

EXPERT

# ICD-10-CM Expert for Hospitals

The complete official code set

Codes valid from October 1, 2021 through September 30, 2022

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#### **External Causes Index**

The External Causes Index is arranged in alphabetic order by main terms that describe the cause, the intent, the place of occurrence, the activity, and the status of the patient at the time the injury occurred or health condition arose.

#### **Index Notations**

#### With

The word "with" or "in" should be interpreted to mean "associated with" or "due to." The classification presumes a causal relationship between the two conditions linked by these terms in the index. These conditions should be coded as related even in the absence of provider documentation explicitly linking them unless the documentation clearly states the conditions are unrelated or when another guideline specifically requires a documented linkage between two conditions (e.g., the sepsis guideline for "acute organ dysfunction that is not clearly associated with the sepsis"). For conditions not specifically linked by these relational terms in the classification or when a guideline requires explicit documentation of a linkage between two conditions, provider documentation must link the conditions to code them as related.

The word "with" in the index is sequenced immediately following the main term, not in alphabetical order.

#### Dermatopolymyositis M33.90

with myopathy M33.92 respiratory involvement M33.91 specified organ involvement NEC M33.99 in neoplastic disease — *see also* Neoplasm D49.9 [*M36.0*]

#### See

When the instruction "see" follows a term in the index, it indicates that another term must be referenced to locate the correct code.

Hematoperitoneum — see Hemoperitoneum

#### See Also

The instructional note "see also" simply provides alternative terms the coder may reference that may be useful in determining the correct code but are not necessary to follow if the main term supplies the appropriate code.

Hematinuria — see also Hemaglobinuria malarial B50.8

#### **Default Codes**

In the index, the default code is the code listed next to the main term and represents the condition most commonly associated with that main term. This code may be assigned when documentation does not support reporting a more specific code. Alternatively, it may provide an unspecified code for the condition.

> Hemiatrophy R68.89 cerebellar G31.9 face, facial, progressive (Romberg) G51.8 tongue K14.8

#### Parentheses

Parentheses in the indexes enclose nonessential modifiers, supplementary words that may be present or absent in the statement of a disease without affecting the code.

> Pseudomeningocele (cerebral) (infective) (post-traumatic) G96.198 postprocedural (spinal) G97.82

L\_\_\_\_\_

#### Brackets

ICD-10-CM has a coding convention addressing code assignment for manifestations that occur as a result of an underlying condition. This convention requires the underlying condition to be sequenced first, followed by the code or codes for the associated manifestation. In the index, italicized codes in brackets identify manifestation codes.

> Polyneuropathy (peripheral) G62.9 alcoholic G62.1 amyloid (Portuguese) E85.1 [G63] transthyretin-related (ATTR) familial E85.1 [G63]

#### Shaded Guides

Exclusive vertical shaded guides in the Index to Diseases and Injuries and External Causes Index help the user easily follow the indent levels for the subentries under a main term. Sequencing rules may apply depending on the level of indent for separate subentries.

#### Hemicrania

congenital malformation Q00.0
continua G44.51
meaning migraine — see also Migraine G43.909
paroxysmal G44.039
chronic G44.Ø49
intractable G44.Ø41
not intractable G44.049
episodic G44.039
intractable G44.031
not intractable G44.039
intractable G44.031
not intractable G44.039

#### Following References

The Index to Diseases and Injuries includes following references to assist in locating out-of-sequence codes in the tabular list. Out-of-sequence codes contain an alphabetic character (letter) in the third- or fourth-character position. These codes are placed according to the classification rules — according to condition — not according to alphabetic or numeric sequencing rules.

> Carcinoma (malignant) — see also Neoplasm, by site, malignant neuroendocrine — see also Tumor, neuroendocrine high grade, any site C7A.1 (following C75) poorly differentiated, any site C7A.1 (following C75)

#### **Additional Character Required**

The Index to Diseases and Injuries, Neoplasm Table, and External Causes Index provide an icon after certain codes to signify to the user that additional characters are required to make the code valid. The tabular list should be consulted for appropriate character selection.

> Fall, falling (accidental) W19 ☑ building W20.1 ☑

## Tabular List of Diseases

ICD-10-CM codes and descriptions are arranged numerically within the tabular list of diseases with 19 separate chapters providing codes associated with a particular body system or nature of injury or disease. There is also a chapter providing codes for external causes of an injury or health conditions, a chapter for codes that address encounters with healthcare facilities for circumstances other than a disease or injury, and finally a chapter for codes that capture special circumstances such as new diseases of uncertain etiology or emergency use codes.

#### Code and Code Descriptions

ICD-10-CM is an alphanumeric classification system that contains categories, subcategories, and valid codes. The first character is always a letter with any additional characters represented by either a letter or number. A three-character category without further subclassification is equivalent to a valid three-character code. Valid codes may be three, four, five, six, or seven characters in length, with each level of subdivision after a three-character category representing a subcategory. The final level of subdivision is a valid code.

#### Boldface

Boldface type is used for all codes and descriptions in the tabular list.

#### Italics

Italicized type is used to identify manifestation codes, those codes that should not be reported as first-listed diagnoses.

#### Deleted Text

Strikethrough on a code and code description indicates a deletion from the classification for the current year.

#### Key Word

Green font is used throughout the Tabular List of Diseases to differentiate the key words that appear in similar code descriptions in a given category or subcategory. The key word convention is used only in those categories in which there are multiple codes with very similar descriptions with only a few words that differentiate them.

For example, refer to the list of codes below from category H55:

4th	H55	Nystagmus and other irregular eye movements		
	$\sqrt{5}^{th}$	H55.Ø	Nystagn	nus
			H55.ØØ	Unspecified nystagmus
			H55.Ø1	Congenital nystagmus
			H55.Ø2	Latent nystagmus
			H55.Ø3	Visual deprivation nystagmus
			H55.Ø4	Dissociated nystagmus
			H55.Ø9	Other forms of nystagmus

The portion of the code description that appears in **green font** in the tabular list helps the coder quickly identify the key terms and the correct code. This convention is especially useful when the codes describe laterality, such as the following codes from subcategory H40.22:

#### H40.22 Chronic angle-closure glaucoma

- Chronic primary angle-closure glaucoma √7<sup>th</sup> H4Ø.221 Chronic angle-closure glaucoma, right eye
- √7<sup>th</sup> H4Ø.222 Chronic angle-closure glaucoma, left eye
- H40.223 Chronic angle-closure glaucoma, bilateral
- H40.229 Chronic angle-closure glaucoma, unspecified
  - eye

#### **Tabular Notations**

Official parenthetical notes as well as Optum360's supplementary notations are provided at the chapter, code block, category, subcategory, and individual code level to help the user assign proper codes. The information in the notation can apply to one or more codes depending on where the citation is placed.

#### **Official Notations**

#### **Includes Notes**

The word **INCLUDES** appears immediately under certain categories to further define, clarify, or give examples of the content of a code category.

#### Inclusion Terms

Lists of inclusion terms are included under certain codes. These terms indicate some of the conditions for which that code number may be used. Inclusion terms may be synonyms with the code title, or, in the case of "other specified" codes, the terms may also provide a list of various conditions included within a classification code. The inclusion terms are not exhaustive. The index may provide additional terms that may also be assigned to a given code.

#### **Excludes Notes**

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

#### **Excludes 1**

An **EXCLUDES 1** note is a "pure" excludes. It means "NOT CODED HERE!" An Excludes 1 note indicates mutually exclusive codes: two conditions that cannot be reported together. 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.

