2017 ICD-10-CM Diagnosis Codes
Related to Speech, Language, and Swallowing Disorders
General Information

This ASHA document provides a listing of the 2017 *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.

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/ICD-10/](http://www.asha.org/Practice/reimbursement/coding/ICD-10/).

A listing of new ICD-10-CM codes effective October 1, 2016, is available at [www.asha.org/Practice/reimbursement/coding/New-and-Revised-ICD-10-CM-Codes-for-SLP/](http://www.asha.org/Practice/reimbursement/coding/New-and-Revised-ICD-10-CM-Codes-for-SLP/).

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 Diagnostic Codes

Overview
On October 1, 2015, the International Classification of Diseases, 10th Revision (ICD-10) replaced the 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 ICD-10 includes 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 (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) was October 1, 2015.

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. Speech-language pathologists whose services may be billed to third-party payers may also be required to report ICD-10-CM codes.

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-10-CM Diagnosis Codes for Audiology and Speech-Language Pathology: www.asha.org/Practice/reimbursement/coding/ICD-10/
- ICD-10-CM Coding FAQs for Audiologists and SLPS: www.asha.org/Practice/reimbursement/coding/ICD-10-CM-Coding-FAQs-for-Audiologists-and-SLPS/
- 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-9 to ICD-10 Mapping Tool for Audiologists and Speech-Language Pathologists: www.asha.org/icdmapping.aspx
ICD-10-CM Tabular List of Diseases and Injuries
Related to speech, language, and swallowing disorders

<table>
<thead>
<tr>
<th>Ch. 2 Neoplasms (C00-D49)</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Malignant neoplasms of lip, oral cavity, and pharynx (C00-C14)</strong></td>
<td></td>
</tr>
<tr>
<td>C00 Malignant neoplasm of lip</td>
<td></td>
</tr>
<tr>
<td>C00.0 Malignant neoplasm of external upper lip</td>
<td></td>
</tr>
<tr>
<td>C00.1 Malignant neoplasm of external lower lip</td>
<td></td>
</tr>
<tr>
<td>C00.2 Malignant neoplasm of external lip, unspecified</td>
<td></td>
</tr>
<tr>
<td>C00.3 Malignant neoplasm of upper lip, inner aspect</td>
<td></td>
</tr>
<tr>
<td>C00.4 Malignant neoplasm of lower lip, inner aspect</td>
<td></td>
</tr>
<tr>
<td>C00.5 Malignant neoplasm of lip, unspecified, inner aspect</td>
<td></td>
</tr>
<tr>
<td>C00.6 Malignant neoplasm of commissure of lip, unspecified</td>
<td></td>
</tr>
<tr>
<td>C00.8 Malignant neoplasm of overlapping sites of lip</td>
<td></td>
</tr>
<tr>
<td>C00.9 Malignant neoplasm of lip, unspecified</td>
<td></td>
</tr>
<tr>
<td>C01 Malignant neoplasm of base of tongue</td>
<td></td>
</tr>
<tr>
<td>C02 Malignant neoplasm of other and unspecified parts of tongue</td>
<td></td>
</tr>
<tr>
<td>C02.0 Malignant neoplasm of dorsal surface of tongue</td>
<td></td>
</tr>
<tr>
<td>C02.1 Malignant neoplasm of border of tongue</td>
<td></td>
</tr>
<tr>
<td>C02.2 Malignant neoplasm of ventral surface of tongue</td>
<td></td>
</tr>
<tr>
<td>C02.3 Malignant neoplasm of anterior two-thirds of tongue, part unspecified</td>
<td></td>
</tr>
<tr>
<td>C02.4 Malignant neoplasm of lingual tonsil</td>
<td></td>
</tr>
<tr>
<td>C02.8 Malignant neoplasm of overlapping sites of tongue</td>
<td></td>
</tr>
<tr>
<td>C02.9 Malignant neoplasm of tongue, unspecified</td>
<td></td>
</tr>
<tr>
<td>◊ C03 Malignant neoplasm of gum</td>
<td></td>
</tr>
<tr>
<td>◊ C04 Malignant neoplasm of floor of mouth</td>
<td></td>
</tr>
<tr>
<td>C05 Malignant neoplasm of palate</td>
<td></td>
</tr>
<tr>
<td>C05.0 Malignant neoplasm of hard palate</td>
<td></td>
</tr>
<tr>
<td>C05.1 Malignant neoplasm of soft palate</td>
<td></td>
</tr>
<tr>
<td>C05.2 Malignant neoplasm of uvula</td>
<td></td>
</tr>
<tr>
<td>◊ C06 Malignant neoplasm of other and unspecified parts of mouth</td>
<td></td>
</tr>
<tr>
<td>◊ C08 Malignant neoplasm of other and unspecified major salivary glands</td>
<td></td>
</tr>
<tr>
<td>◊ C09 Malignant neoplasm of tonsil</td>
<td></td>
</tr>
<tr>
<td>C10 Malignant neoplasm of oropharynx</td>
<td></td>
</tr>
<tr>
<td>C10.1 Malignant neoplasm of anterior surface of epiglottis</td>
<td></td>
</tr>
<tr>
<td>C10.2 Malignant neoplasm of lateral wall of oropharynx</td>
<td></td>
</tr>
<tr>
<td>C10.3 Malignant neoplasm of posterior wall of oropharynx</td>
<td></td>
</tr>
<tr>
<td>C11 Malignant neoplasm of nasopharynx</td>
<td></td>
</tr>
<tr>
<td>C11.0 Malignant neoplasm of superior wall of nasopharynx</td>
<td></td>
</tr>
</tbody>
</table>

