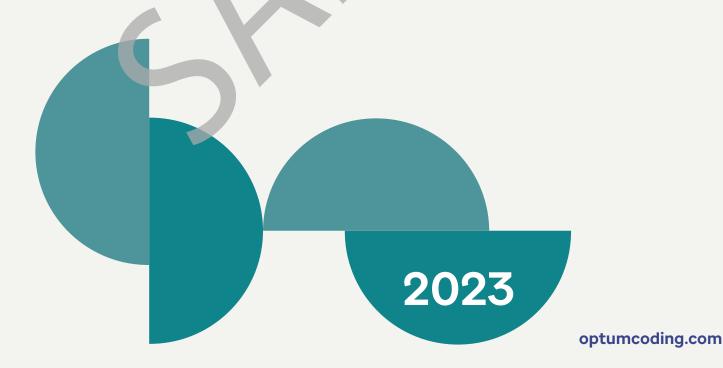


# ICD-10-CM Expert for Skilled Nursing Facilities and Inpatient Rehabilitation Facilities

The complete official code set

Codes valid from October 1, 2022 through September 30, 2023



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New for 2023

New for 2023

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### 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.

# **Code Also**

A "code also" note alerts the coder that more than one code may be required to fully describe the condition. The sequencing depends on the circumstances of the encounter. Factors that may determine sequencing include severity and reason for the encounter.

### **Revised Text**

The revised text ▶ ◀ "bow ties" alert the user to changes in official notations for the current year. Revised text may include the following:

- A change in a current parenthetical description
- A change in the code(s) associated with a current parenthetical note
- A change in how a current parenthetical note is classified (e.g., an Excludes 1 note that changed to an Excludes 2 note)
- · Addition of a new parenthetical note(s) to a code

### Deleted Text

Strikethrough on official notations indicate a deletion from the classification for the current year.

# **Optum360 Notations**

# **AHA Coding Clinic Citations**

Coding Clinics are official American Hospital Association (AHA) publications that provide coding advice specific to ICD-10-CM and ICD-10-PCS.

Coding Clinic citations included in this manual are current up to the second quarter of 2021.

These citations identify the year, quarter, and page number of one or more *Coding Clinic* publications that may have coding a dvice relevant to a particular code or group of codes. With the most current citation listed first, these notations are preceded by the symbol **AHA**: and appear in purple type.

I15.1 Hypertension secondary to other renal disorders AHA: 2016, 3Q, 22

### **Definitions**

Definitions explain a specific term, condition, or disease process in layman's terms. These notations are preceded by the symbol **DEF**: and appear in purple type.

M51.4 Schmorl's nodes

**DEF:** Irregular bone defect in the margin of the vertebral body that causes herniation into the end plate of the vertebral body.

# **Coding Tips**

The tips in the tabular list offer coding advice that is not readily available within the ICD-10-CM classification. They may relate to official coding guidelines, indexing nuances, or advice from AHA's Coding Clinic for ICD-10-CM/PCS. These notations are preceded by the symbol **TIP:** and appear in brown type.

**B97.2** Coronavirus as the cause of diseases classified elsewhere TIP: Do not report a code from this subcategory for COVID-19; refer to UØ7.1.

### **RIC Excl**

Rehabilitation impairment categories (RIC) may qualify for a payment adjustment when certain conditions, considered comorbid conditions (CC), are assigned as secondary diagnoses. However, there are certain CCs that are excluded from specific RIC categories. This reference identifies those RIC categories to which the C is excluded. These notations are preceded by the symbol RIC Excl. and appear in blue type.

140.8 Other acute myocarditis
RIC Excl: 14 Cardiac

If no RIC exclusion reference is listed, the CC may qualify a case for payment adjustment regardless of the RIC category assigned. For a comprehensive list of all RIC CCs and the RIC categories from which they may be excluded, please refer to appendix K, "RIC Comorbid Conditions," at the back of this book.

### Icons

Note: The following icons are placed to the left of the code.

New Code

Codes that have been added to the classification effective October 1, 2022.

New Code – Mid-year

Codes that have been added to the classification effective April 1, 2022.

# Revised Code

Codes that have had a change to their description or validity effective October 1, 2022. For additional information on codes with validity changes, see the "What's New" section.

Revised Code – Mid-year

Codes that have had a change to their description or validity effective April 1, 2022.

Additional Characters Required

This symbol indicates that the code requires a 4th character.

This symbol indicates that the code requires a 5th character.

This symbol indicates that the code requires a 6th character.

This symbol indicates that the code requires a 7th character.

H6Ø.3 Other infective otitis externa

H6Ø.31 Diffuse otitis externa

H6Ø.311 Diffuse otitis externa, right ear

H6Ø.312 Diffuse otitis externa, left ear

H6Ø.313 Diffuse otitis externa, bilateral

H6Ø.319 Diffuse otitis externa, unspecified ear

# 71 Placeholder Alert

This symbol indicates that the code requires a 7th character following the placeholder "X". Codes with fewer than six characters that require a 7th character must contain placeholder "X" to fill in the empty character(s).

T16.1 Foreign body in right ear

**Note:** The following icons are placed at the end of the code description.

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# **10 Steps to Correct Coding**

Follow the 10 steps below to correctly code encounters for health care services.

# Step 1: Identify the reason for the visit or encounter (i.e., a sign, symptom, diagnosis and/or condition).

The medical record documentation should accurately reflect the patient's condition, using terminology that includes specific diagnoses and symptoms or clearly states the reasons for the encounter.

Choosing the main term that best describes the reason chiefly responsible for the service provided is the most important step in coding. If symptoms are present and documented but a definitive diagnosis has not yet been determined, code the symptoms. For outpatient cases, do not code conditions that are referred to as "rule out," "suspected," "probable," or "questionable." Diagnoses often are not established at the time of the initial encounter/visit and may require two or more visits to be established. Code only what is documented in the available outpatient records and only to the highest degree of certainty known at the time of the patient's visit. For inpatient medical records, uncertain diagnoses may be reported if documented at the time of discharge.

# Step 2: After selecting the reason for the encounter, consult the alphabetic index.

The most critical rule is to begin code selection in the alphabetic index. Never turn first to the tabular list. The index provides cross-references, essential and nonessential modifiers, and other instructional notations that may not be found in the tabular list.

# Step 3: Locate the main term entry.

The alphabetic index lists conditions, which may be expressed as nouns or eponyms, with critical use of adjectives. Some conditions known by several names have multiple main entries. Reasons for encounters may be located under general terms such as admission, encounter, and examination. Other general terms such as history, status (post), or presence (of) can be used to locate other factors influencing health.

# Step 4: Scan subterm entries.

Scan the subterm entries, as appropriate, being sure to review continued lines and additional subterms that may appear in the next column or on the next page. Shaded vertical guidelines in the index indicate the indentation level for each subterm in relation to the main terms.

# Step 5: Pay close attention to index instructions.

- Parentheses () enclose nonessential modifiers, terms that are supplementary words or explanatory information that may or may not appear in the diagnostic statement and do not affect code selection.
- Brackets [] enclose manifestation codes that can be used only as secondary codes to the underlying condition code immediately preceding it. If used, manifestation codes must be reported with the appropriate etiology codes.
- Default codes are listed next to the main term and represent the condition most commonly associated with the main term or the unspecified code for the main term.
- "See" cross-references, identified by italicized type and "code by" cross-references indicate that another term must be referenced to locate the correct code.
- "See also" cross-references, identified by italicized type, provide alternative terms that may be useful to look up but are not mandatory.
- "Omit code" cross-references identify instances when a code is not applicable depending on the condition being coded.
- "With" subterms are listed out of alphabetic order and identify a presumed causal relationship between the two conditions they link.
- "Due to" subterms identify a relationship between the two conditions they link.