An exception to the Excludes 1 definition is when the two conditions are unrelated to each other. If it is not clear whether the two conditions involving an Excludes 1 note are related or not, guery the provider. For example, code F45.8 Other somatoform disorders, has an Excludes 1 note for "sleep related teeth grinding (G47.63)" because "teeth grinding" is an inclusion term under F45.8. Only one of these two codes should be assigned for teeth grinding. However, psychogenic dysmenorrhea is also an inclusion term under F45.8, and a patient could have both this condition and sleep-related teeth grinding. In this case, the two conditions are clearly unrelated to each other, so it would be appropriate to report F45.8 and G47.63 together.

#### **Excludes 2**

An **EXCLUDES 2** note means "NOT INCLUDED HERE." An Excludes 2 note indicates that although the excluded condition is not part of the condition it is excluded from, a patient may have both conditions at the same time. Therefore, when an Excludes 2 note appears under a code, it may be acceptable to use both the code and the excluded code together if supported by the medical documentation.

#### Note

The term "NOTE" appears as an icon and precedes the instructional information. These notes function as alerts to highlight coding instructions within the text.

#### Code First/Use additional code

These instructional notes provide sequencing instruction. They may appear independently of each other or to designate certain etiology/manifestation paired codes. These instructions signal the coder that an additional code should be reported to provide a more complete picture of that diagnosis.

In etiology/manifestation coding, ICD-10-CM requires the underlying condition to be sequenced first, followed by the manifestation. In these situations, codes with "In diseases classified elsewhere" in the code description are never permitted as a first-listed or principal diagnosis code and must be sequenced following the underlying condition code.

#### What's New for 2022

What's New for 2021

D72.118	Other hypereosinophilic syndrome	
D72.119	Hypereosinophilic syndrome [HES], unspecified	
D72.12	Drug rash with eosinophilia and systemic symptoms syndrome	
D72.18	Eosinophilia in diseases classified elsewhere	
D72.19	Other eosinophilia	
D84.81	Immunodeficiency due to conditions classified elsewhere	
D84.821	Immunodeficiency due to drugs	
D84.822	Immunodeficiency due to external causes	
D84.89	Other immunodeficiencies	
D89.831	Cytokine release syndrome, grade 1	
D89.832	Cytokine release syndrome, grade 2	
D89.833	Cytokine release syndrome, grade 3	
D89.834	Cytokine release syndrome, grade 4	
D89.835	Cytokine release syndrome, grade 5	
D89.839	Cytokine release syndrome, grade unspecified	
E7Ø.81	Aromatic L-amino acid decarboxylase deficiency	
E7Ø.89	Other disorders of aromatic amino-acid metabolism	
E74.81Ø	Glucose transporter protein type 1 deficiency	
E74.818	Other disorders of glucose transport	
E74.819	Disorders of glucose transport, unspecified	
E74.89	Other specified disorders of carbohydrate metabolism	
F1Ø.13Ø	Alcohol abuse with withdrawal, uncomplicated	
F1Ø.131	Alcohol abuse with withdrawal delirium	
F1Ø.132	Alcohol abuse with withdrawal with perceptual disturbance	
F1Ø.139	Alcohol abuse with withdrawal, unspecified	
F1Ø.93Ø	Alcohol use, unspecified with withdrawal, uncomplicated	
F1Ø.931	Alcohol use, unspecified with withdrawal delirium	
F1Ø.932	Alcohol use, unspecified with withdrawal with perceptual disturbance	
F1Ø.939	Alcohol use, unspecified with withdrawal, unspecified	
F11.13	Opioid abuse with withdrawal	
F12.13	Cannabis abuse with withdrawal	
F13.13Ø	Sedative, hypnotic or anxiolytic abuse with withdrawal, uncomplicated	
F13.131	Sedative, hypnotic or anxiolytic abuse with withdrawal delirium	
F13.132	Sedative, hypnotic or anxiolytic abuse with withdrawal with perceptual disturbance	
F13.139	Sedative, hypnotic or anxiolytic abuse with withdrawal, unspecified	
F14.13	Cocaine abuse, unspecified with withdrawal	
F14.93	Cocaine use, unspecified with withdrawal	
F15.13	Other stimulant abuse with withdrawal	
F19.13Ø	Other psychoactive substance abuse with withdrawal, uncomplicated	
F19.131	Other psychoactive substance abuse with withdrawal delirium	
F19.132	Other psychoactive substance abuse with withdrawal with perceptual disturbance	
F19.139	Other psychoactive substance abuse with withdrawal, unspecified	
G11.1Ø	Early-onset cerebellar ataxia, unspecified	
G11.11	Friedreich ataxia	

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G11.19	Other early-onset cerebellar ataxia
G4Ø.42	Cyclin-Dependent Kinase-Like 5 Deficiency Disorder
G4Ø.833	Dravet syndrome, intractable, with status epilepticus
G4Ø.834	Dravet syndrome, intractable, without status epilepticus
G71.2Ø	Congenital myopathy, unspecified
G71.21	Nemaline myopathy
G71.22Ø	X-linked myotubular myopathy
G71.228	Other centronuclear myopathy
G71.29	Other congenital myopathy
G96.ØØ	Cerebrospinal fluid leak, unspecified
G96.Ø1	Cranial cerebrospinal fluid leak, spontaneous
G96.Ø2	Spinal cerebrospinal fluid leak, spontaneous
G96.Ø8	Other cranial cerebrospinal fluid leak
G96.Ø9	Other spinal cerebrospinal fluid leak
G96.191	Perineural cyst
G96.198	Other disorders of meninges, not elsewhere classified
G96.81Ø	Intracranial hypotension, unspecified
G96.811	Intracranial hypotension, spontaneous
G96.819	Other intracranial hypotension
G96.89	Other specified disorders of central nervous system
G97.83	Intracranial hypotension following lumbar cerebrospinal fluid shunting
G97.84	Intracranial hypotension following other procedure
H18.5Ø1	Unspecified hereditary corneal dystrophies, right eye
H18.5Ø2	Unspecified hereditary corneal dystrophies, left eye
H18.5Ø3	Unspecified hereditary corneal dystrophies, bilateral
H18.5Ø9	Unspecified hereditary corneal dystrophies, unspecified
1110.505	eye
H18.511	Endothelial corneal dystrophy, right eye
H18.512	Endothelial corneal dystrophy, left eye
H18.513	Endothelial corneal dystrophy, bilateral
H18.519	Endothelial corneal dystrophy, unspecified eye
H18.521	Epithelial (juvenile) corneal dystrophy, right eye
H18.522	Epithelial (juvenile) corneal dystrophy, left eye
H18.523	Epithelial (juvenile) corneal dystrophy, bilateral
H18.529	Epithelial (juvenile) corneal dystrophy, unspecified eye
H18.531	Granular corneal dystrophy, right eye
H18.532	Granular corneal dystrophy, left eye
H18.533	Granular corneal dystrophy, bilateral
H18.539	Granular corneal dystrophy, unspecified eye
H18.541	Lattice corneal dystrophy, right eye
H18.542	Lattice corneal dystrophy, left eye
H18.543	Lattice corneal dystrophy, bilateral
H18.549	Lattice corneal dystrophy, unspecified eye
H18.551	Macular corneal dystrophy, right eye
H18.552	Macular corneal dystrophy, left eye
H18.553	Macular corneal dystrophy, bilateral
H18.559	Macular corneal dystrophy, unspecified eye
H18.591	Other hereditary corneal dystrophies, right eye
H18.592	Other hereditary corneal dystrophies, left eye
H18.593	Other hereditary corneal dystrophies, bilateral
H18.599	Other hereditary corneal dystrophies, unspecified eye
H55.82	Deficient smooth pursuit eye movements
	. ,