**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 [www.cdc.gov/nchs/icd/icd10cm.htm](http://www.cdc.gov/nchs/icd/icd10cm.htm).
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>C11.1</td>
<td>Malignant neoplasm of posterior wall of nasopharynx</td>
</tr>
<tr>
<td>C11.2</td>
<td>Malignant neoplasm of lateral wall of nasopharynx</td>
</tr>
<tr>
<td>C11.3</td>
<td>Malignant neoplasm of anterior wall of nasopharynx</td>
</tr>
<tr>
<td>C11.8</td>
<td>Malignant neoplasm of overlapping sites of nasopharynx</td>
</tr>
<tr>
<td>C14</td>
<td>Malignant neoplasm of other and ill-defined sites in the lip, oral cavity and pharynx</td>
</tr>
<tr>
<td>C14.0</td>
<td>Malignant neoplasm of pharynx, unspecified</td>
</tr>
<tr>
<td>C15</td>
<td>Malignant neoplasm of esophagus</td>
</tr>
</tbody>
</table>

**Malignant neoplasms of digestive organs (C15-C26)**

**Malignant neoplasms of respiratory and intrathoracic organs (C30-C39)**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>C30</td>
<td>Malignant neoplasm of nasal cavity and middle ear</td>
</tr>
<tr>
<td>C32</td>
<td>Malignant neoplasm of larynx</td>
</tr>
<tr>
<td>C32.0</td>
<td>Malignant neoplasm of glottis</td>
</tr>
<tr>
<td>C32.1</td>
<td>Malignant neoplasm of supraglottis</td>
</tr>
<tr>
<td>C32.2</td>
<td>Malignant neoplasm of subglottis</td>
</tr>
<tr>
<td>C32.3</td>
<td>Malignant neoplasm of laryngeal cartilage</td>
</tr>
<tr>
<td>C32.8</td>
<td>Malignant neoplasm of overlapping sites of larynx</td>
</tr>
<tr>
<td>C32.9</td>
<td>Malignant neoplasm of larynx, unspecified</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>C33</td>
<td>Malignant neoplasm of trachea</td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>C71</td>
<td>Malignant neoplasm of brain</td>
</tr>
</tbody>
</table>

**In situ neoplasms (D00-D09)**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D00</td>
<td>Carcinoma in situ of oral cavity, esophagus and stomach</td>
</tr>
<tr>
<td>D00.07</td>
<td>Carcinoma in situ of tongue</td>
</tr>
<tr>
<td>D02</td>
<td>Carcinoma in situ of middle ear and respiratory system</td>
</tr>
<tr>
<td>D02.1</td>
<td>Carcinoma in situ of trachea</td>
</tr>
</tbody>
</table>

**Benign neoplasms, except benign neuroendocrine tumors (D10-D36)**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>D10</td>
<td>Benign neoplasm of mouth and pharynx</td>
</tr>
<tr>
<td>D14</td>
<td>Benign neoplasm of middle ear and respiratory system</td>
</tr>
<tr>
<td>D14.1</td>
<td>Benign neoplasm of larynx</td>
</tr>
<tr>
<td>D14.2</td>
<td>Benign neoplasm of trachea</td>
</tr>
<tr>
<td>D33</td>
<td>Benign neoplasm of brain and other parts of central nervous system</td>
</tr>
<tr>
<td>D38</td>
<td>Neoplasm of uncertain behavior of middle ear and respiratory and intrathoracic organs</td>
</tr>
</tbody>
</table>

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

**Mental disorders due to known physiological conditions (F01-F09)**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>F01</td>
<td>Vascular dementia</td>
</tr>
<tr>
<td>F01.5</td>
<td>Vascular dementia</td>
</tr>
<tr>
<td>F01.50</td>
<td>Vascular dementia without behavioral disturbance</td>
</tr>
<tr>
<td>F01.51</td>
<td>Vascular dementia with behavioral disturbance</td>
</tr>
</tbody>
</table>