- "NEC," abbreviation for "not elsewhere classified," follows some main terms or subterms and indicates that there is no specific code for the condition even though the medical documentation may be very specific.
- "NOS," abbreviation for "not otherwise specified," follows some main terms or subterms and is the equivalent of unspecified; NOS signifies that the information in the medical record is insufficient for assigning a more specific code.
- Following references help coders locate alphanumeric codes that are out of sequence in the tabular section.
- Check-additional-character symbols flag codes that require additional characters to make the code valid; the characters available to complete the code should be verified in the tabular section.

# Step 6: Choose a potential code and locate it in the tabular list.

To prevent coding errors, always use both the alphabetic index (to identify a code) and the tabular list (to verify a code), as the index does not include the important instructional notes found in the tabular list. An added benefit of using the tabular list, which groups like things together, is that while looking at one code in the list, a coder might see a more specific one that would have been missed had the coder relied solely on the alphabetic index. Additionally, many of the codes require a fourth, fifth, sixth, or seventh character to be valid, and many of these characters can be found only in the tabular list.

# Step 7: Read all instructional material in the tabular section.

The coder must follow any Includes, Excludes 1 and Excludes 2 notes, and other instructional notes, such as "Code first" and "Use additional code," listed in the tabular list for the chapter, category, subcategory, and subclassification levels of code selection that direct the coder to use a different or additional code. Any codes in the tabular range AØØ.Ø-through T88.9- may be used to identify the diagnostic reason for the encounter. The tabular list encompasses many codes describing disease and injury classifications (e.g., infectious and parasitic diseases, neoplasms, symptoms, nervous and circulatory system etc.).

Codes that describe symptoms and signs, as opposed to definitive diagnoses, should be reported when an established diagnosis has not been made (confirmed) by the physician. Chapter 18 of the ICD-10-CM code book, "Symptoms, Signs, and Abnormal Clinical and Laboratory Findings, Not Elsewhere Classified" (codes RØØ.—R99), contains many, but not all, codes for symptoms.

ICD-10-CM classifies encounters with health care providers for circumstances other than a disease or injury in chapter 21, "Factors Influencing Health Status and Contact with Health Services" (codes ZØØ–Z99). Circumstances other than a disease or injury often are recorded as chiefly responsible for the encounter.

A code is invalid if it does not include the full number of characters (greatest level of specificity) required. Codes in ICD-10-CM can contain from three to seven alphanumeric characters. A three-character code is to be used only if the category is not further subdivided into four-, five-, six-, or seven-character codes. Placeholder character X is used as part of an alphanumeric code to allow for future expansion and as a placeholder for empty characters in a code that requires a seventh character but has no fourth, fifth, or sixth character. Note that certain categories require seventh characters that apply to all codes in that category. Always check the category level for applicable seventh characters for that category.

# Step 8: Consult the official ICD-10-CM conventions and quidelines.

The ICD-10-CM Official Guidelines for Coding and Reporting govern the use of certain codes. These guidelines provide both general and chapter-specific coding guidance.

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Adhesions, adhesive — continued	Adiposis — see also Obesity	Admission — continued
mesenteric — see Adhesions, peritoneum	cerebralis E23.6	adjustment — continued
nasal (septum) (to turbinates) J34.89 ocular muscle — <i>see</i> Strabismus, mechanical	dolorosa E88.2 <b>Adiposity</b> — see also Obesity	neuropacemaker (brain) (peripheral nerve) (spinal cord) Z46.2
omentum — see Adhesions, peritoneum	heart — see Degeneration, myocardial	implanted Z45.42
ovary N73.6	localized E65	orthodontic device Z46.4
congenital (to cecum, kidney or omentum) Q50.39 paraovarian N73.6	Adiposogenital dystrophy E23.6 Adjustment	orthopedic (brace) (cast) (device) (shoes) Z46.89 pacemaker (cardiac resynchronization therapy (CRT-
pelvic (peritoneal)	disorder — see Disorder, adjustment	P))
female N73.6	implanted device — see Encounter (for), adjustment	cardiac Z45.018
postprocedural N99.4 male — <i>see</i> Adhesions, peritoneum	(of)	pulse generator Z45.010 nervous system Z46.2
postpartal (old) N73.6	prosthesis, external — see Fitting reaction — see Disorder, adjustment	implanted Z45.42
tuberculous A18.17	Administration of tPA (rtPA) in a different facility within	portacath (port-a-cath) Z45.2
penis to scrotum (congenital) Q55.8 periappendiceal — <i>see also</i> Adhesions, peritoneum	the last 24 hours prior to admission to current facil-	prosthesis Z44.9 arm — see Admission, adjustment, artificial, arm
pericardium (nonrheumatic) 131.0	ity Z92.82 <b>Admission</b> (for) — see also Encounter (for)	breast Z44.3 <b>☑</b>
focal I31.8	adjustment (of)	dental Z46.3
rheumatic 109.2 tuberculous A18.84	artificial	eye Z44.2 🗹 leg — see Admission, adjustment, artificial, leg
pericholecystic K82.8	arm Z44.00- ☑ complete Z44.01- ☑	specified NEC Z44.8
perigastric — <i>see</i> Adhesions, peritoneum	partial Z44.02- ✓	spectacles Z46.0
periovarian N73.6 periprostatic N42.89	eye Z44.2 🔽	aftercare — see also Aftercare Z51.89 postpartum
periprostatic 1442.09 perirectal — see Adhesions, peritoneum	leg Z44.10- ✓	immediately after delivery Z39.0
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peritoneum, peritoneal (postinfective) K66.0	brain neuropacemaker Z46.2	radiation therapy (antineoplastic) Z51.Ø
with obstruction (intestinal) K56.5Ø complete K56.52	implanted Z45.42	attention to artificial opening (of) Z43.9 artificial vagina Z43.7
incomplete K56.51	breast implant Z45.81 ☑	colostomy Z43.3
partial K56.51	prosthesis (external) Z44.3	cystostomy Z43.5
congenital Q43.3 pelvic, female N73.6	colostomy belt Z46.89	enterostomy Z43.4 gastrostomy Z43.1
postprocedural N99.4	contact lenses Z46.0 cystostomy device Z46.6	ileostomy Z43.2
postpartal, pelvic N73.6	dental prosthesis Z46.3	jejunostomy Z43.4
postprocedural K66.0 to uterus N73.6	device NEC	nephrostomy Z43.6 specified site NEC Z43.8
peritubal N73.6	abdominal Z46.89	intestinal tract Z43.4
periureteral N28.89	implanted Z45.89 cardiac Z45.09	urinary tract Z43.6
periuterine N73.6 perivesical N32.89	defibrillator (with synchronous cardiac	tracheostomy Z43.0 ureterostomy Z43.6
perivesical N32.89 perivesicular (seminal vesicle) N5Ø.89	pacemaker) Z45.02	urethrostomy Z43.6
pleura, pleuritic J94.8	pacemaker (cardiac resynchronization therapy (CRT-P)) Z45.018	breast augmentation or reduction Z41.1
tuberculous NEC A15.6 pleuropericardial J94.8	pulse generator Z45.Ø1Ø	breast reconstruction following mastectomy Z42.1 change of
postoperative (gastrointestinal tract) K66.Ø	resynchronization therapy defibrillator	dressing (nonsurgical) Z48.00
with obstruction — see also Obstruction, intestine,	(CRT-D) Z45.02 hearing device Z45.328	neuropacemaker device (brain) (peripheral nerve)
postoperative K91.30 due to foreign body accidentally left in wound —	bone conduction Z45.320	(spinal cord) Z46.2 implanted Z45.42
see Foreign body, accidentally left during a	cochlear Z45.321	surgical dressing Z48.01
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postpartal, old (vulva or perineum) N9Ø.89	device implanted, hearing	contraceptive management Z3Ø.9
preputial, prepuce N47.5 pulmonary J98.4	neuropacemaker Z45.42	cosmetic surgery NEC Z41.1 counseling — see also Counseling
pylorus — see Adhesions, peritoneum	visual substitution Z45.31	dietary Z71.3
sciatic nerve — see Lesion, nerve, sciatic	specified NEC Z45.89 vascular access Z45.2	gestational carrier Z31.7
seminal vesicle N50.89 shoulder (joint) — see Capsulitis, adhesive	visual substitution Z45.31	HIV Z71.7 human immunodeficiency virus Z71.7
sigmoid flexure — see Adhesions, peritoneum	nervous system Z46.2 implanted — see Admission, adjustment,	nonattending third party Z71.0
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tunica vaginalis N44.8	implanted — see Admission, adjustment,	donor (potential) ZØØ.5 ear ZØ1.1Ø
uterus N73.6 internal N85.6	device, implanted, hearing device	with abnormal findings NEC ZØ1.118
to abdominal wall N73.6	nervous system Z46.2 implanted — see Admission, adjustment,	eye Z01.00
vagina (chronic) N89.5	device, implanted, nervous system	with abnormal findings ZØ1.Ø1 following failed vision screening ZØ1.Ø2Ø
postoperative N99.2 vitreomacular H43.82- <b>▼</b>	visual Z46.2	with abnormal findings ZØ1.Ø21
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Adiaspiromycosis B48.8 Adie (-Holmes) pupil or syndrome — see Anomaly,	implanted — see Admission, adjustment, device,	infant or child (over 28 days old) ZØØ.129
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Adiponecrosis neonatorum P83.88	intestinal appliance or device NEC Z46.89	postpartum checkup Z39.2