Admission — continued

examination at health care facility — see also Examination — continued vision — continued infant or child (over 28 days old) ZØØ.129 with abnormal findings ZØØ.121 fitting (of) artificial arm — see Admission, adjustment, artificial, arm eye Z44.2 🗹 leg — see Admission, adjustment, artificial, leg brain neuropacemaker Z46.2 implanted Z45.42 breast prosthesis (external) Z44.3 colostomy belt Z46.89 contact lenses Z46.Ø cystostomy device Z46.6 dental prosthesis Z46.3 dentures Z46.3 device NEC abdominal Z46.89 nervous system Z46.2 implanted — see Admission, adjustment, device, implanted, nervous system orthodontic Z46.4 prosthetic Z44.9 breast Z44.3 🗹 dental Z46.3 eye Z44.2 🗹 substitution auditory Z46.2 implanted — see Admission, adjustment, device, implanted, hearing device nervous system Z46.2 implanted — see Admission, adjustment, device, implanted, nervous system visual Z46.2 implanted Z45.31 hearing aid Z46.1 ileostomy device Z46.89 intestinal appliance or device NEC Z46.89 neuropacemaker (brain) (peripheral nerve) (spinal cord) Z46.2 implanted Z45.42 orthodontic device Z46.4 orthopedic device (brace) (cast) (shoes) Z46.89 prosthesis Z44.9 arm — see Admission, adjustment, artificial, arm breast Z44.3 🗹 dental Z46.3 eye Z44.2 🗹 leg — see Admission, adjustment, artificial, leg specified type NEC Z44.8 spectacles Z46.Ø follow-up examination ZØ9 intrauterine device management Z30.431 initial prescription Z30.014 mental health evaluation ZØØ.8 requested by authority ZØ4.6 observation — see Observation Papanicolaou smear, cervix Z12.4 for suspected malignant neoplasm Z12.4 plastic and reconstructive surgery following medical procedure or healed injury NEC Z42.8 plastic surgery, cosmetic NEC Z41.1 postpartum observation immediately after delivery Z39.0 routine follow-up Z39.2 poststerilization (for restoration) Z31.0 aftercare Z31.42 procreative management Z31.9 prophylactic (measure) — see also Encounter, prophylactic measures organ removal Z40.00 breast Z4Ø.Ø1 fallopian tube(s) Z4Ø.Ø3 with ovary(s) Z4Ø.Ø2 ovary(s) Z40.02 specified organ NEC Z40.09 testes Z40.09 vaccination Z23 psychiatric examination (general) ZØØ.8 requested by authority ZØ4.6 radiation therapy (antineoplastic) Z51.Ø reconstructive surgery following medical procedure or healed injury NEC Z42.8 removal of cystostomy catheter Z43.5 Aftercare — see also Care Z51.89

Admission — continued removal of — continued drains Z48.03 dressing (nonsurgical) Z48.ØØ implantable subdermal contraceptive Z3Ø.46 intrauterine contraceptive device Z30.432 neuropacemaker (brain) (peripheral nerve) (spinal . cord) Z46.2 implanted Z45.42 staples Z48.02 surgical dressing Z48.01 sutures Z48.Ø2 ureteral stent Z46.6 respirator [ventilator] use during power failure Z99.12 restoration of organ continuity (poststerilization) Z31.0 aftercare Z31.42 sensitivity test — *see also* Test, skin allergy NEC ZØ1.82 Mantoux Z11.1 tuboplasty following previous sterilization Z31.Ø aftercare Z31.42 vasoplasty following previous sterilization Z31.Ø aftercare Z31.42 vision examination ZØ1.ØØ with abnormal findings ZØ1.Ø1 following failed vision screening ZØ1.020 with abnormal findings ZØ1.020 infant or child (over 28 days old) ZØ0.129 with abnormal findings ZØØ.121 waiting period for admission to other facility Z75.1 Adnexitis (suppurative) — see Salpingo-oophoritis Adolescent X-linked adrenoleukodystrophy E71.521 Adrenal (gland) — see condition Adrenalism, tuberculous A18.7 Adrenalitis, adrenitis E27.8 autoimmune E27.1 meningococcal, hemorrhagic A39.1 Adrenarche, premature E27.0 Adrenocortical syndrome — see Cushing's, syndrome Adrenogenital syndrome E25.9 acquired E25 congenital E25.Ø salt loss E25.Ø Adrenogenitalism, congenital E25.0 Adrenoleukodystrophy E71.529 eonatal E71.511 X-linked E71.529 Addison only phenotype E71.528 Addison-Schilder E71.528 adolescent E71.521 adrenomyeloneuropathy E71.522 childhood cerebral E71.520 other specified E71.528 Adrenomyeloneuropathy E71.522 Adventitious bursa — see Bursopathy, specified type Adverse effect — see Table of Drugs and Chemicals, categories T36-T50, with 6th character 5 Advice — see Counseling Adynamia (episodica) (hereditary) (periodic) G72.3 Aeration lung imperfect, newborn — see Atelectasis Aerobullosis T70.3 Aerocele — see Embolism, air Aerodermectasia subcutaneous (traumatic) T79.7 🗹 Aerodontalgia T70.29 🗹 Aeroembolism T7Ø.3 Aerogenes capsulatus infection A48.0 Aero-otitis media T70.0 🗹 Aerophagy, aerophagia (psychogenic) F45.8 Aerophobia F4Ø.228 Aerosinusitis T70.1 Aerotitis T70.0 🗹 Affection — see Disease Afibrinogenemia — see also Defect, coagulation D68.8 acquired D65 congenital D68.2 following ectopic or molar pregnancy OØ8.1 in abortion — see Abortion, by type, complicated by, afibrinogenemia puerperal 072.3 African sleeping sickness B56.9 tick fever A68.1 trypanosomiasis B56.9 gambian B56.Ø rhodesian B56.1

Aftercare — continued

following surgery (for) (on) amputation Z47.81 attention to drains Z48.Ø3 dressings (nonsurgical) Z48.00 surgical Z48.01 sutures Z48.Ø2 circulatory system Z48.812 delayed (planned) wound closure Z48.1 digestive system Z48.815 explantation of joint prosthesis (staged procedure) hip Z47.32 knee Z47.33 shoulder Z47.31 genitourinary system Z48.816 joint replacement Z47.1 neoplasm Z48.3 nervous system Z48.811 oral cavity Z48.814 organ transplant bone marrow Z48.290 heart Z48.21 heart-lung Z48.28Ø kidney Z48.22 liver Ź48.23 nver 246.23 lung Z48.24 multiple organs NEC Z48.288 specified NEC Z48.298 orthopedic NEC Z47.89 planned wound closure Z48.1 removal of internal fixation device Z47.2 respiratory system Z48.813 scoliosis Z47.82 sense organs Z48.81Ø skin and subcutaneous tissue Z48.817 specified body system circulatory Z48.812 digestive Z48.815 genitourinary Z48.816 nervous Z48.811 oral cavity Z48.814 respiratory Z48.813 sense organs Z48.81Ø skin and subcutaneous tissue Z48.817 teeth Z48.814 specified NEC Z48.89 spinal Z47.89 teeth Z48.814 fracture — code to fracture with seventh character D involving removal of drains Z48.Ø3 dressings (nonsurgical) Z48.00 staples Z48.02 surgical dressings Z48.Ø1 sutures Z48.Ø2 neuropacemaker (brain) (peripheral nerve) (spinal cord) Z46.2 implanted Z45.42 orthopedic NEC Z47.89 postprocedural — see Aftercare, following surgery After-cataract — see Cataract, secondary Agalactia (primary) 092.3 elective, secondary or therapeutic 092.5 Agammaglobulinemia (acquired (secondary) (nonfamilial) D8Ø.1 with immunoglobulin-bearing B-lymphocytes D80.1 lymphopenia D81.9 autosomal recessive (Swiss type) D8Ø.Ø Bruton's X-linked D80.0 common variable (CVAgamma) D8Ø.1 congenital sex-linked D80.0 hereditary D80.0 lymphopenic D81.9 Świss type (autosomal recessive) D8Ø.Ø X-linked (with growth hormone deficiency) (Bruton) Aganglionosis (bowel) (colon) Q43.1 Age (old) — see Senility Agenesis adrenal (gland) Q89.1 alimentary tract (complete) (partial) NEC Q45.8 upper Q4Ø.8 anus, anal (canal) Q42.3 with fistula Q42.2 aorta Q25.41