**Dementia in other diseases classified elsewhere**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>F02</td>
<td>Dementia in other diseases classified elsewhere</td>
</tr>
</tbody>
</table>

**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

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

NEW

F80.82 Social pragmatic communication disorder
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 (F81.8)
   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

See bit.ly/2jqM1DL for more information on use of this new code.
**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.-)

temporary mutism a part of separation anxiety in young children (F93.0)
F98  Other behavioral and emotional disorders with onset usually occurring in childhood and adolescence
    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
  - 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, 169.11, 169.21, 169.31, 169.81, 169.91)
  - 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
        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

Diseases of myoneural junction and muscle (G70-G73)

G71  Primary disorders of muscles
    Excludes2: arthrogryposis multiplex congenita (Q74.3)
        metabolic disorders (E70-E88)
        myositis (M60.-)
    G71.0  Muscular dystrophy

✓ Code typically used by SLPs  ◊ Additional digits not listed here
Autosomal recessive, childhood type, muscular dystrophy resembling Duchenne or Becker muscular dystrophy
Benign [Becker] muscular dystrophy
Benign scapuloperoneal muscular dystrophy with early contractures [Emery-Dreifuss]
Congenital muscular dystrophy NOS
Congenital muscular dystrophy with specific morphological abnormalities of the muscle fiber
Distal muscular dystrophy
Facioscapulohumeral muscular dystrophy
Limb-girdle muscular dystrophy
Ocular muscular dystrophy
Oculopharyngeal muscular dystrophy
Scapuloperoneal muscular dystrophy
Severe [Duchenne] muscular dystrophy

G71.1  Myotonic disorders
G71.11  Myotonic muscular dystrophy
        Dystrophia myotonica [Steinert]
        Myotonia atrophica
        Myotonic muscular dystrophy
        Dystrophia myotonica [Steinert]
        Myotonia atrophica
G71.12  Myotonia congenital
        Acetazolamide responsive myotonia congenita
        Dominant myotonia congenita [Thomsen disease]
        Myotonia levior
        Recessive myotonia congenita [Becker disease]
G71.13  Myotonic chondrodystrophy
        Chondrodystrophic myotonias
        Congenital myotonic chondrodystrophy
        Schwartz-Jampel disease
G71.14  Drug induced myotonia
        Use additional code for adverse effect, if applicable, to identify drug (T36-T50 with fifth or sixth character 5)
G71.19  Other specified myotonic disorders
        Myotonia fluctuans
        Myotonia permanens
        Neuromyotonia [Isaacs]
        Paramyotonia congenita (of von Eulenburg)
        Pseudomyotonia
        Symptomatic myotonia

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
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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 sequelae 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
      NEW I69.010 Attention and concentration deficit following nontraumatic subarachnoid hemorrhage
      NEW I69.011 Memory deficit following nontraumatic subarachnoid hemorrhage
      NEW I69.012 Visuospatial deficit and spatial neglect following nontraumatic subarachnoid hemorrhage
      NEW I69.013 Psychomotor deficit following nontraumatic subarachnoid hemorrhage

The original ICD-9 series for cerebrovascular disease (438) is expanded under ICD-10. Check the entire I69.- category before assigning a code.

See bit.ly/2jaMIDL for more information on use of the new codes related to cognitive deficits.
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>I69.014</td>
<td>Frontal lobe and executive function deficit following nontraumatic subarachnoid hemorrhage</td>
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<td>I69.022</td>
<td>Dysarthria following nontraumatic subarachnoid hemorrhage</td>
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<td>I69.023</td>
<td>Fluency disorder following nontraumatic subarachnoid hemorrhage</td>
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<td>I69.028</td>
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<td>I69.112</td>
<td>Visuospatial deficit and spatial neglect following nontraumatic intracerebral hemorrhage</td>
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<td>I69.113</td>
<td>Psychomotor deficit following nontraumatic intracerebral hemorrhage</td>
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<td>I69.114</td>
<td>Frontal lobe and executive function deficit following nontraumatic intracerebral hemorrhage</td>
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<td>I69.115</td>
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<td>I69.118</td>
<td>Other symptoms and signs involving cognitive functions following nontraumatic intracerebral hemorrhage</td>
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<tr>
<td>I69.119</td>
<td>Unspecified symptoms and signs involving cognitive functions following nontraumatic intracerebral hemorrhage</td>
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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