# Chapter 4. Endocrine, Nutritional and Metabolic Diseases (EØØ–E89)

# **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. Diabetes mellitus

The diabetes mellitus codes are combination codes that include the type of diabetes mellitus, the body system affected, and the complications affecting that body system. As many codes within a particular category as are necessary to describe all of the complications of the disease may be used. They should be sequenced based on the reason for a particular encounter. Assign as many codes from categories EØ8–E13 as needed to identify all of the associated conditions that the patient has.

Patient is admitted with reported diagnoses of uncontrolled diabetes, type 2, with diabetic polyneuropathy and diabetic retinopathy with macular edema. Endocrinology clinical notes indicate a current HgA1c of 12.4 and "persistent blood glucose elevations over 300."

E11.65 Type 2 diabetes mellitus with hyperglycemia

E11.311 Type 2 diabetes mellitus with unspecified diabetic retinopathy with macular edema

E11.42 Type 2 diabetes mellitus with diabetic polyneuropathy

Explanation: Use as many codes to describe the diabetic complications as needed. Many are combination codes that describe more than one condition. "Uncontrolled" may reference hypo- or hyperglycemia, based upon available provider documentation.

### 1) Type of diabetes

The age of a patient is not the sole determining factor, though most type 1 diabetics develop the condition before reaching puberty. For this reason, type 1 diabetes mellitus is also referred to as juvenile diabetes.

A 45-year-old patient is diagnosed with type 1 diabetes.

## E10.9 Type 1 diabetes mellitus without complications

Explanation: Although most type 1 diabetics are diagnosed in childhood or adolescence, it can also begin in adults.

# 2) Type of diabetes mellitus not documented

If the type of diabetes mellitus is not documented in the medical record the default is E11.-, Type 2 diabetes mellitus.

Patient is referred with new diagnoses reported in provider clinical notes as diabetes and hypertension.

E11.9 Type 2 diabetes mellitus without complications

I10 Essential (primary) hypertension

*Explanation*: Since the type of diabetes was not documented and no complications were noted, the default code is E11.9.

# 3) Diabetes mellitus and the use of insulin, oral hypoglycemics, and injectable non-insulin drugs

If the documentation in a medical record does not indicate the type of diabetes but does indicate that the patient uses insulin, code E11-, Type 2 diabetes mellitus, should be assigned. **Additional** code(s) should be assigned from category Z79 to identify the long-term (current) use of insulin, oral hypoglycemic drugs, or injectable non-insulin antidiabetic, as follows:

If the patient is treated with both oral medications and insulin, **both** code **Z79.4**, **Long term (current) use of insulin, and code Z79.84**, **Long term (current) use of oral hypoglycemic drugs**, should be assigned

If the patient is treated with both insulin and an injectable non-insulin antidiabetic drug, assign codes Z79.4, Long term (current) use of insulin, and Z79.899, Other long term (current) drug therapy.

If the patient is treated with both oral hypoglycemic drugs and an injectable non-insulin antidiabetic drug, assign codes Z79.84, Long term (current) use of oral hypoglycemic drugs, and Z79.899, Other long term (current) drug therapy.

Code Z79.4 should not be assigned if insulin is given temporarily to bring a type 2 patient's blood sugar under control during an encounter.

Patient is referred with documented provider notes reporting 10-year history of diabetes requiring daily insulin use.

# E11.9 Type 2 diabetes mellitus without complications

# Z79.4 Long term (current) use of insulin

*Explanation*: Do not assume that a patient requiring insulin use must have type 1 diabetes. The default for diabetes without further specification defaults to type 2. Add the code for long-term use of insulin.

# 4) Diabetes mellitus in pregnancy and gestational diabetes

See Section I.C.15. Diabetes mellitus in pregnancy. See Section I.C.15. Gestational (pregnancy induced) diabetes

5) Complications due to insulin pump malfunction

# (a) Underdose of insulin due to insulin pump failure

An underdose of insulin due to an insulin pump failure should be assigned to a code from subcategory T85.6, Mechanical complication of other specified internal and external prosthetic devices, implants and grafts, that specifies the type of pump malfunction, as the principal or first-listed code, followed by code T38.3X6-, Underdosing of insulin and oral hypoglycemic [antidiabetic] drugs. Additional codes for the type of diabetes mellitus and any associated complications due to the underdosing should also be assigned.

A 76-year-old male with diabetic ESRD is admitted for rehabilitation and develops hyperglycemia. He has had an insulin pump for 14 years and after returning from the ER, is noted to have experienced hyperglycemia due to a pump malfunction that caused too little insulin to be administered. On return, the blood sugars are resolving.

T85.614D Breakdown (mechanical) of insulin pump, subsequent encounter

T38.3X6D Underdosing of insulin and oral hypoglycemic [antidiabetic] drugs, subsequent encounter

Type 2 diabetes mellitus with hyperglycemia

E11.22 Type 2 diabetes mellitus with diabetic chronic kidney disease

# N18.6 End stage renal disease

Explanation: The complication code for the mechanical breakdown of the pump is sequenced first, followed by the underdosing code and code for the type of diabetes with complication. If other diabetic complications are present, assign all codes needed to capture each complication. The seventh character D is used for a subsequent encounter.

