# **Optum**



# ICD-10-CM Professional for Hospitals

The complete official code set

Codes valid from October 1, 2024 through September 30, 2025



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#### **Age Edits**

#### Newborn Age: 0

These diagnoses are intended for newborns and neonates and the patient's age must be 0 years.

N47.Ø Adherent prepuce, newborn

#### Pediatric Age: 0-17

These diagnoses are intended for children and the patient's age must be between 0 and 17 years.

L21.1 Seborrheic infantile dermatitis

#### Maternity Age: 9-64

These diagnoses are intended for childbearing patients between the age of 9 and 64 years.

#### Adult Age: 15-124

These diagnoses are intended for patients between the age of 15 and 124 years.

# R54 Age-related physical debility Frailty Old age Senescence Senile asthenia Senile debility EXGLUDISS\*1 age-related cognitive decline (R41.81) sarcopenia (M62.84) senile psychosis (FØ3) senility NOS (R41.81)

#### **Sex Edits**

#### ♂ Male diagnosis only

Q98.Ø Klinefelter syndrome karyotype 47, XXY

Female diagnosis only

N35.12 Postinfective urethral stricture, not elsewhere classified, female

#### H1-H14 Hospital Acquired Condition (HAC)

These icons identify codes that are high-cost and/or high-volume (CC or MCC) that when assigned as a secondary diagnosis result in assignment of a case to a higher-paying MS-DRG. The condition or diagnosis represented by these codes is considered reasonably preventable through the application of evidence-based guidelines. If the condition is not present on admission (meaning it developed during the hospital admission), the case will not group to the higher-paying MS-DRG based solely upon the reporting of the HAC code. Many of these HACs are conditional and are based on reporting the specific diagnosis code(s) in combination with certain procedure codes.

**Note:** Hospital-acquired conditions do not impact MS-LTC-DRG assignment.

N15.1 Renal and perinephric abscess

#### CC Condition

This icon identifies a complication or comorbidity diagnosis that may affect DRG assignment. A complication or comorbidity diagnosis, CC condition, is defined as a significant acute disease, a significant acute manifestation of a chronic disease, an advanced or end-stage chronic disease, or a chronic disease associated with systemic physiological decompensation and debility that have consistently greater impact on hospital resources.

G90.59 Complex regional pain syndrome I of other specified site

#### MCC Condition

This icon identifies a major complication or comorbidity diagnosis that may affect DRG assignment. An MCC condition meets the same criteria as a CC condition but is associated with a higher acuity level and hospital resource consumption is expected to be higher than that for a CC condition. There are fewer conditions that meet the criteria as an MCC than those for a CC condition.

S35.238 Other injury of inferior mesenteric artery

**Note:** The assignment of an MS-DRG or MS-LTC-DRG often depends on the presence or absence of a secondary diagnosis code that is designated as an MCC or CC. However, in some instances the MCC or CC designation for that secondary diagnosis code is negated due to its relationship with the principal diagnosis; this is referred to as CC exclusion. The ICD-10 MS-DRG Definitions Manual included with the IPPS final rule provides a list of all principal diagnosis codes that would render ineffective the MCC/CC designation for a particular ICD-10-CM code when used as a secondary diagnosis. Optum has provided this CC exclusion list in an easily searchable data file, which can be accessed at the following:

https://www.optumcoding.com/ProductUpdates/ Title: "2024 ICD-10-CM for Hospitals CC Excludes Data File" Password: XXXXXX

#### **Unspecified Site**

This icon identifies codes that are considered an MCC or CC but lack specificity in regard to their anatomical location. The medical record documentation should be reviewed carefully, to ensure that no other code within the same category or subcategory can be assigned for greater specificity.

G81.00 Flaccid hemiplegia affecting unspecified side COLONS HOCO

#### Unacceptable Principal Diagnosis

This icon identifies codes that should not be assigned as principal diagnosis for *inpatient* admissions. Codes with an unacceptable principal diagnosis edit are considered supplementary — describing circumstances that influence an individual's health status or an additional code — identifying conditions that are not specific manifestations but may be due to an underlying cause.

T48.5X5 Adverse effect of other anti-common-cold drugs

#### HIV-related Condition

This icon identifies codes that are considered a major HIV-related diagnosis. When the condition is coded in combination with a diagnosis of human immunodeficiency virus (HIV), code B2Ø, the case will move from MS-DRG/MS-LTC-DRG 977 to MS-DRGs/MS-LTC-DRGs 974-976.

G96.9 Disorder of central nervous system, unspecified

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# **Conversion Table of ICD-10-CM Codes**

The FY 2025 (October 1,2024-September 30, 2025) Conversion Table for new ICD-10-CM codes is provided to assist users in data retrieval. For each new code the table shows its previously assigned code equivalent. Asterisks identify new codes added to the classification April 1, 2024.