<u> Admission — Agenesis</u>

## ICD-10-CM Tabular List of Diseases and Injuries

## Chapter 1. Certain Infectious and Parasitic Diseases (AØØ–B99), UØ7.1

#### **Chapter-specific Guidelines with Coding Examples**

The chapter-specific guidelines from the ICD-10-CM Official Guidelines for Coding and Reporting have been provided below. Along with these guidelines are coding examples, contained in the shaded boxes, that have been developed to help illustrate the coding and/or sequencing guidance found in these guidelines.

#### a. Human immunodeficiency virus (HIV) infections

#### 1) Code only confirmed cases

Code only confirmed cases of HIV infection/illness. This is an exception to the hospital inpatient guideline Section II, H.

In this context, "confirmation" does not require documentation of positive serology or culture for HIV; the provider's diagnostic statement that the patient is HIV positive, or has an HIV-related illness is sufficient.

Patient admitted with anemia with possible HIV infection

#### D64.9 Anemia, unspecified

*Explanation:* Only the anemia is coded in this scenario because it has not been confirmed that an HIV infection is present. This is an exception to the guideline Section II, H for hospital inpatient coding.

#### 2) Selection and sequencing of HIV codes

#### (a) Patient admitted for HIV-related condition

If a patient is admitted for an HIV-related condition, the principal diagnosis should be B2Ø, Human immunodeficiency virus [HIV] disease followed by additional diagnosis codes for all reported HIV-related conditions.

#### (b) Patient with HIV disease admitted for unrelated condition

If a patient with HIV disease is admitted for an unrelated condition (such as a traumatic injury), the code for the unrelated condition (e.g., the nature of injury code) should be the principal diagnosis. Other diagnoses would be B2Ø followed by additional diagnosis codes for all reported HIV-related conditions.

Unstable angina, native coronary artery atherosclerosis, HIV

## I25.110 Atherosclerotic heart disease of native coronary artery with unstable angina pectoris

#### B20 Human immunodeficiency virus [HIV] disease

*Explanation*: The arteriosclerotic coronary artery disease and the unstable angina are not related to HIV, so those conditions are reported first using a combination code, and HIV is reported secondarily.

#### (c) Whether the patient is newly diagnosed

Whether the patient is newly diagnosed or has had previous admissions/encounters for HIV conditions is irrelevant to the sequencing decision.

#### (d) Asymptomatic human immunodeficiency virus

Z21, Asymptomatic human immunodeficiency virus [HIV] infection status, is to be applied when the patient without any documentation of symptoms is listed as being "HIV positive," "known HIV," "HIV test positive," or similar terminology. Do not use this code if the term "AIDS" is used or if the patient is treated for any HIV-related illness or is described as having any condition(s) resulting from his/her HIV positive status; use B20 in these cases.

Patient admitted with acute appendicitis. Status positive HIV test on Atripla, with no prior symptoms

#### K35.8Ø Unspecified acute appendicitis

#### Z21 Asymptomatic human immunodeficiency virus [HIV] infection status

*Explanation:* Code Z21 is sequenced second since documentation indicates that the patient has had a positive HIV test but has been asymptomatic. Being on medication for HIV is not an indication that code B2Ø is used instead of Z21. Unless there has been documentation that the patient has had current or prior symptoms or HIV-related complications, code B2Ø is not used. The appendicitis is not an AIDS-related complication and is sequenced first.

#### (e) Patients with inconclusive HIV serology

Patients with inconclusive HIV serology, but no definitive diagnosis or manifestations of the illness, may be assigned code R75, Inconclusive laboratory evidence of human immunodeficiency virus [HIV].

#### (f) Previously diagnosed HIV-related illness

Patients with any known prior diagnosis of an HIV-related illness should be coded to B20. Once a patient has developed an HIV-related illness, the patient should always be assigned code B20 on every subsequent admission/encounter. Patients previously diagnosed with any HIV illness (B20) should never be assigned to R75 or Z21, Asymptomatic human immunodeficiency virus [HIV] infection status.

#### (g) HIV infection in pregnancy, childbirth and the puerperium

During pregnancy, childbirth or the puerperium, a patient admitted (or presenting for a health care encounter) because of an HIV-related illness should receive a principal diagnosis code of O98.7-, Human immunodeficiency [HIV] disease complicating pregnancy, childbirth and the puerperium, followed by B20 and the code(s) for the HIVrelated illness(es). Codes from Chapter 15 always take sequencing priority.

Patients with asymptomatic HIV infection status admitted (or presenting for a health care encounter) during pregnancy, childbirth, or the puerperium should receive codes of O98.7- and Z21.

#### (h) Encounters for testing for HIV

If a patient is being seen to determine his/her HIV status, use code Z11.4, Encounter for screening for human immunodeficiency virus [HIV]. Use additional codes for any associated high-risk behavior.

If a patient with signs or symptoms is being seen for HIV testing, code the signs and symptoms. An additional counseling code Z71.7, Human immunodeficiency virus [HIV] counseling, may be used if counseling is provided during the encounter for the test.

When a patient returns to be informed of his/her HIV test results and the test result is negative, use code Z71.7, Human immunodeficiency virus [HIV] counseling.

If the results are positive, see previous guidelines and assign codes as appropriate.

#### b. Infectious agents as the cause of diseases classified to other chapters

Certain infections are classified in chapters other than Chapter 1 and no organism is identified as part of the infection code. In these instances, it is necessary to use an additional code from Chapter 1 to identify the organism. A code from category B95, Streptococcus, Staphylococcus, and Enterococcus as the cause of diseases classified to other chapters, B96, Other bacterial agents as the cause of diseases classified to other chapters, or B97, Viral agents as the cause of diseases classified to other chapters, is to be used as an additional code to identify the organism. An instructional note will be found at the infection code advising that an additional organism code is required.

#### E. coli UTI

N39.Ø Urinary tract infection, site not specified

## B96.20 Unspecified Escherichia coli [E.coli] as the cause of diseases classified elsewhere

*Explanation:* An instructional note under the code for the urinary tract infection indicates to code also the specific organism.

#### c. Infections resistant to antibiotics

Many bacterial infections are resistant to current antibiotics. It is necessary to identify all infections documented as antibiotic resistant. Assign a code from category Z16, Resistance to antimicrobial drugs, following the infection code only if the infection code does not identify drug resistance.

#### d. Sepsis, severe sepsis, and septic shock

#### 1) Coding of sepsis and severe sepsis (a) Sepsis

#### (a) Seps

For a diagnosis of sepsis, assign the appropriate code for the underlying systemic infection. If the type of infection or causal organism is not further specified, assign code A41.9, Sepsis, unspecified organism.

A code from subcategory R65.2, Severe sepsis, should not be assigned unless severe sepsis or an associated acute organ dysfunction is documented.