### (b) Overdose of insulin due to insulin pump failure

The principal or first-listed code for an encounter due to an insulin pump malfunction resulting in an overdose of insulin, should also be T85.6-, Mechanical complication of other specified internal and external prosthetic devices, implants and grafts, followed by code T38.3X1-, Poisoning by insulin and oral hypoglycemic [antidiabetic] drugs, accidental (unintentional).

A 70-year-old female with type 2 diabetes is found on the floor and sent to the ER. On her return from the hospital, it is noted that a failure of her insulin pump that caused excess insulin administration was the cause of hypoglycemia, resulting in her condition.

T85.614D Breakdown (mechanical) of insulin pump, subsequent encounter

T38.3X1D Poisoning by insulin and oral hypoglycemic [antidiabetic] drugs, accidental (unintentional), subsequent encounter

E11.649 Type 2 diabetes mellitus with hypoglycemia without coma

Explanation: The complication code for the mechanical breakdown of the pump is sequenced first, followed by the code for poisoning and type of diabetes with any associated complications. The seventh character D is used for subsequent encounter.

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			182.6Ø3	Acute embolism and thrombosis of unspecified veins of left upper extremity  Acute embolism and thrombosis of unspecified veins of upper extremity, bilateral  Acute embolism and thrombosis of unspecified veins of unspecified upper		√6 <sup>th</sup>	182.72	Chronic embolism and thrombosis of deep veins of upper extremity Chronic embolism and thrombosis of brachial vein Chronic embolism and thrombosis of radial vein Chronic embolism and thrombosis of ulnar vein 182.721 Chronic embolism and thrombosis of deep veins of right upper extremity
	√6 <sup>th</sup>	I82.61	Acute en Acute en Acute en	extremity bolism and thrombosis of superficial veins extremity mbolism and thrombosis of antecubital vein mbolism and thrombosis of basilic vein mbolism and thrombosis of cephalic vein				I82.722 Chronic embolism and thrombosis of deep veins of left upper extremity I82.723 Chronic embolism and thrombosis of deep veins of upper extremity, bilateral I82.729 Chronic embolism and thrombosis of
			I82.611	Acute embolism and thrombosis of superficial veins of right upper extremity	√5th	182.A	Embolisi	deep veins of unspecified upper extremity RP RIC mand thrombosis of axillary vein
			182.612	Acute embolism and thrombosis of		√6 <sup>th</sup>		Acute embolism and thrombosis of axillary vein
				superficial veins of left upper		, ,	102.741	182.A11 Acute embolism and thrombosis of right
			I82.613	extremity  Acute embolism and thrombosis of superficial veins of upper extremity, bilateral				axillary vein RC  182.A12 Acute embolism and thrombosis of left axillary vein RC
			182.619	Acute embolism and thrombosis of superficial veins of unspecified upper extremity				I82.A13 Acute embolism and thrombosis of axillary vein, bilateral  I82.A19 Acute embolism and thrombosis of
	√6th	182.62	Acute em	bolism and thrombosis of deep veins of				unspecified axillary vein
			<b>upper ex</b> t Acute ex			√6th	182.A2	Chronic embolism and thrombosis of axillary vein 182.A21 Chronic embolism and thrombosis of right axillary vein
			Acute e	mbolism and thrombosis of ulnar vein				182.A22 Chronic embolism and thrombosis of left
			<b>I82.621</b>	Acute embolism and thrombosis of deep				axillary vein RIC
			102 622	veins of right upper extremity				182.A23 Chronic embolism and thrombosis of
			102.022	Acute embolism and thrombosis of deep veins of left upper extremity				axillary vein, bilateral  182.A29 Chronic embolism and thrombosis of
			182.623	Acute embolism and thrombosis of deep				unspecified axillary vein
				veins of upper extremity, bilateral RIC	√5 <sup>th</sup>	182.B	Embolis	m and thrombosis of subclavian vein
			182.629	Acute embolism and thrombosis of deep veins of unspecified upper		√6 <sup>th</sup>	182.B1	Acute embolism and thrombosis of subclavian vein
				extremity				182.B11 Acute embolism and thrombosis of right
√5 <sup>th</sup>	182.7	extremit	y	and thrombosis of veins of upper de, if applicable, for associated long-term				subclavian vein  I82.B12 Acute embolism and thrombosis of left subclavian vein
				of anticoagulants (Z79.Ø1)				182.B13 Acute embolism and thrombosis of
	(O1)	EXCLUDE	<b>§1</b> perso	nal history of venous embolism and thrombosis (Z86.718)				subclavian vein, bilateral  RE2.B19 Acute embolism and thrombosis of unspecified subclavian vein
	√6 <sup>th</sup>	182.70		embolism and thrombosis of unspecified upper extremity		$\sqrt{6}^{\text{th}}$	182.B2	Chronic embolism and thrombosis of subclavian
				Chronic embolism and thrombosis of unspecified veins of right upper extremity				vein 182.B21 Chronic embolism and thrombosis of right subclavian vein
			182.702	Chronic embolism and thrombosis of unspecified veins of left upper extremity				I82.B22 Chronic embolism and thrombosis of left subclavian vein I82.B23 Chronic embolism and thrombosis of
			182.7Ø3	Chronic embolism and thrombosis of unspecified veins of upper extremity, bilateral				subclavian vein, bilateral  182.B29 Chronic embolism and thrombosis of unspecified subclavian vein
			182.709		√5 <sup>th</sup>	182.C	Embolis	m and thrombosis of internal jugular vein
				unspecified veins of unspecified upper		$\sqrt{6}^{th}$	182.C1	Acute embolism and thrombosis of internal jugular
	$\sqrt{6}^{\text{th}}$	I82.71		extremity RP RIC embolism and thrombosis of superficial upper extremity				vein I82.C11 Acute embolism and thrombosis of right internal jugular vein
				embolism and thrombosis of antecubital vein				182.C12 Acute embolism and thrombosis of left
			Chronic	embolism and thrombosis of basilic vein embolism and thrombosis of cephalic vein Chronic embolism and thrombosis of				internal jugular vein  I82.C13 Acute embolism and thrombosis of internal jugular vein, bilateral
			I82.712	superficial veins of right upper extremity  Chronic embolism and thrombosis of				I82.C19 Acute embolism and thrombosis of unspecified internal jugular vein
				superficial veins of left upper		$\sqrt{6}^{\text{th}}$	182.C2	Chronic embolism and thrombosis of internal
			I82.713	extremity Chronic embolism and thrombosis of				jugular vein I82.C21 Chronic embolism and thrombosis of
				superficial veins of upper extremity, bilateral				right internal jugular vein
			182.719	Chronic embolism and thrombosis of				182.C22 Chronic embolism and thrombosis of left internal jugular vein
				superficial veins of unspecified upper				182.C23 Chronic embolism and thrombosis of
				extremity RP RIC				internal jugular vein, bilateral
								l82.C29 Chronic embolism and thrombosis of unspecified internal jugular
								vein RP RIC
Non-ther	apy Anci	llary <b>S</b>	Speech-La	anguage Pathology Return To Provider	RIC DX	PDx Pi	rimary Dx	RIC CC Condition Manifestation Unspecified Dx ICD-10-CM 2023