Code Assignment Beginning 10/1/2023	Previous Code(s) Assignment										
A41.54	A41.59	G2Ø.C	G2Ø	H5Ø.671	H5Ø.69	M8Ø.ØB1G	M8Ø.ØAXG	NØ6.29	NØ6.2	T56.821D	T56.891D
B96.83	B96.89	G23.3	G23.8	H5Ø.672	H5Ø.69	M8Ø.ØB1K	M8Ø.ØAXK	026.641	026.611	T56.821S	T56.891S
D13.91	D13.9	G31.8Ø	G31.89	H5Ø.679	H5Ø.69	M8Ø.ØB1P	M8Ø.ØAXP	026.642	026.612	T56.822A	T56.892A
D13.99	D13.9	G31.86	G31.89	H5Ø.681	H5Ø.69	M8Ø.ØB1S	M8Ø.ØAXS	026.643	026.613	T56.822D	T56.892D
D48.110	D48.1	G37.81	G37.8	H5Ø.682	H5Ø.69	M8Ø.ØB2A	M8Ø.ØAXA	026.649	026.619	T56.822S	T56.892S
D48.111	D48.1	G37.89	G37.8	H5Ø.689	H5Ø.69	M8Ø.ØB2D	M8Ø.ØAXD	090.41	O9Ø.4	T56.823A	T56.893A
D48.112	D48.1	G4Ø.CØ1	G4Ø.3Ø1	H57.8A1	H57.89	M8Ø.ØB2G	M8Ø.ØAXG	090.49	O9Ø.4	T56.823D	T56.893D
D48.113	D48.1	G4Ø.CØ9	G4Ø.3Ø9	H57.8A2	H57.89	M8Ø.ØB2K	M8Ø.ØAXK	Q44.7Ø	Q44.7	T56.823S	T56.893S
D48.114	D48.1	G4Ø.C11	G4Ø.311	H57.8A3	H57.89	M8Ø.ØB2P	M8Ø.ØAXP	Q44.71	Q44.7	T56.824A	T56.894A
D48.115	D48.1	G4Ø.C19	G4Ø.319	H57.8A9	H57.89	M8Ø.ØB2S	M8Ø.ØAXS	Q44.79	Q44.7	T56.824D	T56.894D
D48.116	D48.1	G43.EØ1	G43.8Ø1	I1A.Ø	l1Ø-l15.9	M8Ø.ØB9A	M8Ø.ØAXA	Q75.ØØ1	Q75.Ø	T56.824S	T56.894S
D48.117	D48.1	G43.EØ9	G43.8Ø9	120.81	120.8	M8Ø.ØB9D	M8Ø.ØAXD	Q75.ØØ2	Q75.Ø	*T74.A1XA	T74.91XA
D48.118	D48.1	G43.E11	G43.811	120.89	120.8	M8Ø.ØB9G	M8Ø.ØAXG	Q75.ØØ9	Q75.Ø	*T74.A1XD	T74.91XD
D48.119	D48.1	G43.E19	G43.819	I21.B	I21.A9	M8Ø.ØB9K	M8Ø.ØAXK	Q75.Ø1	Q75.Ø	*T74.A1XS	T74.91XS
D48.19	D48.1	G9Ø.B	G9Ø.8	124.81	124.8	M8Ø.ØB9P	M8Ø.ØAXP	Q75.Ø21	Q75.Ø	*T74.A2XA	T74.92XA
D57.Ø4	D57.Ø9	G93.42	G93.49	124.89	124.8	M8Ø.ØB9S	M8Ø.ØAXS	Q75.Ø22	Q75.Ø	*T74.A2XD	T74.92XD
D57.214	D57.218	G93.43	G93.49	125.85	125.89	M8Ø.8B1A	M8Ø.8ØXA	Q75.Ø29	Q75.Ø	*T74.A2XS	T74.92XS
D57.414	D57.418	G93.44	G93.49	147.10	147.1	M8Ø.8B1D	M8Ø.8AXD	Q75.Ø3	Q75.Ø	*T76.A1XA	T76.91XA
D57.434	D57.438	H36.811	H35.2Ø-	147.11	147.1	M8Ø.8B1G	M8Ø.8AXG	Q75.Ø41	Q75.Ø	*T76.A1XD	T76.91XD
D57.454	D57.458		H35.23	147.19	147.1	M8Ø.8B1K	M8Ø.8AXK	Q75.Ø42	Q75.Ø	*T76.A1XS	T76.91XS
D57.814	D57.818	H36.812	H35.20- H35.23	J15.61	J15.6	M8Ø.8B1P	M8Ø.8AXP	Q75.Ø49	Q75.Ø	*T76.A2XA	T76.92XA