NEW 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
  NEW I69.210 Attention and concentration deficit following other nontraumatic intracranial hemorrhage
  NEW I69.211 Memory deficit following other nontraumatic intracranial hemorrhage
  NEW I69.212 Visuospatial deficit and spatial neglect following other nontraumatic intracranial hemorrhage
  NEW I69.213 Psychomotor deficit following other nontraumatic intracranial hemorrhage
  NEW I69.214 Frontal lobe and executive function deficit following other nontraumatic intracranial hemorrhage
  NEW I69.215 Cognitive social or emotional deficit following other nontraumatic intracranial hemorrhage
  NEW I69.218 Other symptoms and signs involving cognitive functions following other nontraumatic intracranial hemorrhage
  NEW I69.219 Unspecified symptoms and signs involving cognitive functions following other nontraumatic intracranial hemorrhage

NEW 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
  Stuttering following nontraumatic subarachnoid hemorrhage
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<td>Sequelae of stroke NOS</td>
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<td>Cognitive deficits following cerebral infarction</td>
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</table>

**Code typically used by SLPs**

**Additional digits not listed here**
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

NEW  ✓ I69.810  Attention and concentration deficit following other cerebrovascular disease

NEW  ✓ I69.811  Memory deficit following cerebral infarction following other cerebrovascular disease

NEW  I69.812  Visuospatial deficit and spatial neglect following other cerebrovascular disease

NEW  I69.813  Psychomotor deficit following other cerebrovascular disease

NEW  ✓ I69.814  Frontal lobe and executive function deficit following other cerebrovascular disease

NEW  ✓ I69.815  Cognitive social or emotional deficit following other cerebrovascular disease

NEW  I69.818  Other symptoms and signs involving cognitive functions following other cerebrovascular disease

NEW  I69.819  Unspecified symptoms and signs involving cognitive functions following other cerebrovascular disease

I69.82  Speech and language deficits following other cerebrovascular disease

✓ I69.820  Aphasia following other cerebrovascular disease

✓ 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

✓ Code typically used by SLPs  ○ Additional digits not listed here
NEW  I69.910  Attention and concentration deficit following unspecified cerebrovascular disease
NEW  I69.911  Memory deficit following cerebral infarction following unspecified cerebrovascular disease
NEW  I69.912  Visuospatial deficit and spatial neglect following unspecified cerebrovascular disease
NEW  I69.913  Psychomotor deficit following unspecified cerebrovascular disease
NEW  I69.914  Frontal lobe and executive function deficit following unspecified cerebrovascular disease
NEW  I69.915  Cognitive social or emotional deficit following unspecified cerebrovascular disease
NEW  I69.918  Other symptoms and signs involving cognitive functions following unspecified cerebrovascular disease
NEW  I69.919  Unspecified symptoms and signs involving cognitive functions 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
   ✔ 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
   ✔ I69.993  Ataxia following unspecified cerebrovascular disease
I69.998  Other sequelae following unspecified cerebrovascular disease
   ✔ I69.998  Other sequelae 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
   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)
               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 polyp (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

---

- Code typically used by SLPs
- Additional digits not listed here

There is no longer a code for vocal cord paresis. Consider using the J38.0 series instead. Documentation should indicate the paresis.

J38.5 is an option for a diagnosis of paradoxical vocal fold motion.
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

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 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