. I – IVI 12.3/ I			Chapter 13. Diseases	or the muscu	IIOSK	teletai 3yst	eiii aiic	Connect	tive rissue ICD-10-CW 2023
√5 <sup>™</sup> M12.1							$\sqrt{6}^{th}$	M12.25	Villonodular synovitis (pigmented), hip
			hrosis deformans endemic Beck disease, unspecified		SIC.				M12.251 Villonodular synovitis (pigmented), right hip
√6 <sup>th</sup>		Kaschin-l	Beck disease, shoulder						M12.252 Villonodular synovitis (pigmented), left
		M12.111	Kaschin-Beck disease, r shoulder		RIC				M12.259 Villonodular synovitis (pigmented), unspecified hip
		M12.112	Kaschin-Beck disease, le shoulder		RIC		$\sqrt{6}$ <sup>th</sup>	M12.26	Villonodular synovitis (pigmented), knee
		M12.119	Kaschin-Beck disease, u shoulder	_					M12.261 Villonodular synovitis (pigmented), right knee
√6 <sup>th</sup>	M12.12	Kaschin-l	Beck disease, elbow						M12.262 Villonodular synovitis (pigmented), left
		M12.121	Kaschin-Beck disease, r	<b>J</b>	RIC				knee M12.269 Villonodular synovitis (pigmented),
			Kaschin-Beck disease, le Kaschin-Beck disease, u		RIC				unspecified knee RP RIC
_			elbow	RP F	RIC		√6th	M12.27	Villonodular synovitis (pigmented), ankle and foot M12.271 Villonodular synovitis (pigmented), right
√6 <sup>th</sup>	M12.13		Beck disease, wrist Kaschin-Beck disease, r	iahtuwist <b>[</b>	RIC				ankle and foot
			Kaschin-Beck disease, le		RIC				M12.272 Villonodular synovitis (pigmented), left
		M12.139	Kaschin-Beck disease, u wrist	nspecified	RIC				M12.279 Villonodular synovitis (pigmented),
√6 <sup>th</sup>	M12.14	Kaschin-l	Beck disease, hand					M12.28	unspecified ankle and foot  Villonodular synovitis (pigmented), other specified
			Kaschin-Beck disease, r	3	RIC				villonodular synovitis (pigmented), vertebrae
			Kaschin-Beck disease, le Kaschin-Beck disease, u		RIC			M12.29	Villonodular synovitis (pigmented), multiple
(Csb.	M12.15	Vacabin	hand Beck disease, hip	RP F	RIC	/5th	M12.0	Dalindra	sites omic rheumatism
<b>V</b> 0	W112.13		Kaschin-Beck disease, ri	ight hin	RIC	, ,	WILZ		udden and recurring attacks of moderate to severe joint
			Kaschin-Beck disease, le		RIC			pain ar	nd swelling generally occurring in the hands or feet of
		M12.159	Kaschin-Beck disease, u					normal	wn etiology. After the attack subsides, the joints appear
√6 <sup>th</sup>	M12.16	Kaschin-l	hip Beck disease, knee	RP F	NIC				Palindromic rheumatism, unspecified
			Kaschin-Beck disease, r	ight knee	RIC		√6 <sup>th</sup>	M12.31	Palindromic rheumatism, shoulder
			Kaschin-Beck disease, le		RIC		7		M12.311 Palindromic rheumatism, right
		WI 12.109	Kaschin-Beck disease, u knee	nspecified RP	RIC			,	shoulder M12.312 Palindromic rheumatism, left
√6 <sup>th</sup>	M12.17		Beck disease, ankle and f						shoulder
		M12.1/1	Kaschin-Beck disease, r foot	ignt ankle and	RIC				M12.319 Palindromic rheumatism, unspecified shoulder
		M12.172	Kaschin-Beck disease, le		RIC		√6th	M12.32	Palindromic rheumatism, elbow
		M12.179	Kaschin-Beck disease, u						M12.321 Palindromic rheumatism, right elbow
	M12.18	Kaschin-l	and foot Beck disease, vertebrae	RP 5	RIC		Ť		M12.322 Palindromic rheumatism, left elbow
			Beck disease, multiple si	tes	RIC				M12.329 Palindromic rheumatism, unspecified
			vitis (pigmented) ular synovitis (pigmente	d) unenecified	.		√6th	M12.33	elbow RP RIC Palindromic rheumatism, wrist
		site		RP F	RIC.				M12.331 Palindromic rheumatism, right
√6 <sup>th</sup>	M12.21		ular synovitis (pigmente Villonodular synovitis (p		ht				wrist M12.332 Palindromic rheumatism, left
			shoulder		RIC				wrist
		M12.212	Villonodular synovitis ( shoulder	pigmented), le	ft RIC				M12.339 Palindromic rheumatism, unspecified wrist RP RIC
		M12.219	Villonodular synovitis ( unspecified shoulder	pigmented),	ole.		$\sqrt{6}$ <sup>th</sup>	M12.34	Palindromic rheumatism, hand
√6 <sup>th</sup>	M12.22	Villonodi	ular synovitis (pigmente						M12.341 Palindromic rheumatism, right hand
		M12.221	Villonodular synovitis (pelbow	oigmented), rig	jht ic				M12.342 Palindromic rheumatism, left hand
		M12.222	Villonodular synovitis (						M12.349 Palindromic rheumatism, unspecified
		M12.229	elbow Villonodular synovitis (	_	RIC		√6th	M12.35	hand RP RIC Palindromic rheumatism, hip
_			unspecified elbow	RP F	RIC				M12.351 Palindromic rheumatism, right
√6 <sup>th</sup>	M12.23		ular synovitis (pigmente Villonodular synovitis (p		ıht				hip M12.352 Palindromic rheumatism, left hip
			wrist	F	RIC				M12.359 Palindromic rheumatism, unspecified
		M12.232	Villonodular synovitis (		ft		√6 <sup>th</sup>	M12.36	hip Palindromic rheumatism, knee
		M12.239	Villonodular synovitis (	pigmented),					M12.361 Palindromic rheumatism, right
√6 <sup>th</sup>	M12.24	Villonod	unspecified wrist ular synovitis (pigmente	d), hand	(IC				knee M12.362 Palindromic rheumatism, left
			Villonodular synovitis (p	igmented), rig					knee
		M12.242	hand Villonodular synovitis (		ft				M12.369 Palindromic rheumatism, unspecified knee
			hand Villonodular synovitis (		RIC		√6 <sup>th</sup>	M12.37	Palindromic rheumatism, ankle and foot
		W 12.249	unspecified hand	pigmented), RP [	RIC				M12.371 Palindromic rheumatism, right ankle and foot
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CD-10-C	W 2023			Chapter 19. Injury, Poisoning and Certain	n Other Con	sequen	ces of Ex	ternal Ca	uses <b>\$89-\$89.3</b> 09
√4 <sup>th</sup> S89	Other	and unsp	ecified in	juries of lower leg			√7 <sup>th</sup>	S89.119	Salter-Harris Type I physeal fracture of
				•				5031113	lower end of unspecified tibia
	NOT		closed.	indicated as open or closed should be coded		√6th	S89.12	Salter-Ha	rris Type II physeal fracture of lower end
				(6.1)				of tibia	
				specified injuries of ankle and foot (\$99)			√7 <sup>th</sup>	S89.121	Salter-Harris Type II physeal fracture of
				1,3; 2015,3Q,37-39 lignment" is not synonymous with malunion				C90 122	lower end of right tibia Salter-Harris Type II physeal fracture of
				when noted in provider documentation.			V 1	307.122	lower end of left tibia
	The	e appropria	ate 7th cha	racter is to be added to each code from			$\sqrt{7}$ th	S89.129	Salter-Harris Type II physeal fracture of
				9.1, S89.2, and S89.3.					lower end of unspecified tibia
	Α	initial end	counter for	closed fracture		√6 <sup>th</sup>	S89.13		rris Type III physeal fracture of lower end
	D			ter for fracture with routine healing				of tibia	fracture of medial malleolus (adult)