#### Chapter 9. Diseases of the Circulatory System

**Chapter 9. Diseases of the Circulatory System** 

	Chapter 5. Diseases of	the ch	culat	ULA DI A	otem 199-197.9
Chapter 9	9. Diseases of the Circulatory System (IØØ-I99)				hronic rheumatic heart diseases (105-109)
EXCLUDES					monic medinatic near (diseases (105-109)
	certain infectious and parasitic diseases (AØ0-B99) complications of pregnancy, childbirth and the puerperium (OØ0-O9A) congenital malformations, deformations, and chromosomal abnormalities (QØ0-Q99)	√4 <sup>th</sup>	1Ø5	INCLU	whether specified as rheumatic or not
	endocrine, nutritional and metabolic diseases (EØØ-E88) injury, poisoning and certain other consequences of external causes (SØØ-T88)				mitral valve disease specified as nonrheumatic (134) mitral valve disease with aortic and/or tricuspid valve involvement (108)
	neoplasms (CØØ-D49) symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified (RØØ-R94) systemic connective tissue disorders (M3Ø-M36) transient cerebral ischemic attacks and related syndromes (G45)			105.0	Rheumatic mitral stenosis Mitral (valve) obstruction (rheumatic) DEF: Narrowing of the mitral valve between the left atrium and left ventricle due to rheumatic fever. Symptoms include shortness of breath during or after exercise, fatigue, palpitations, chest
This cha	pter contains the following blocks:				discomfort, and swelling of feet or legs.
100-102	Acute rheumatic fever			IØ5.1	Rheumatic mitral insufficiency
105-109 110-116 120-125	Chronic rheumatic heart diseases Hypertensive diseases Ischemic heart diseases				Rheumatic mitral incompetence           Rheumatic mitral regurgitation           EXGLUDEST
126-128 13Ø-152	Pulmonary heart disease and diseases of pulmonary circulation Other forms of heart disease			lø5.2	Rheumatic mitral stenosis with insufficiency Rheumatic mitral stenosis with incompetence or regurgitation
160-169 170-179 180-189	Cerebrovascular diseases Diseases of arteries, arterioles and capillaries Diseases of veins, lymphatic vessels and lymph nodes, not elsewhere			lø5.8	Other rheumatic mitral valve diseases Rheumatic mitral (valve) failure
195-199	classified Other and unspecified disorders of the circulatory system			lø5.9	Rheumatic mitral valve disease, unspecified Rheumatic mitral (valve) disorder (chronic) NOS
	Acute rheumatic fever (IØØ-IØ2)	√4 <sup>th</sup>	<b>IØ6</b>	Rheun	natic aortic valve diseases
streptoc	lammatory disease that can follow a throat infection by group A occi. Complications can involve the joints (arthritis), subcutaneous tissue s), skin (erythema marginatum), heart (carditis), or brain (chorea).			TEXCEL	DEST aortic valve disease not specified as rheumatic (135) aortic valve disease with mitral and/or tricuspid valve involvement (108)
	heumatic fever without heart involvement			IØ6.Ø	Rheumatic aortic stenosis
	[INCLUDES] arthritis, rheumatic, acute or subacute			1000	Rheumatic aortic (valve) obstruction
	<b>EXCLUDES 1</b> rheumatic fever with heart involvement (101.0-101.9)			106.1	Rheumatic aortic insufficiency Rheumatic aortic incompetence
<mark>√4"</mark> IØ1 R	heumatic fever with heart involvement				Rheumatic aortic regurgitation
	<b>EXCLUDEST</b> chronic diseases of rheumatic origin (105-109) unless rheumatic fever is also present or there is evidence of reactivation are the interaction of the chrome the reactivation				Rheumatic aortic stenosis with insufficiency Rheumatic aortic stenosis with incompetence or regurgitation
I	or activity of the rheumatic process <b>01.0 Acute rheumatic pericarditis</b>				Other rheumatic aortic valve diseases
	Any condition in IØØ with pericarditis			100.9	Rheumatic aortic valve disease, unspecified Rheumatic aortic (valve) disease NOS
	Rheumatic pericarditis (acute)	√4 <sup>th</sup>	<b>IØ7</b>	Rheun	natic tricuspid valve diseases
	EXCLUDES 1 acute pericarditis not specified as rheumatic (130) 01.1 Acute rheumatic endocarditis			INCLU	
13	Ø1.1 Acute rheumatic endocarditis Any condition in IØØ with endocarditis or valvulitis Acute rheumatic valvulitis			EXCLU	or unspecified <u>DES 1</u> tricuspid valve disease specified as nonrheumatic (136)
li	Ø1.2         Acute rheumatic myocarditis         Image: Comparison of the myocarditis           Any condition in 100 with myocarditis         Image: Comparison of the myocarditis         Image: Comparison of the myocarditis			107.0	tricuspid valve disease with aortic and/or mitral valve involvement (108)
10	01.8 Other acute rheumatic heart disease			0.701	Rheumatic tricuspid stenosis Tricuspid (valve) stenosis (rheumatic)
	Any condition in IØØ with other or multiple types of heart involvement Acute rheumatic pancarditis			lø7.1	Rheumatic tricuspid insufficiency Tricuspid (valve) insufficiency (rheumatic)
10	81.9 Acute rheumatic heart disease, unspecified				Rheumatic tricuspid stenosis and insufficiency
	Any condition in IØØ with unspecified type of heart involvement				Other rheumatic tricuspid valve diseases Rheumatic tricuspid valve disease, unspecified
	Rheumatic carditis, acute Rheumatic heart disease, active or acute				Rheumatic tricuspid valve disorder NOS
<mark>√4ª</mark> 1ø2 R	heumatic chorea	√4 <sup>th</sup>	108		ble valve diseases
	[INCLUDES] Sydenham's chorea			INCLU	JUES multiple valve diseases specified as rheumatic or unspecified
	EXCLUDES 1 chorea NOS (G25.5) Huntington's chorea (G1Ø)			EXCLU	DES 1 endocarditis, valve unspecified (138)
10	<b>02.0</b> Rheumatic chorea with heart involvement         CC           Chorea NOS with heart involvement         CC				multiple valve disease specified a nonrheumatic (134, 135, 136, 137, 138, Q22, Q23, Q24.8-)
	Rheumatic chorea with heart involvement of any type			108.0	rheumatic valve disease NOS (109.1) Rheumatic disorders of both mitral and aortic valves
10	classifiable under IØ1 <b>Ø2.9 Rheumatic chorea without heart involvement</b>				Involvement of both mitral and aortic valves specified as rheumatic or unspecified
	Kileuniauc chorea NOS			100 0	<b>AHA:</b> 2019,2Q,5
				IØ8.2	Rheumatic disorders of both mitral and tricuspid valves Rheumatic disorders of both aortic and tricuspid valves Combined rheumatic disorders of mitral, aortic and tricuspid
					valves
					Other rheumatic multiple valve diseases Rheumatic multiple valve disease, unspecified
		√4th	Ião		rheumatic heart diseases
		V 4**	107		Rheumatic myocarditis
					<b>EXCLUDES 1</b> myocarditis not specified as rheumatic (151.4)