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)
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>M26.00</td>
<td>Unspecified anomaly of jaw size</td>
</tr>
<tr>
<td>M26.01</td>
<td>Maxillary hyperplasia</td>
</tr>
<tr>
<td>M26.02</td>
<td>Maxillary hypoplasia</td>
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<tr>
<td>M26.03</td>
<td>Mandibular hyperplasia</td>
</tr>
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<td>M26.04</td>
<td>Mandibular hypoplasia</td>
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<td>M26.05</td>
<td>Macroelia</td>
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<tr>
<td>M26.06</td>
<td>Microelia</td>
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<tr>
<td>M26.07</td>
<td>Excessive tuberosity of jaw</td>
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<tr>
<td></td>
<td>Entire maxillary tuberosity</td>
</tr>
<tr>
<td>M26.09</td>
<td>Other specified anomalies of jaw size</td>
</tr>
<tr>
<td>M26.1</td>
<td>Anomalies of jaw-cranial base relationship</td>
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<tr>
<td>M26.10</td>
<td>Unspecified anomaly of jaw-cranial base relationship</td>
</tr>
<tr>
<td>M26.11</td>
<td>Maxillary asymmetry</td>
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<tr>
<td>M26.12</td>
<td>Other jaw asymmetry</td>
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<tr>
<td>M26.19</td>
<td>Other specified anomalies of jaw-cranial base relationship</td>
</tr>
<tr>
<td>M26.2</td>
<td>Anomalies of dental arch relationship</td>
</tr>
<tr>
<td>M26.20</td>
<td>Unspecified anomaly of dental arch relationship</td>
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<tr>
<td>M26.21</td>
<td>Malocclusion, Angle's class</td>
</tr>
<tr>
<td></td>
<td>M26.211 Malocclusion, Angle's class I</td>
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<tr>
<td></td>
<td>Neutro-occlusion</td>
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<td></td>
<td>M26.212 Malocclusion, Angle's class II</td>
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<tr>
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<td>Disto-occlusion Division I</td>
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<tr>
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<td>Disto-occlusion Division II</td>
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<td>M26.213 Malocclusion, Angle's class III</td>
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<tr>
<td></td>
<td>Mesio-occlusion</td>
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<td>M26.219 Malocclusion, Angle's class, unspecified</td>
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<tr>
<td>M26.22</td>
<td>Open occlusal relationship</td>
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<td>M26.220 Open anterior occlusal relationship</td>
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<td>Anterior openbite</td>
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<tr>
<td></td>
<td>M26.221 Open posterior occlusal relationship</td>
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<tr>
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<td>Posterior openbite</td>
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<tr>
<td>M26.23</td>
<td>Excessive horizontal overlap</td>
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<td>Excessive horizontal overjet</td>
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<tr>
<td>M26.24</td>
<td>Reverse articulation</td>
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<td>Crossbite (anterior) (posterior)</td>
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<tr>
<td>M26.25</td>
<td>Anomalies of interarch distance</td>
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<tr>
<td>M26.29</td>
<td>Other anomalies of dental arch relationship</td>
</tr>
<tr>
<td></td>
<td>Midline deviation of dental arch</td>
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<tr>
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<td>Overbite (excessive) deep</td>
</tr>
<tr>
<td></td>
<td>Overbite (excessive) horizontal</td>
</tr>
<tr>
<td></td>
<td>Overbite (excessive) vertical</td>
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<tr>
<td></td>
<td>Posterior lingual occlusion of mandibular teeth</td>
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<tr>
<td>M26.3</td>
<td>Anomalies of tooth position of fully erupted tooth or teeth</td>
</tr>
<tr>
<td>M26.4</td>
<td>Malocclusion, unspecified</td>
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<tr>
<td>M26.5</td>
<td>Dentofacial functional abnormalities</td>
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<tr>
<td></td>
<td>Excludes1: bruxism (F45.8)</td>
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<tr>
<td></td>
<td>teeth-grinding NOS (F45.8)</td>
</tr>
<tr>
<td></td>
<td>M26.50 Dentofacial functional abnormalities, unspecified</td>
</tr>
</tbody>
</table>
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
    **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
    NEW M26.601 Right temporomandibular joint disorder
    NEW M26.602 Left temporomandibular joint disorder
    NEW M26.603 Bilateral temporomandibular joint disorder
    NEW M26.609 Unspecified temporomandibular joint disorder
    NEW M26.61 Adhesions and ankylosis of temporomandibular joint
        NEW M26.611 Adhesions and ankylosis of right temporomandibular joint
        NEW M26.612 Adhesions and ankylosis of left temporomandibular joint
        NEW M26.613 Adhesions and ankylosis of bilateral temporomandibular joint
        NEW M26.619 Adhesions and ankylosis of temporomandibular joint, unspecified side
    NEW M26.62 Arthralgia of temporomandibular joint
        NEW M26.621 Arthralgia of right temporomandibular joint
        NEW M26.622 Arthralgia of left temporomandibular joint
        NEW M26.623 Arthralgia of bilateral temporomandibular joint
        NEW M26.629 Arthralgia of temporomandibular joint, unspecified side
    NEW M26.63 Articular disc disorder of temporomandibular joint
        NEW M26.631 Articular disc disorder of right temporomandibular joint
        NEW M26.632 Articular disc disorder of left temporomandibular joint
        NEW M26.633 Articular disc disorder of bilateral temporomandibular joint
        NEW M26.639 Articular disc disorder of temporomandibular joint, unspecified side
    NEW M26.69 Other specified disorders of temporomandibular joint

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

- Code typically used by SLPs
- Additional digits not listed here
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

Ch. 16 Certain conditions originating in the perinatal period (P00-P96)

Newborn affected by maternal factors and by complications of pregnancy, labor, and delivery (P00-P04)

Note: These codes are for use when the listed maternal conditions are specified as the cause of confirmed morbidity or potential morbidity which have their origin in the perinatal period (before birth through the first 28 days after birth)

P04 Newborn affected by noxious substances transmitted via placenta or breast milk
   Includes: nonteratogenic effects of substances transmitted via placenta
   Excludes2: congenital malformations
       encounter for observation of newborn for suspected diseases and conditions ruled out (Z05.-)
       neonatal jaundice from excessive hemolysis due to drugs or toxins transmitted from mother (P58.4)
       newborn in contact with and (suspected) exposures hazardous to health not transmitted via placenta or breast milk (Z77.-)
P04.3 Newborn (suspected to be) affected by maternal use of alcohol
   Excludes1: fetal alcohol syndrome (Q86.0)