	G			ter for fracture with delayed healing				EXCLUBE	(S82.5-)
	K P			ter for fracture with nonunion ter for fracture with malunion			$\sqrt{7}$ th	S89.131	Salter-Harris Type III physeal fracture of
	S	seguela	.ne cheodii	ter for fracture with fridation					lower end of right tibia
√5¹	\$89.Ø	Physeal	fracture o	f upper end of tibia			√7 <sup>th</sup>	589.132	Salter-Harris Type III physeal fracture of lower end of left tibia
	505.5		019,4Q,56	apper end of tibid			√7 <sup>th</sup>	S89.139	Salter-Harris Type III physeal fracture of
	√6 <sup>th</sup>		, -,	ied physeal fracture of upper end of tibia					lower end of unspecified tibia
			•	Unspecified physeal fracture of upper end		√6 <sup>th</sup>	S89.14		rris Type IV physeal fracture of lower end
				of right tibia				of tibia	fracture of modial malloclus (adult)
		<b>√7</b> th	S89.ØØ2	Unspecified physeal fracture of upper end				EXCLUDES	fracture of medial malleolus (adult) (S82.5-)
			COO 646	of left tibia			<b>√7</b> th	589.141	Salter-Harris Type IV physeal fracture of
		<b>√7</b> <sup>th</sup>	289.009	Unspecified physeal fracture of upper end of unspecified tibia					lower end of right tibia
	√6th	S89.Ø1	Salter-Ha	nris Type I physeal fracture of upper end			$\sqrt{7}$ <sup>th</sup>	\$89.142	Salter-Harris Type IV physeal fracture of
			of tibia	, per projection and a supplied and			/70	500 140	lower end of left tibia
		√7 <sup>th</sup>	S89.Ø11	Salter-Harris Type I physeal fracture of			<b>√</b> T <sup>th</sup>	309.149	Salter-Harris Type IV physeal fracture of lower end of unspecified tibia
		/7th	COO #12	upper end of right tibia  Salter-Harris Type I physeal fracture of		√6th	\$89.19	Other ph	yseal fracture of lower end of tibia
		<b>√</b> 1 <sup>ca</sup>	309.012	upper end of left tibia				•	Other physeal fracture of lower end of
		<b>√7</b> th	S89.Ø19	Salter-Harris Type I physeal fracture of					right tibia
				upper end of unspecified tibia	\		<b>√</b> 7 <sup>th</sup>	589.192	Other physeal fracture of lower end of
	$\sqrt{6}^{th}$	S89.Ø2		arris Type II physeal fracture of upper end			-/7th	589 100	left tibia Other physeal fracture of lower end of
		./7th	of tibia	Salter-Harris Type II physeal fracture of				303.133	unspecified tibia
		V 1	307.021	upper end of right tibia	√5 <sup>th</sup>	<b>S89.2</b>	Physeal	fracture o	f upper end of fibula
		<b>√7</b> th	S89.Ø22	Salter-Harris Type II physeal fracture of			AHA:2	019,4Q,56	
		_		upper end of left tibia		6 <sup>th</sup>	S89.2Ø	Unspecif	ied physeal fracture of upper end of fibula
		<b>√7</b> <sup>th</sup>	589.029	Salter-Harris Type II physeal fracture of upper end of unspecified tibia			$\sqrt{7}$ <sup>th</sup>	S89.2Ø1	Unspecified physeal fracture of upper end
	√6th	S89.Ø3	Salter-Ha	arris Type III physeal fracture of upper end			(7th	C00 202	of right fibula Unspecified physeal fracture of upper end
			of tibia				V 1	307.202	of left fibula
		√7 <sup>th</sup>	S89.Ø31	Salter-Harris Type III physeal fracture of upper end of right tibia	_		$\sqrt{7}$ <sup>th</sup>	S89.2Ø9	Unspecified physeal fracture of upper end
		<b>√7</b> th	\$89,032	Salter-Harris Type III physeal fracture of		_			of unspecified fibula
				upper end of left tibia		√6 <sup>th</sup>	389.21	of fibula	rris Type I physeal fracture of upper end
		<b>√7</b> th	\$89.039	Salter-Harris Type III physeal fracture of			$\sqrt{7}$ <sup>th</sup>		Salter-Harris Type I physeal fracture of
	√6th	C90 Ø4	Caltor-Us	upper end of unspecified tibia RP RIC Irris Type IV physeal fracture of upper end					upper end of right fibula
	V	302.04	of tibia	ints Type IV physear fracture of appel end			<b>√7</b> <sup>th</sup>	589.212	Salter-Harris Type I physeal fracture of upper end of left fibula
		√7ub	\$89.041	Salter-Harris Type IV physeal fracture of			√7 <u>th</u>	\$89.219	upper end of left fibula Salter-Harris Type I physeal fracture of
			500 F	upper end of right tibia					upper end of unspecified
		<b>√7</b> th	589.042	Salter-Harris Type IV physeal fracture of upper end of left tibia					fibula RP RIC
		√7th	\$89,049	Salter-Harris Type IV physeal fracture of		$\sqrt{6}$ <sup>th</sup>	589.22	Salter-Ha of fibula	rris Type II physeal fracture of upper end
				upper end of unspecified tibia			√7 <sup>th</sup>		Salter-Harris Type II physeal fracture of
	√6 <sup>th</sup>	\$89.09	Other ph	yseal fracture of upper end of tibia					upper end of right fibula
		<b>√7</b> th	S89.Ø91	Other physeal fracture of upper end of			$\sqrt{7}$ <sup>th</sup>	\$89.222	Salter-Harris Type II physeal fracture of
			500 500	right tibia Ric				C00 220	upper end of left fibula
		√7 <sup>th</sup>	589.092	Other physeal fracture of upper end of left tibia			√7 <sup>th</sup>	589.229	Salter-Harris Type II physeal fracture of upper end of unspecified
		√7 <sup>th</sup>	589.099	Other physeal fracture of upper end of					fibula RP RIC
				unspecified tibia		$\sqrt{6}^{th}$	S89.29	Other ph	yseal fracture of upper end of fibula
√5 <sup>t</sup>	S89.1	Physeal	fracture o	f lower end of tibia			$\sqrt{7}$ <sup>th</sup>	S89.291	Other physeal fracture of upper end of
		AHA: 2	019,4Q,56				_		right fibula
	$\sqrt{6}^{th}$	S89.1Ø	Unspecif	ied physeal fracture of lower end of tibia			$\sqrt{7}$ <sup>th</sup>	589.292	Other physeal fracture of upper end of left fibula
		<b>√7</b> th	S89.1Ø1	Unspecified physeal fracture of lower end			√7 <sup>th</sup>	S89.299	Other physeal fracture of upper end of
		√7th	\$89,102	of right tibia Unspecified physeal fracture of lower end					unspecified fibula
		-		of left tibia	√5 <sup>th</sup>	<b>S89.3</b>	•		f lower end of fibula
		$\sqrt{7}$ <sup>th</sup>	\$89.109	Unspecified physeal fracture of lower end				019,4Q,56	
	√6 <sup>th</sup>	CQO 11	Calton He	of unspecified tibia		√6 <sup>th</sup>			ied physeal fracture of lower end of fibula
	<b>4</b> 0™	307.11	of tibia	rris Type I physeal fracture of lower end			√7 <sup>th</sup>	589.301	Unspecified physeal fracture of lower end of right fibula
		$\sqrt{7}^{\text{th}}$		Salter-Harris Type I physeal fracture of			$\sqrt{7}$ th	S89.3Ø2	Unspecified physeal fracture of lower end
		/7/4	COO 112	lower end of right tibia					of left fibula
		✓ Im	309.112	Salter-Harris Type I physeal fracture of lower end of left tibia			$\sqrt{7}$ <sup>th</sup>	589.309	Unspecified physeal fracture of lower end
									of unspecified fibula