0.601-001

#### Chapter 11. Diseases of the Digestive System

Chapter 11. Diseases of the Digestive System (KØØ-K95)	KØØ.5 Hereditary disturbances in tooth structure, not elsewhere
EXCHUDUSSE2       certain conditions originating in the perinatal period (PØ4-P96)         certain infectious and parasitic diseases (AØ0-B99)       complications of pregnancy, childbirth and the puerperium (OØ0-O9A)         congenital malformations, deformations and chromosomal abnormalities (QØ0-Q99)       endocrine, nutritional and metabolic diseases (EØ0-E88)         injury, poisoning and certain other consequences of external causes (SØ0-T88)       neoplasms (CØ0-D49)         symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified (RØ0-R94)       content of the consequences of external causes	classified Amelogenesis imperfecta Dentinogenesis imperfecta Odontogenesis imperfecta Dentinal dysplasia Shell teeth KØØ.6 Disturbances in tooth eruption Dentia praecox Natal tooth Neonatal tooth Bermeture gruntion of tooth
This chapter contains the following blocks:	Premature eruption of tooth Premature shedding of primary [deciduous] tooth
KØØ-K14 Diseases of oral cavity and salivary glands	Prenatal teeth
K2Ø-K31 Diseases of esophagus, stomach and duodenum K35-K38 Diseases of appendix	Retained [persistent] primary tooth
K4Ø-K46 Hernia	KØØ.7 Teething syndrome
K50-K52 Noninfective enteritis and colitis K55-K64 Other diseases of intestines	KØØ.8 Other disorders of tooth development
K65-K68 Diseases of peritoneum and retroperitoneum K70-K77 Diseases of liver	Color changes during tooth formation Intrinsic staining of teeth NOS
K8Ø-K87 Disorders of gallbladder, biliary tract and pancreas	EXCLUDES 2 posteruptive color changes (KØ3.7)
K9Ø-K95 Other diseases of the digestive system	KØØ.9 Disorder of tooth development, unspecified
Diseases of oral cavity and salivary glands (KØØ-K14)	Disorder of odontogenesis NOS KØ1 Embedded and impacted teeth
🜠 KØØ Disorders of tooth development and eruption	EXCLUDES 1 abnormal position of fully erupted teeth (M26.3-)
EXCLUDES 2 embedded and impacted teeth (KØ1)	KØ1.Ø Embedded teeth
KØØ.Ø Anodontia Hypodontia	KØ1.1 Impacted teeth
Oligodontia	KØ2 Dental caries
<b>EXCLUDES 1</b> acquired absence of teeth (KØ8.1-)	[INCLUDES] caries of dentine dental cavities
<b>DEF:</b> Partial or complete absence of teeth due to a congenital defect involving the tooth bud.	early childhood caries
KØØ.1 Supernumerary teeth	pre-eruptive caries recurrent caries (dentino enamel junction) (enamel) (to
Distomolar Fourth molar	the pulp)
Mesiodens	tooth decay DEF: Localized section of tooth decay that begins on the tooth surface
Paramolar Supplementary teeth	with destruction of the calcified enamel, allowing bacterial destruction
EXCLUDES 2 supernumerary roots (KØØ.2)	to continue and form cavities and may extend to the dentin and pulp.
KØØ.2 Abnormalities of size and form of teeth Concrescence of teeth	Tooth Anatomy
Fusion of teeth	
Gemination of teeth	Enamel
Dens evaginatus Dens in dente	Dentin
Dens invaginatus	Pulp chamber
Enamel pearls Macrodontia	rup chamber
Microdontia	
Peg-shaped [conical] teeth Supernumerary roots	Root canal
Taurodontism	
Tuberculum paramolare	Cementum
(A5Ø.5)	
tuberculum Carabelli, which is regarded as a normal variation and should not be coded	
KØØ.3 Mottled teeth	
Dental fluorosis	870.53 112 2020 a.R.
Mottling of enamel Nonfluoride enamel opacities	KØ2.3 Arrested dental caries
EXCLUDES 2 deposits [accretions] on teeth (KØ3.6)	Arrested coronal and root caries
KØØ.4 Disturbances in tooth formation Aplasia and hypoplasia of cementum	KØ2.5 Dental caries on pit and fissure surface
Dilaceration of tooth	Dental caries on chewing surface of tooth KØ2.51 Dental caries on pit and fissure surface limited to
Enamel hypoplasia (neonatal) (postnatal) (prenatal)	enamel
Regional odontodysplasia Turner's tooth	White spot lesions [initial caries] on pit and fissure surface of tooth
<b>EXCLUDES 1</b> Hutchinson's teeth and mulberry molars in congenital	KØ2.52 Dental caries on pit and fissure surface penetrating
syphilis (A5Ø.5) EXCLUDES 2 mottled teeth (KØØ.3)	into dentin Primary dental caries, cervical origin
	KØ2.53 Dental caries on pit and fissure surface penetrating
	into pulp

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Additional Character Required ICD-10-CM 2022

✓x7<sup>th</sup> Placeholder

Unacceptable PDx

Questionable PDx

Manifestation

Unspecified Dx

H1 - H14 HAC

HCC CMS-HCC Dx

J-10-CIV	2022	Chapter 18. Symptoms, Signs and Abn	Roz.z-Ro4.4
	D02.2	Diliunia	
		Biliuria	
	R82.3	Hemoglobinuria	Abnormal findings on examination of other body fluids,
		<b>EXCLUDES 1</b> hemoglobinuria due to hemolysis from external	substances and tissues, without diagnosis (R83-R89)
		<b>3</b> ,	
		causes NEC (D59.6)	<b>EXCLUDES 1</b> abnormal findings on antenatal screening of mother (O28)
		hemoglobinuria due to paroxysmal nocturnal	diagnostic abnormal findings classified elsewhere — see Alphabetical
		[Marchiafava-Micheli] (D59.5)	
			Index
		DEF: Free hemoglobin in blood due to rapid hemolysis of red	<b>EXCLUDES 2</b> abnormal findings on examination of blood, without diagnosis
		blood cells. Causes include burns, crushed injury, sickle cell	(R7Ø-R79)
		anemia, thalassemia, parasitic infections, or kidney infections.	
			abnormal findings on examination of urine, without diagnosis
	R82.4	Acetonuria	(R8Ø-R82)
		Ketonuria	abnormal tumor markers (R97)
		<b>DEF:</b> Excessive excretion of acetone in urine that commonly occurs	
			R83 Abnormal findings in cerebrospinal fluid
		in diabetic acidosis.	
	R82.5	Elevated urine levels of drugs, medicaments and biological	
		substances	Cerebrospinal Fluid
		Elevated urine levels of catecholamines	
		Elevated urine levels of indoleacetic acid	AT THE PARTY
		Elevated urine levels of 17-ketosteroids	
		Elevated urine levels of steroids	
	D926	Abnormal urine levels of substances chiefly nonmedicinal	
	102.0		
		as to source	Ventricles
		Abnormal urine level of heavy metals	Ventricles
/5th	D92 7	Abnormal findings on microbiological examination of urine	
V J	N02./	Abilotinal multips on microbiological examination of unite	alw -
		EXCLUDES 1 colonization status (Z22)	Cerebrospinal
			fluid
		AHA: 2016,4Q,65	
		R82.71 Bacteriuria	
		R82.79 Other abnormal findings on microbiological	447
		examination of urine	
		Positive culture findings of urine	
	<b>Baa a</b>	5	Spinal
√ 5 <sup>th</sup>	K82.8	Abnormal findings on cytological and histological	cord
		examination of urine	4
		AHA: 2019,4Q,16	G
		R82.81 Pyuria	
		Sterile pyuria	YAF.
		R82.89 Other abnormal findings on cytological and	ar Fi
		histological examination of urine	
√5 <sup>th</sup>	R82.9	Other and unspecified abnormal findings in urine	C C C
		R82.90 Unspecified abnormal findings in urine	
		R82.91 Other chromoabnormalities of urine	
		Chromoconversion (dipstick)	
		Idiopathic dipstick converts positive for blood with	
			۳ × ۵ کر
		no cellular forms in sediment	(° ~~ ~
		EXCLUDES 1 hemoglobinuria (R82.3)	(°)
		myoglobinuria (R82.1)	
			A A A A A A A A A A A A A A A A A A A
	√6 <sup>th</sup>	R82.99 Other abnormal findings in urine	5
		AHA: 2018,4Q,29-30	
			R83.Ø Abnormal level of enzymes in cerebrospinal fluid
		R82.991 Hypocitraturia	
		R82.992 Hyperoxaluria	R83.1 Abnormal level of hormones in cerebrospinal fluid
			R83.2 Abnormal level of other drugs, medicaments and biological
		<b>EXCLUDES 1</b> primary hyperoxaluria (E72.53)	substances in cerebrospinal fluid
		R82.993 Hyperuricosuria	•
		R82.994 Hypercalciuria	R83.3 Abnormal level of substances chiefly nonmedicinal as to
			source in cerebrospinal fluid
		Idiopathic hypercalciuria	R83.4 Abnormal immunological findings in cerebrospinal fluid
		R82.998 Other abnormal findings in urine	• • •
		Cells and casts in urine	R83.5 Abnormal microbiological findings in cerebrospinal fluid
			Positive culture findings in cerebrospinal fluid
		Crystalluria	EXCLUDES 1 colonization status (Z22)
		Melanuria	
			R83.6 Abnormal cytological findings in cerebrospinal fluid
			R83.8 Other abnormal findings in cerebrospinal fluid
			Abnormal chromosomal findings in cerebrospinal fluid
			D22.0 Unenceified abnormal finding in corebroeninal fluid
			R83.9 Unspecified abnormal finding in cerebrospinal fluid
			R84 Abnormal findings in specimens from respiratory organs and
			thorax
			<b>INCLUDES</b> abnormal findings in bronchial washings
			abnormal findings in nasal secretions
			5
			abnormal findings in pleural fluid
			abnormal findings in sputum
			5 1
			abnormal findings in throat scrapings
			<b>EXCLUDES 1</b> blood-stained sputum (RØ4.2)
			R84.Ø Abnormal level of enzymes in specimens from respiratory
			organs and thorax
			R84.1 Abnormal level of hormones in specimens from respiratory
			organs and thorax
			R84.2 Abnormal level of other drugs, medicaments and biological
			substances in specimens from respiratory organs and thorax
			R84.3 Abnormal level of substances chiefly nonmedicinal as to
			source in specimens from respiratory organs and thorax
			R84.4 Abnormal immunological findings in specimens from
			respiratory organs and thorax