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)
   P92.1 Regurgitating and rumination of newborn
   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)**

**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
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 Laryngoecele
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:** cheilosophisis
    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

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

Other Congenital Malformations (Q80-Q89)

Q86 Congenital malformation syndromes due to known exogenous causes, not elsewhere classified
  Excludes2: iodine-deficiency-related hypothyroidism (E00-E02)
  nonteratogenic effects of substances transmitted via placenta or breast milk (P04.-)
  Q86.0 Fetal alcohol syndrome (dysmorphic)

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

Q99  Other chromosome abnormalities, not elsewhere classifiable
   Q99.2  Fragile X chromosome
          Fragile X syndrome

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

Symptoms and signs involving the circulatory and respiratory systems (R00-R09)

R05  Cough
    Excludes1: cough with hemorrhage (R04.2)
               smoker’s cough (J41.0)

R06  Abnormalities of breathing
    Excludes1: acute respiratory distress syndrome (J80)
               respiratory arrest (R09.2)
               respiratory arrest of newborn (P28.81)
               respiratory distress syndrome of newborn (P22.-)
               respiratory failure (J96.-)
               respiratory failure of newborn (P28.5)

R06.0  Dyspnea
    Excludes1: tachypnea NOS (R06.82)
               transient tachypnea of newborn (P22.1)

R06.00  Dyspnea, unspecified
R06.01  Orthopnea
R06.02  Shortness of breath
R06.09  Other forms of dyspnea

R06.1  Stridor
    Excludes1: congenital laryngeal stridor
               (P28.89) laryngismus (stridulus) (J38.5)

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)**

**Excludes2:** symptoms and signs constituting part of a pattern of mental disorder (F01-F99)

R40  Somnolence, stupor and coma
    **Excludes1:** neonatal coma (P91.5)
    somnolence, stupor and coma in diabetes (E08-E13)
    somnolence, stupor and coma in hepatic failure (K72.-)
    somnolence, stupor and coma in hypoglycemia (nondiabetic) (E15)
  R40.2  Coma
    **Code first** any associated:
      Fracture of skull (S02.-)
      Intracranial injury (S06.-)
R40.24 Glasgow coma scale, total score

**Note:** Assign a code from subcategory R40.24, when only the total coma score is documented.

The following appropriate 7th character is to be added to subcategory R40.24-:
- 0 – unspecified time
- 1 – in the field [EMT or ambulance]
- 2 – at arrival to emergency department
- 3 – at hospital admission
- 4 – 24 hours or more after hospital admission

R40.241 Glasgow coma scale score 13-15
R40.242 Glasgow coma scale score 9-12
R40.243 Glasgow coma scale score 3-8
R40.244 Other coma, without documented Glasgow coma scale score, or with partial score reported

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

**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

☑ Code typically used by SLPs  ◊ Additional digits not listed here
R41.84 Other specified cognitive deficit
- R41.840 Attention and concentration deficit
- 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
- R44.0 Auditory hallucinations

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

R47 Speech disturbances, not elsewhere classified
- R47.0 Dysphasia and aphasia
  - R47.01 Aphasia
- R47.1 Dysarthria and anarthria
- R47.8 Other speech disturbances
  - 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)
- 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  Aphonia

- Loss of voice

R49.2  Hypernasality and hyponasality

✓ R49.21  Hypernasality

✓ R49.22  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)

- 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 periorcular area
  injuries of scalp
  injuries of temporomandibular joint area
  injuries of tongue
  injuries of tooth

**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

**Excludes1**: diffuse cerebral contusion (S06.2-)
focal cerebral contusion (S06.3-)
injury of eye and orbit (S05.-)
open wound of head (S01.-)

The appropriate 7th character is to be added to each code from category S00
A - initial encounter
D - subsequent encounter
S - sequela

◊ S00.5  Superficial injury of lip and oral cavity

S01  Open wound of head

**Code also** any associated:
injury of cranial nerve (S04.-)
injury of muscle and tendon of head (S09.1-)
intracranial injury (S06.-)
wound infection

**Excludes1**: open skull fracture (S02.- with 7th character B)

**Excludes2**: injury of eye and orbit (S05.-)
traumatic amputation of part of head (S08.-)

The appropriate 7th character is to be added to each code from category S01
A - initial encounter
D - subsequent encounter
S - sequela

◊ S01.5  Open wound of lip and oral cavity

◊ S02  Fracture of skull and facial bones

**Note**: A fracture not indicated as open or closed should be coded to closed

**Code also** any associated intracranial injury (S06.-)

The appropriate 7th character is to be added to each code from category S02
A - initial encounter for closed fracture
B - initial encounter for open fracture
D - subsequent encounter for fracture with routine healing
G - subsequent encounter for fracture with delayed healing
K - subsequent encounter for fracture with nonunion
S - sequela

S06  Intracranial injury

**Includes**: traumatic brain injury

**Code also** any associated:
open wound of head (S01.-)

**Excludes1**: head injury NOS (S09.90)

The appropriate 7th character is to be added to each code from category S06
A - initial encounter
D - subsequent encounter
S - sequela
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

**Note:** A fracture not indicated as displaced or nondisplaced should be coded to displaced.
A fracture not indicated as open or closed should be coded to closed.