# **Appendixes**

# **Appendix A: Valid 3-character ICD-10-CM Codes**

		I	
AØ9	Infectious gastroenteritis and colitis, unspecified	E43	Unspecified severe protein-calorie malnutrition
A33	Tetanus neonatorum	E45	Retarded development following protein-calorie malnutrition
A34	Obstetrical tetanus	E46	Unspecified protein-calorie malnutrition
A35	Other tetanus	E52	Niacin deficiency [pellagra]
A46	Erysipelas	E54	Ascorbic acid deficiency
A55	Chlamydial lymphogranuloma (venereum)	E58	Dietary calcium deficiency
A57	Chancroid	E59	Dietary selenium deficiency
A58	Granuloma inguinale	E6Ø	Dietary zinc deficiency
A64	Unspecified sexually transmitted disease	E65	Localized adiposity
A65	Nonvenereal syphilis	E68	Sequelae of hyperalimentation
A7Ø	Chlamydia psittaci infections	FØ4	Amnestic disorder due to known physiological condition
A78	Q fever	FØ5	Delirium due to known physiological condition
A86	Unspecified viral encephalitis	FØ9	Unspecified mental disorder due to known physiological condition
A89	Unspecified viral infection of central nervous system	F21	Schizotypal disorder
A9Ø	Dengue fever [classical dengue]	F22	Delusional disorders
A91	Dengue hemorrhagic fever	F23	Brief psychotic disorder
A94	Unspecified arthropod-borne viral fever	F24	Shared psychotic disorder
A99	Unspecified viral hemorrhagic fever	F28	Other psychotic disorder not due to a substance or known physiological
BØ3	Smallpox	F20	condition
BØ4	Monkeypox	F29	Unspecified psychosis not due to a substance or known physiological condition
BØ9	Unspecified viral infection characterized by skin and mucous membrane	F39	Unspecified mood [affective] disorder
Dα	lesions	F42	Obsessive-compulsive disorder
B2Ø B49	Human immunodeficiency virus [HIV] disease	F54	Psychological and behavioral factors associated with disorders or
В54	Unspecified mycosis	134	diseases classified elsewhere
B59	Unspecified malaria	F59	Unspecified behavioral syndromes associated with physiological
B64	Pneumocystosis Unspecified protozoal disease	. 3.	disturbances and physical factors
B72	Dracunculiasis	F66	Other sexual disorders
B75	Trichinellosis	F69	Unspecified disorder of adult personality and behavior
B79	Trichuriasis	F7Ø	Mild intellectual disabilities
B8Ø	Enterobiasis	F71	Moderate intellectual disabilities
B86	Scabies	F72	Severe intellectual disabilities
B89	Unspecified parasitic disease	F73	Profound intellectual disabilities
B91	Sequelae of poliomyelitis	F79	Unspecified intellectual disabilities
B92	Seguelae of leprosy	F82	Specific developmental disorder of motor function
CØ1	Malignant neoplasm of base of tongue	F88	Other disorders of psychological development
CØ7	Malignant neoplasm of parotid gland	F89	Unspecified disorder of psychological development
C12	Malignant neoplasm of pyriform sinus	F99	Mental disorder, not otherwise specified
C19	Malignant neoplasm of rectosigmoid junction	GØ1	Meningitis in bacterial diseases classified elsewhere
C2Ø	Malignant neoplasm of rectum	GØ2	Meningitis in other infectious and parasitic diseases classified elsewhere
C23	Malignant neoplasm of gallbladder	GØ7	Intracranial and intraspinal abscess and granuloma in diseases classified
C33	Malignant neoplasm of trachea		elsewhere
C37	Malignant neoplasm of thymus	GØ8	Intracranial and intraspinal phlebitis and thrombophlebitis
C52	Malignant neoplasm of vagina	GØ9	Sequelae of inflammatory diseases of central nervous system
C55	Malignant neoplasm of uterus, part unspecified	G1Ø	Huntington's disease
C58	Malignant neoplasm of placenta	G14	Postpolio syndrome
C61	Malignant neoplasm of prostate	G2Ø	Parkinson's disease
C73	Malignant neoplasm of thyroid gland	G26	Extrapyramidal and movement disorders in diseases classified elsewhere
D34	Benign neoplasm of thyroid gland	G35	Multiple sclerosis
D45	Polycythemia vera	G53	Cranial nerve disorders in diseases classified elsewhere
D62	Acute posthemorrhagic anemia	G55 G59	Nerve root and plexus compressions in diseases classified elsewhere  Mononeuropathy in diseases classified elsewhere
D65	Disseminated intravascular coagulation [defibrination A syndrome]	G63	·
D66	Hereditary factor VIII deficiency	G64	Polyneuropathy in diseases classified elsewhere Other disorders of peripheral nervous system
D67	Hereditary factor IX deficiency	G94	Other disorders of peripheral nervous system  Other disorders of brain in diseases classified elsewhere
D71	Functional disorders of polymorphonuclear neutrophils	H22	Disorders of iris and ciliary body in diseases classified elsewhere
D77	Other disorders of blood and blood-forming organs in diseases classified	H28	Cataract in diseases classified elsewhere
	elsewhere	H32	Chorioretinal disorders in diseases classified elsewhere
EØ2	Subclinical iodine-deficiency hypothyroidism	пз2 Н36	Retinal disorders in diseases classified elsewhere
E15	Nondiabetic hypoglycemic coma	H42	Glaucoma in diseases classified elsewhere
E35	Disorders of endocrine glands in diseases classified elsewhere	100	Rheumatic fever without heart involvement
E4Ø	Kwashiorkor	I1Ø	Essential (primary) hypertension
E41	Nutritional marasmus	132	Pericarditis in diseases classified elsewhere
E42	Marasmic kwashiorkor		· · · · · · · · · · · · · · · · · · ·