HIV HIV Dx 951

## Appendix A: Valid 3-Character ICD-10-CM Codes

AØ9	Infectious gastroenteritis and colitis, unspecified
A33	Tetanus neonatorum
A34	Obstetrical tetanus
A35	Other tetanus
A46	Erysipelas
A55	Chlamydial lymphogranuloma (venereum)
A57	Chancroid
A58	Granuloma inguinale
A64	Unspecified sexually transmitted disease
A65	Nonvenereal syphilis
A7Ø	Chlamydia psittaci infections
A78	Q fever
A86	Unspecified viral encephalitis
A89	Unspecified viral infection of central nervous system
A9Ø	Dengue fever [classical dengue]
A91	Dengue hemorrhagic fever
A94	Unspecified arthropod-borne viral fever
A99	Unspecified viral hemorrhagic fever
BØ3	Smallpox
BØ4	Monkeypox
BØ9	Unspecified viral infection characterized by skin and mucous membrane lesions
B2Ø	Human immunodeficiency virus [HIV] disease
B49	Unspecified mycosis
B54	Unspecified malaria
B59	Pneumocystosis
B64	Unspecified protozoal disease
B72	Dracunculiasis
B75	Trichinellosis
B79	Trichuriasis
B8Ø	Enterobiasis
B86	Scabies
B89	Unspecified parasitic disease
B91	Sequelae of poliomyelitis
B92	Sequelae of leprosy
CØ1	Malignant neoplasm of base of tongue
CØ7	Malignant neoplasm of parotid gland
C12	Malignant neoplasm of pyriform sinus
C19	Malignant neoplasm of rectosigmoid junction
C2Ø	Malignant neoplasm of rectum
C23	Malignant neoplasm of gallbladder
C33	Malignant neoplasm of trachea
C37	Malignant neoplasm of thymus
C52	Malignant neoplasm of vagina
C55	Malignant neoplasm of uterus, part unspecified
C58	Malignant neoplasm of placenta
C61	Malignant neoplasm of prostate
C73	Malignant neoplasm of thyroid gland
D34	Benign neoplasm of thyroid gland
D45	Polycythemia vera
D62	Acute posthemorrhagic anemia
D65	Disseminated intravascular coagulation [defibrination A syndrome]
D66	Hereditary factor VIII deficiency
D67	Hereditary factor IX deficiency
D71	Functional disorders of polymorphonuclear neutrophils
D77	Other disorders of blood and blood-forming organs in diseases classified elsewhere
EØ2	Subclinical iodine-deficiency hypothyroidism
E15	Nondiabetic hypoglycemic coma
E35	Disorders of endocrine glands in diseases classified elsewhere
E4Ø	Kwashiorkor
E41	Nutritional marasmus
E42	Marasmic kwashiorkor
E43	Unspecified severe protein-calorie malnutrition
E45	Retarded development following protein-calorie malnutrition

E46	Unspecified protein-calorie malnutrition
E52	Niacin deficiency [pellagra]
E54	Ascorbic acid deficiency
E58	Dietary calcium deficiency
E59	Dietary selenium deficiency
E6Ø	Dietary zinc deficiency
E65	Localized adiposity
E68	Sequelae of hyperalimentation
FØ4	Amnestic disorder due to known physiological condition
FØ5	Delirium due to known physiological condition
FØ9	Unspecified mental disorder due to known physiological condition
F21	Schizotypal disorder
F22	Delusional disorders
F23	Brief psychotic disorder
F24	Shared psychotic disorder
F28	Other psychotic disorder not due to a substance or known physiological condition
F29	Unspecified psychosis not due to a substance or known physiological condition
F39	Unspecified mood [affective] disorder
F54	Psychological and behavioral factors associated with disorders or diseases classified elsewhere
F59	Unspecified behavioral syndromes associated with physiological disturbances and physical factors
F66	Other sexual disorders
F69	Unspecified disorder of adult personality and behavior
F7Ø	Mild intellectual disabilities
F71	Moderate intellectual disabilities
F72	Severe intellectual disabilities
F73	Profound intellectual disabilities
F78	Other intellectual disabilities
F79	Unspecified intellectual disabilities
F82	Specific developmental disorder of motor function
F88	Other disorders of psychological development
F89	Unspecified disorder of psychological development
F99	Mental disorder, not otherwise specified
GØ1	Meningitis in bacterial diseases classified elsewhere
GØ2	Meningitis in other infectious and parasitic diseases classified elsewhere
GØ7	Intracranial and intraspinal abscess and granuloma in diseases classified elsewhere
GØ8	Intracranial and intraspinal phlebitis and thrombophlebitis
GØ9	Sequelae of inflammatory diseases of central nervous system
G1Ø	Huntington's disease
G14	Postpolio syndrome
G2Ø	Parkinson's disease
G26	Extrapyramidal and movement disorders in diseases classified elsewhere
G35	Multiple sclerosis
G53	Cranial nerve disorders in diseases classified elsewhere
G55	Nerve root and plexus compressions in diseases classified elsewhere
G59	Mononeuropathy in diseases classified elsewhere
G63	Polyneuropathy in diseases classified elsewhere
G64	Other disorders of peripheral nervous system
G92	Toxic encephalopathy
G94	Other disorders of brain in diseases classified elsewhere
H22	Disorders of iris and ciliary body in diseases classified elsewhere
H28	Cataract in diseases classified elsewhere
H32	Chorioretinal disorders in diseases classified elsewhere
H36	Retinal disorders in diseases classified elsewhere
H42	Glaucoma in diseases classified elsewhere
IØØ	Rheumatic fever without heart involvement