**Includes:**
- fracture of cervical neural arch
- fracture of cervical spine
- fracture of cervical spinous process
- fracture of cervical transverse process
- fracture of cervical vertebral arch
- fracture of neck

**Code first** any associated cervical spinal cord injury (S14.0, S14.1-)

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)**

**Effects of foreign body entering through natural orifice (T15-T19)**

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

☑ Code typically used by SLPs  ◊ Additional digits not listed here
T18  Foreign body in alimentary tract
  \textbf{Excludes2:} foreign body in pharynx (T17.2-)
  ♦ T18.1  Foreign body in esophagus

\textit{Poisoning by, adverse effects of and underdosing of drugs, medicaments and biological substances} (T36-T50)

\textbf{Includes:} adverse effect of correct substance properly administered
poisoning by overdose of substance
poisoning by wrong substance given or taken in error
underdosing by (inadvertently) (deliberately) taking less substance than prescribed or instructed

\textbf{Code first,} for adverse effects, the nature of the adverse effect, such as:
adverse effect NOS (T88.7)
aspirin gastritis (K29.-)
blood disorders (D56-D76)
contact dermatitis (L23-L25)
dermatitis due to substances taken internally (L27.-)
nephropathy (N14.0-N14.2)

\textbf{Note:} The drug giving rise to the adverse effect should be identified by use of codes from categories T36-T50 with fifth or sixth character 5.

\textbf{Use additional code(s) to specify:}
manifestations of poisoning
underdosing or failure in dosage during medical and surgical care (Y63.6, Y63.8-Y63.9)
underdosing of medication regimen (Z91.12-, Z91.13-)

\textbf{Excludes1:} toxic reaction to local anesthesia in pregnancy (O29.3-)
\textbf{Excludes2:} abuse and dependence of psychoactive substances (F10-F19)
abuse of non-dependence-producing substances (F55.-)
drug reaction and poisoning affecting newborn (P00-P96)
pathological drug intoxication (inebriation) (F10-F19)

\textit{Toxic effects of substances chiefly nonmedicinal as to source} (T51-T65)

\textbf{Note:} When no intent is indicated code to accidental. Undetermined intent is only for use when there is specific documentation in the record that the intent of the toxic effect cannot be determined.

\textbf{Use additional code(s):}
for all associated manifestations of toxic effect, such as:
respiratory conditions due to external agents (J60-J70)
personal history of foreign body fully removed (Z87.821)
to identify any retained foreign body, if applicable (Z18.-)

\textbf{Excludes1:} contact with and (suspected) exposure to toxic substances (Z77.-)

The appropriate 7th character is to be added to each code from category T51-T65
  A - initial encounter
  D - subsequent encounter
  S – sequela

\textit{Other and unspecified effects of external causes} (T66-T78)

The appropriate 7th character is to be added to each code from category T66-T78
  A - initial encounter
D - subsequent encounter  
S – sequela

**Certain early complications of trauma (T79)**

The appropriate 7th character is to be added to each code from category T79  
A - initial encounter  
D - subsequent encounter  
S – sequela

**Complications of surgical and medical care, not elsewhere classified (T80-T88)**

**Use additional** code for adverse effect, if applicable, to identify drug (T36-T50 with fifth or sixth character 5)  
**Use additional** code(s) to identify the specified condition resulting from the complication  
**Use additional** code to identify devices involved and details of circumstances (Y62-Y82)  
**Excludes2:** any encounters with medical care for postprocedural conditions in which no complications are present, such as:
- artificial opening status (Z93.-)  
- closure of external stoma (Z43.-)  
- fitting and adjustment of external prosthetic device (Z44.-)  
- burns and corrosions from local applications and irradiation (T20-T32)  
- complications of surgical procedures during pregnancy, childbirth and the puerperium (O00-O9A)  
- mechanical complication of respirator [ventilator] (J95.850)  
- poisoning and toxic effects of drugs and chemicals (T36-T65 with fifth or sixth character 1-4 or 6)  
- postprocedural fever (R50.82) specified complications classified elsewhere, such as:  
- cerebrospinal fluid leak from spinal puncture (G97.0)  
- colostomy malfunction (K94.0-) disorders of fluid and electrolyte imbalance (E86-E87) functional disturbances following cardiac surgery (I97.0-I97.1) intraoperative and postprocedural complications of specified body systems (D78.-, E36.-, E89.-, G97.3-, G97.4, H59.3-, H59.-, H95.2-, H95.3, I97.4, I97.5, J95.6-, J95.7, K91.6-, L76.-, M96.-, N99.-)  
- ostomy complications (J95.0-, K94.-, N99.5-) postgastric surgery syndromes (K91.1) postlaminectomy syndrome NEC (M96.1) postmastectomy lymphedema syndrome (I97.2) postsurgical blind-loop syndrome (K91.2) ventilator associated pneumonia (J95.851)

