering 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

The original ICD-9 series for cerebrovascular disease (438) is expanded under ICD-10. Check the entire I69.- category before assigning a code.
### 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 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
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>I69.821</td>
<td>Dysphasia following other cerebrovascular disease</td>
</tr>
<tr>
<td>I69.822</td>
<td>Dysarthria following other cerebrovascular disease</td>
</tr>
<tr>
<td>I69.823</td>
<td>Fluency disorder following other cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Stuttering following nontraumatic subarachnoid hemorrhage</td>
</tr>
<tr>
<td>I69.828</td>
<td>Other speech and language deficits following other cerebrovascular disease</td>
</tr>
</tbody>
</table>

**I69.89 Other sequelae of other cerebrovascular disease**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>I69.890</td>
<td>Apraxia following other cerebrovascular disease</td>
</tr>
<tr>
<td>I69.891</td>
<td>Dysphagia following other cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Use additional code to identify the type of dysphagia, if known (R13.1-)</td>
</tr>
<tr>
<td>I69.892</td>
<td>Facial weakness following other cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Facial droop following other cerebrovascular disease</td>
</tr>
<tr>
<td>I69.893</td>
<td>Ataxia following other cerebrovascular disease</td>
</tr>
<tr>
<td>I69.898</td>
<td>Other sequelae of other cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Alteration of sensation following other cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Disturbance of vision following other cerebrovascular disease</td>
</tr>
</tbody>
</table>

**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.-)

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>I69.90</td>
<td>Unspecified sequelae of unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.91</td>
<td>Cognitive deficits following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.92</td>
<td>Speech and language deficits following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Aphasia following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.921</td>
<td>Dysphasia following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.922</td>
<td>Dysarthria following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.923</td>
<td>Fluency disorder following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Stuttering following nontraumatic subarachnoid hemorrhage</td>
</tr>
<tr>
<td>I69.928</td>
<td>Other speech and language deficits following unspecified cerebrovascular disease</td>
</tr>
</tbody>
</table>

**I69.99 Other sequelae of unspecified cerebrovascular disease**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>I69.990</td>
<td>Apraxia following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.991</td>
<td>Dysphagia following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Use additional code to identify the type of dysphagia, if known (R13.1-)</td>
</tr>
<tr>
<td>I69.992</td>
<td>Facial weakness following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Facial droop following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.993</td>
<td>Ataxia following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td>I69.998</td>
<td>Other sequelae following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Alteration in sensation following unspecified cerebrovascular disease</td>
</tr>
<tr>
<td></td>
<td>Disturbance of vision following unspecified cerebrovascular disease</td>
</tr>
</tbody>
</table>

**Use additional code to identify the sequelae**

---

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

**Acute upper respiratory infections (J00-J06)**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>J02</td>
<td>Acute pharyngitis</td>
</tr>
</tbody>
</table>

**Code typically used by SLPs**

**Additional digits not listed here**
◊ 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
◊ J35 Chronic diseases of tonsils and adenoids

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)

J37.1 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

✓ Code typically used by SLPs  ◊ Additional digits not listed here
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  Other diseases of larynx

J39.3  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)

J39.8  Other specified diseases of upper respiratory tract

J39.9  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
ever elsewhere classified (J95)**

J95  Intraoperative and postprocedural complications and disorders of respiratory system, not
ever 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
      Glossoceles
      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

☑ Code typically used by SLPs  ◊ Additional digits not listed here
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)
M26.00 Unspecified anomaly of jaw size
M26.01 Maxillary hyperplasia
M26.02 Maxillary hypoplasia
M26.03 Mandibular hyperplasia
M26.04 Mandibular hypoplasia
M26.05 Macrogenia
M26.06 Microgenia
M26.07 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
M26.11 Maxillary asymmetry
M26.12 Other jaw asymmetry
M26.19 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
M26.211 Malocclusion, Angle's class I
Neutro-occlusion
M26.212 Malocclusion, Angle's class II
Disto-occlusion Division I
Disto-occlusion Division II
M26.213 Malocclusion, Angle's class III
Mesio-occlusion
M26.219 Malocclusion, Angle's class, unspecified
M26.22 Open occlusal relationship
M26.220 Open anterior occlusal relationship
Anterior openbite
M26.221 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