ICD-10-CM 2023 Appendixes-1

Code	Title	Tier	RIC Excl	Code	Title
B18.2	Chr viral hep C	3	03	C95.Ø2	Ac leuk of unsp cell type, in
B19.Ø	Unsp viral hep w hep coma	3	03	D57.Ø-	Hb-SS w crisis
B19.11	Unsp viral hep B w hep coma	3	03	D57.1	Sickle-cell wo crisis
B19.21	Unsp viral hep C w hep coma	3	03	D57.20	Sickle-cell/Hb-C wo crisis
B2Ø	HIV	3		D57.21-	Sickle-cell/Hb-C w crisis
B25.Ø	Cytomegaloviral pneumonitis	3	15	D57.411	Sickle-cell thal unsp w ac ch
B26.1	Mumps meningitis	3	03, 05	D57.8Ø	Oth sickle-cell d/o wo crisis
B26.2	Mumps encephalitis	3	03	D57.81-	Oth sickle-cell d/o w crisis
B26.3	Mumps pancreatitis	3		D60	Acq pure red cell aplasia
B33.3	Retrovirus infections, nec	3		D61	Oth aplastic anemia/oth bo
B37.1	Pulmonary candidiasis	3	15	D64.81	Anemia d/t antineo chemo
B37.5	Candidal meningitis	3	03, 05	D65	DIC
B37.6	Candidal endocarditis	3	14	D66	Hereditary factor VIII deficie
B37.7	Candidal sepsis	3		D67	Hereditary factor IX deficien
B37.81	Candidal esophagitis	3	02.05	EØ8.2-	DM d/t underlying conditio complications
B38.4	Coccidioidomycosis meningitis	3	03, 05	EØ8.31-	DM d/t underlying condition
B39.0	Ac pulmonary histoplasm capsulati  Chr pulmonary histoplasm capsulati	3	15		retinopathy
B39.1 B39.2	Pulm histoplasm capsulati, unsp	3	15 15	EØ8.32-	DM d/t underlying condition
B39.2	Disseminated histoplasm capsulati	3	15	F60.22	diabetic retinopathy
B39.4	Histoplasm capsulati, unsp	3	15	EØ8.33-	DM d/t underlying conditio diabetic retinopathy
B39.5	Histoplasm duboisii	3	13	EØ8.34-	DM d/t underlying condition
B39.9	Histoplasm, unsp	3	1,5		diabetic retinopathy
B44.Ø	Invasive pulm aspergillosis	3	15	EØ8.351-	DM d/t underlying conditio
B45.1	Cerebral cryptococcosis	3	03, 05	F60 350	retinopathy w macular ede
B58.2	Toxoplasma meningoencephalitis	3	03,05	EØ8.359-	DM d/t underlying condition retinopathy w/o macular education
B58.3	Pulmonary toxoplasmosis	3	15	EØ8.36	DM d/t underlying conditio
B58.81	Toxoplasma myocarditis	3	14	EØ8.39	DM d/t underlying conditio
B59	Pneumocystosis	3	15		compl
B77.81	Ascariasis pneumonia	3	15	EØ8.40	DM d/t underlying conditio
B94.1	Sequelae of viral encephalitis	3	03	EØ8.41	DM d/t underlying conditio
B96.5	Pseudomonas as the cause of dis class elwh	2		EØ8.42	DM d/t underlying conditio
B97.3Ø	Unsp retrovirus as the cause of dis class elwh	3		EØ8.43	DM d/t underlying conditio (poly)neuro
B97.31	Lentivirus as the cause of dis class elwh	3		EØ8.44	DM d/t underlying conditio
B97.32	Oncovirus as the cause of dis class elwh	3		EØ8.49	DM d/t underlying conditio
B97.33	HTLV-I as the cause of dis class elwh	3	06		complication
B97.34	HTLV-II as the cause of dis class elwh	3	06	EØ8.5-	DM d/t underlying conditio
B97.35	HIV 2 as the cause of dis class elwh	3		F@0 61	complications  DM d/t underlying conditio
B97.39	Oth retrovirus as cause of dis class elwh	3		EØ8.61-	DM d/t underlying condition
C91.ØØ	Ac lymphoblastic leuk no remis	3		EØ8.63-	DM d/t underlying condition
C91.Ø2	Ac lymphoblastic leuk, in relapse	3		EØ8.649	DM d/t underlying condition
C92.ØØ	Ac myeloblastic leuk, no remis	3		L00.047	coma
C92.Ø2	Ac myeloblastic leuk, in relapse	3		EØ8.65	DM d/t underlying conditio
C92.40	Ac promyelocytic leuk, no remis  Ac promyelocytic leuk, in relapse	3	<u> </u>	EØ8.69	DM d/t underlying conditio
C92.42 C92.5Ø	Ac myelomonocytic leuk, no remis	3			complication
C92.52	Ac myelomonocytic leuk, in relapse	3		EØ9.2-	Drug/chem induced DM w
C92.6Ø	Ac myeloid leuk w 11q23-abnormality no remis	3	<del>                                     </del>	EØ9.31-	Drug/chem induced DM w retinopathy
C92.62	Ac myeloid leuk w 11q23-abnormality in relapse	3		EØ9.32-	Drug/chem induced DM w
C92.AØ	Ac myeloid leuk w multilineage dysplasia, no remis	3			retinopathy
C92.A2	Ac myeloid leuk w multilineage dysplasia, in relapse	3		EØ9.33-	Drug/chem induced DM w
C93.ØØ	Ac monoblastic/monocytic leuk, no remis	3		F60 2 :	retinopathy
C93.Ø2	Ac monoblastic/monocytic leuk, in relapse	3	<u> </u>	EØ9.34-	Drug/chem induced DM w s retinopathy
C94.00	Ac erythroid leuk, no remis	3		EØ9.351-	Drug/chem induced DM w
C94.02	Ac erythroid leuk, in relapse	3			retinopathy w macular ede
C95.ØØ	Ac leuk of unsp cell type no remis	3		EØ9.359-	Drug/chem induced DM w
-					retinopathy w/o macular ed

D57.02	
D57.1 Sickle-cell wo crisis D57.20 Sickle-cell/Hb-C wo crisis D57.21 Sickle-cell/Hb-C w crisis D57.21 Sickle-cell thal unsp w ac chest synd D57.411 Sickle-cell thal unsp w ac chest synd D57.80 Oth sickle-cell d/o wo crisis D57.81 Oth sickle-cell d/o w crisis D60. Acq pure red cell aplasia D61. Oth aplastic anemia/oth bone marrow fail syndr D64.81 Anemia d/t antineo chemo D65 DIC D66 Hereditary factor VIII deficiency D67 Hereditary factor IX deficiency D68.2- DM d/t underlying condition w kidney complications E68.31- DM d/t underlying condition w unsp diabetic retinopathy E68.32- DM d/t underlying condition w mod nonprolif diabetic retinopathy E68.33- DM d/t underlying condition w severe nonprolif diabetic retinopathy E68.34- DM d/t underlying condition w prolif diabetic retinopathy w macular edema E68.35- DM d/t underlying condition w cataract S68.36 DM d/t underlying condition w cataract S68.37- DM d/t underlying condition w cataract S68.38- DM d/t underlying condition w cataract S68.39- DM d/t underlying condition w cataract S68.39- DM d/t underlying condition w cataract S68.39- DM d/t underlying condition w diabetic neuro, unsp S68.40 DM d/t underlying condition w diabetic neuro, unsp S68.41 DM d/t underlying condition w diabetic neuro, unsp S68.42 DM d/t underlying condition w diabetic neuro, unsp S68.43- DM d/t underlying condition w diabetic neuro, unsp S68.44- DM d/t underlying condition w diabetic neuro, unsp S68.43- DM d/t underlying condition w diabetic neuro, unsp	
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ICD-10-CM 2023 Appendixes – 49

ICD-10-CM 2023 Illustrations

# **Arteries**