**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

[There is no national requirement for external cause code reporting. Mandatory reporting is subject to state-based mandates or payer requirements. However, providers are encouraged to voluntarily report.]
conditions, codes from Chapter 20 should be used to provide additional information as to the cause of the condition.

**ASHA Note:** External cause codes are not listed in this product due to the volume of available codes in Chapter 20. For a complete listing of codes, see the official ICD-10 list at [www.cdc.gov/nchs/icd/icd10cm.htm](http://www.cdc.gov/nchs/icd/icd10cm.htm).

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

**Note:** Z codes 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

**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

Z01.8 Encounter for other specified special examinations  
Z01.81 Encounter for preprocedural examinations  
Encounter for preoperative examinations
Encounter for radiological and imaging examinations as part of preprocedural examination
Z01.818 Encounter for other preprocedural examination
Encounter for preprocedural examination NOS
Encounter for examinations prior to antineoplastic chemotherapy

Z02 Encounter for administrative examination
Z02.0 Encounter for examination for admission to educational institution
Encounter for examination for admission to preschool (education)
Encounter for examination for re-admission to school following illness or medical treatment
Z02.1 Encounter for pre-employment examination
Z02.2 Encounter for examination for admission to residential institution
**Excludes1:** examination for admission to prison (Z02.89)
Z02.3 Encounter for examination for recruitment to armed forces
Z02.4 Encounter for examination for driving license
Z02.5 Encounter for examination for participation in sport
**Excludes1:** blood-alcohol and blood-drug test (Z02.83)
Z02.6 Encounter for examination for insurance purposes
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
**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)

🔗 Z71 Persons encountering health services for other counseling and medical advice, not elsewhere classifiable
🔗 Z76 Persons encountering health services in other circumstances
🔗 Z76.5 Malingeringer [conscious simulation]
  Person feigning illness (with obvious motivation)
  **Excludes 1:** factitious patient (F68.1-)
  peregrinating patient (F68.1-)
🔗 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
  **Excludes 2:** 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
  **Excludes 2:** 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
  **Excludes 2:** 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

The following instructional notations are from the published ICD-10-CM Tabular List of Diseases and Injuries (www.cdc.gov/nchs/icd/icd10cm.htm).

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. (ASHA note: In other words, never use the listed codes together)

Excludes2 (Can use the listed codes together)
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. (ASHA note: In other words, codes can be listed 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.
**ASHA Note:** Certain ICD-10-CM categories have applicable 7th characters. The seventh character of ICD-10 is often a required character in codes involving, for example, injuries and poisonings (Chapter 19, S00-T88). The purpose of the 7th character is to communicate to the payer the "type of encounter" such as initial (A), subsequent (D), or sequela (S). Any codes requiring a 7th character will be clearly indicated. Not all codes require a 7th character.
ICD-10-PCS (Procedure Coding System) for Speech-Language Pathologists

ICD-10 includes the ICD-10-CM (clinical modification) and ICD-10-PCS (procedure coding system). 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 (ICD-10-PCS) was developed by the Centers for Medicare and Medicaid Services for use in the U.S. for inpatient hospital settings only. ASHA's resources focus mostly on ICD-10-CM. You may want to check with your facility on use of ICD-10-PCS.

The ICD-10-PCS is a procedure classification published by the United States for classifying procedures performed in hospital inpatient health care settings only.

CMS provides the following link for information about ICD-10-PCS; www.cms.gov/Medicare/Coding/ICD10/2017-ICD-10-PCS-and-GEMs.html

The CMS guidelines for ICD-10-PCS, found on the CMS webpage, are a set of rules that have been developed to accompany and complement the official conventions and instructions provided within the ICD-10-PCS itself.

The instructions and conventions of the classification take precedence over guidelines.

The 2017 Code Tables and Index are the actual codes used in ICD-10-PCS. Speech-language pathology and audiology related codes are found in section F - Physical Rehabilitation and Diagnostic Audiology.