✅ Code typically used by SLPs  ◊ Additional digits not listed here
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  

Excludes1: bruxism (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  

---  

M26.55 Centric occlusion maximum intercuspation discrepancy  

Excludes1: 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  

Excludes2: 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  

---  

M26.82 Posterior soft tissue impingement  

---  

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
         Pachgyria
      Excludes1: congenital malformations of corpus callosum (Q04.0)
   ◊ Q05 Spina bifida

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

Excludes2: cleft lip and cleft palate (Q35-Q37)
   congenital malformation of cervical spine (Q05.0, Q05.5, Q67.5, Q76.0-Q76.4)
   congenital malformation of larynx (Q31.-)
   congenital malformation of lip NEC (Q38.0)
   congenital malformation of nose (Q30.-)
   congenital malformation of parathyroid gland (Q89.2)
   congenital malformation of thyroid gland (Q89.2)

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

✓ Code typically used by SLPs  ◊ Additional digits not listed here
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)
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 Laryngoele
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

✓ Code typically used by SLPs ○ Additional digits not listed here
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: cheiloschisis
  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

✓ Code typically used by SLPs  ◊ Additional digits not listed here
**Excludes 1:** 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

**Excludes 1:** 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

**Excludes 1:** 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

**Excludes 1:** dentofacial anomalies [including malocclusion] (M26-)
- Syphilitic saddle nose (A50.5)

**✓** Code typically used by SLPs  
**◊** Additional digits not listed here
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
   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

✔ Code typically used by SLPs  ☉ Additional digits not listed here
✓ 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

✓ Code typically used by SLPs  ◊ Additional digits not listed here
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

**Excludes1:** 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

**Excludes1:** intellectual disabilities (F70-F79)

R41.84 Other specified cognitive deficit

✓ R41.840 Attention and concentration deficit

**Excludes1:** 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

**Excludes1:** 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)

**Excludes2:** disturbances of skin sensation (R20.-)

R44.0 Auditory hallucinations

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

R47 Speech disturbances, not elsewhere classified

**Excludes1:** 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

**Excludes1:** aphasia following cerebrovascular disease (I69. with final characters -20)
 progressive isolated aphasia (G31.01)

✓ R47.02 Dysphasia

**Excludes1:** dysphasia following cerebrovascular disease (I69. with final characters -21)

✓ Code typically used by SLPs  ◊ Additional digits not listed here
R47.1 Dysarthria and anarthria

Excludes1: dysarthria following cerebrovascular disease (I69. with final characters -22)

R47.8 Other speech disturbances

Excludes1: dysarthria following cerebrovascular disease (I69. 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:

Parkinson’s disease (G20)

Excludes1: adult onset fluency disorder (F98.5)

childhood onset fluency disorder (F80.81)

flueny 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 Aphonia

Loss of voice

R49.2 Hypernasality and hyponasality

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)

Code typically used by SLPs

Additional digits not listed here
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.-)

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

Excludes 2: 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
  Excludes 1: 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
  Hyoid bone
  Larynx
  Thyroid cartilage
  Trachea
  The appropriate 7th character is to be added to code S12.8
  A - initial encounter
  D - subsequent encounter
  S - sequela

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
  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
Excludes2: foreign body in pharynx (T17.2-)
  T18.1 Foreign body in esophagus

Ch. 20 External causes of morbidity (V00-Y99)

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.

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

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.

Persons encountering health services for examinations (Z00-Z13)

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

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

Z01 Encounter for other special examination without complaint, suspected or reported diagnosis

✓ Code typically used by SLPs  ◊ Additional digits not listed here
**Includes:** routine examination of specific system

**Note:** Codes from category Z01 represent the reason for the encounter. A separate procedure code is required to identify any examinations or procedures performed

**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 (Z00.0-)
- 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

**Z13** 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

✔ Code typically used by SLPs  ◊ Additional digits not listed here
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

✔ Code typically used by SLPs  ◊ Additional digits not listed here
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.