- I1Ø Essential (primary) hypertension
- 132 Pericarditis in diseases classified elsewhere
- I38 Endocarditis, valve unspecified

## **Appendix F: Present on Admission (POA) Tutorial**

Present-on-admission (POA) indicators are required on all diagnoses and external cause-of-injury codes for all inpatient acute care hospital discharges, according to the requirement outlined in the Deficit Reduction Act of 2005, section 5001(c). The only exception is for a select group of codes that have been designated as exempt from the POA regulations. Certain facilities are also exempt from this reporting requirement and include the following:

- Critical access hospitals
- Long-term care hospitals
- Cancer hospitals
- Inpatient psychiatric hospitals
- Inpatient rehabilitation facilities
- Veterans Administration/Department of Defense Hospitals
- Children's inpatient facilities

The POA indicator guidelines are not intended to replace the official *ICD-10-CM Official Guidelines for Coding and Reporting*. Nor will the POA guidelines supersede the Uniform Hospital Discharge Data Set (UHDDS) definition of principal diagnosis, defined as the condition established after study to be chiefly responsible for occasioning the admission of the patient to the hospital. The definition of "additional diagnoses" also remains the same. Coders must report all conditions that coexist at the time of admission, develop subsequently, or that affect the treatment received and/or the length of stay. This significantly affects assignment of the POA indicators. Because the POA indicator should accurately reflect the patient's conditions upon admission, review of POA guidelines as well as the *ICD-10-CM Official Guidelines for Coding and Reporting* is required.

The following is a summary of the POA guidelines:

- A POA condition is defined not only as one that is clearly present at the time of admission, but also one that was clearly present but not diagnosed until after the time of admission.
- Present on admission is defined as present at the time the order for an inpatient admission is made, whether or not the patient's episode of care originated in the emergency department, ambulatory surgery area, or other outpatient area. Conditions that develop during an outpatient encounter are considered present on admission.
- A POA indicator is assigned to the principal diagnosis, secondary diagnoses, and the external cause-of-injury codes.
- The POA guidelines are not intended to replace any of the other ICD-10-CM Official Guidelines for Coding and Reporting.
- The intent of the POA guidelines is not to determine whether or not a condition should be coded, but rather how to apply the POA indicator.

The following reporting options, with definitions, should be used to indicate whether a condition was POA:

- Y Yes: Present at the time of inpatient admission
- N No: Not present at the time of inpatient admission
- U Unknown: Documentation is insufficient to determine if condition is present at the time of in patient admission
- W Clinically undetermined: Provider is unable to clinically determine whether condition was or was not present at the time of inpatient admission

Unreported/not used—(exempt from POA reporting)—Most ICD-10-CM codes require the assignment of a POA indicator. However, there are some diagnoses that are considered exempt from the POA reporting. For these exempt conditions the POA field should be left blank.

**Note:** A data file of all POA exempt codes can be accessed at the following: http://www.optum360coding.com/ProductUpdates/ Title: "2021 ICD-10-CM for Hospitals POA Exemption Data File" Password: Hospital21

## Timeframe for POA identification and documentation

There is no timeframe during the encounter in which a condition must be diagnosed to be considered present on admission. In many cases, it is not clinically possible for the provider to determine a definitive diagnosis until well into the admission, after labs, imaging, and other tests have been

performed and results examined. In these cases the condition is still considered to be present on admission, indicated with Y.

#### Examples

 A patient was admitted for diagnostic work-up for cachexia. The final diagnosis was malignant neoplasm of lung with metastasis.

Assign Y in the POA field for the malignant neoplasm, which was clearly present on admission, although it was not diagnosed until after the admission occurred.

A patient with severe cough and difficulty breathing was diagnosed during his hospitalization to have lung cancer.

Assign Y in the POA field for the lung cancer. Even though the cancer was not diagnosed until after admission, it is a chronic condition that was clearly present before the patient's admission.

• A urine culture was obtained on admission. The provider documented urinary tract infection when the culture results became available a few days later.

Assign Y to the urinary tract infection since the diagnosis is based on test results from a specimen obtained on admission. It may not be possible for a provider to make a definitive diagnosis for a period of time after admission. There is no required timeframe as to when a provider must identify or document a condition for it to be considered present on admission.

## Condition is on the "Exempt from Reporting" list

The POA field is left blank for codes excluded from POA reporting.

#### Examples

A patient was admitted for inpatient surgery on a burn contracture from scar tissue from a burn five years ago.

Assign Y for the principal condition of the scar; however, the secondary code for the specifics of the burn injury with a seventh character for sequela is left blank since it is excluded from POA reporting.

A patient was admitted to inpatient for a PEG tube insertion because of residual dysphagia due to a recent CVA.

No POA indicator is assigned for the dysphagia following a CVA. Codes under 169.- are defined as sequela of cerebrovascular disease and are exempt from POA reporting.

A patient was admitted to inpatient for corrective surgery for a malunion of a closed femur fracture.

No POA indicator is assigned for the malunion of a closed fracture because the seventh character for malunion of fractures is considered a subsequent encounter and is exempt from POA reporting.

A patient was admitted for breast reconstruction following mastectomy.

The POA indicator field is left blank as this Z code is exempt from POA reporting.

A patient was admitted for changing the battery in his AICD as it is at the end of its useful life.

The POA indicator field is left blank as this Z code is exempt from POA reporting.

## POA explicitly documented

If the provider explicitly documents a condition as being present on admission or not present on admission, assign the appropriate Y or N POA indicator based on that documentation. If there is conflicting documentation by different providers, query the attending physician.

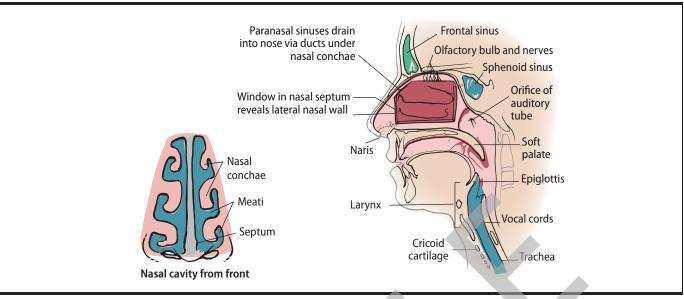
#### Examples

Even though sepsis is not diagnosed in the ED, the discharge summary states that the patient presented with sepsis at the time of admission.

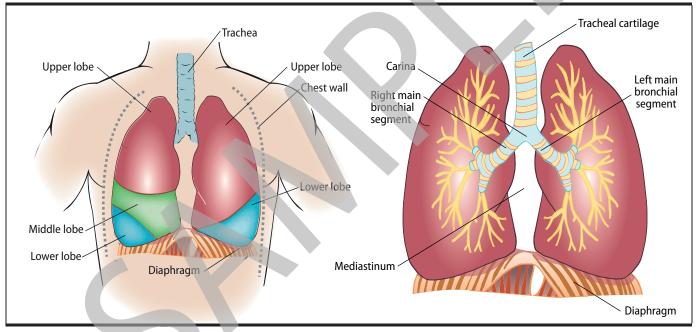
The provider explicitly stated that the sepsis was present on admission and therefore the Y POA indicator should be used.

The patient was admitted with chills and cough and two days later diagnosed with pneumonia. The discharge summary documents that the pneumonia was not present on admission.

Although the patient showed some symptoms of pneumonia on admission, the provider may have clinical evidence (e.g., chest x-rays, cultures) that indicates that pneumonia did not develop until after admission. Since this



Lower Respiratory System



**Paranasal Sinuses** 