D61.Ø2	D61.09	H36.813	H35.20-	J15.69	J15.6	M8Ø.8B1S	M8Ø.8AXS	Q75.Ø51	Q75.Ø	*T76.A2XD	T76.92XD
D89.84	D89.89		H35.23	J44.81	J42	M8Ø.8B2A	M8Ø.8ØXA	Q75.Ø52	Q75.Ø	*T76.A2XS	T76.92XS
E2Ø.81Ø	E2Ø.8	H36.819	H35.2Ø-	J44.89	J42	M8Ø.8B2D	M8Ø.8AXD	Q75.Ø58	Q75.Ø	W44.8XXA	Codes in
E2Ø.811	E2Ø.8	U26 021	H35.23	J4A.Ø	J44.Ø-J44.1,	M8Ø.8B2G	M8Ø.8AXG	Q75.Ø8	Q75.Ø		Categories T15-T19
E2Ø.812	E2Ø.8	H36.821	H35.20- H35.23	144.0	J44.9	M8Ø.8B2K	M8Ø.8AXK	Q87.83	Q87.89	W44.8XXD	Codes in
E2Ø.818	E2Ø.8	H36.822	H35.20-	J4A.8	J44.Ø-J44.1; J44.9	M8Ø.8B2P	M8Ø.8AXP	Q87.84	Q87.89		Categories T15-T19
E2Ø.819	E2Ø.8		H35.23	J4A.9	J44.0-J44.1;	M8Ø.8B2S	M8Ø.8AXS	Q87.85	Q87.89	W44.8XXS	Codes in
E2Ø.89	E2Ø.8	H36.823	H35.20- H35.23		J44.9	M8Ø.8B9A	M8Ø.8ØXA	Q93.52	Q93.59	VV44.0AA3	Categories
E74.05	E74.Ø9	H36.829	H35.20-	K35.2ØØ	K35.2Ø	M8Ø.8B9D	M8Ø.8AXD	RØ9.AØ	RØ9.89		T15-T19
E75.27	E75.29	1130.025	H35.23	K35.2Ø1	K35.2Ø	M8Ø.8B9G	M8Ø.8AXG	RØ9.A1	RØ9.89	W44.9XXA	Codes in Categories
E75.28	E75.29	H36.89	H35.20-	K35.2Ø9	K35.2Ø	M8Ø.8B9K	M8Ø.8AXK	RØ9.A2	RØ9.89		T15-T19
E79.81	E79.8	-	H35.23	K35.21Ø	K35.21	M8Ø.8B9P	M8Ø.8AXP	RØ9.A9	RØ9.89	W44.9XXD	Codes in
E79.82	E79.8	-	H5Ø.69	K35.211	K35.21	M8Ø.8B9S	M8Ø.8AXS	R4Ø.2A	R4Ø.2Ø		Categories T15-T19
E79.89	E79.8	H5Ø.622	H50.69	K35.219	K35.21	NØ2.B1	NØ2.Ø-NØ2.1	R92.30	R92.2	W44.9XXS	Codes in
E88.43	E88.49	H5Ø.629	H5Ø.69	K63.8211	K63.89	NØ2.B2	NØ2.1	R92.311	R92.2		Categories T15-T19
E88.81Ø	E88.81	H5Ø.631	H5Ø.69		K63.89	NØ2.B3	NØ2.2	R92.312	R92.2	W44.AØXA	Codes in
E88.811	E88.81	H5Ø.632	H5Ø.69		K63.89	NØ2.B4	NØ2.2	R92.313	R92.2	VV44.AUAA	Categories
E88.818	E88.81	H5Ø.639	H5Ø.69	K63.822	K63.89	NØ2.B5	NØ2.3	R92.321	R92.2		T15-T19
E88.819	E88.81	H5Ø.641	H5Ø.69	K63.829	K63.89	NØ2.B6	NØ2.5	R92.322	R92.2	W44.AØXD	Codes in Categories
E88.A	R64	H5Ø.642	H5Ø.69		K68.9	NØ2.B9	NØ2.8	R92.323	R92.2		T15-T19
G11.5	G11.8 and	H5Ø.649	H5Ø.69	K68.3	K68.9	NØ4.2Ø	NØ4.2	R92.331	R92.2	W44.AØXS	Codes in
	E23.Ø and KØØ.Ø		H5Ø.69	K9Ø.821	K9Ø.89	NØ4.21	NØ4.2	R92.332	R92.2		Categories T15-T19
G11.6	G11.8	-	H5Ø.69	K9Ø.822	K9Ø.89	NØ4.22	NØ4.2	R92.333	R92.2	W44.A1XA	Codes in
G2Ø.A1	G2Ø	H5Ø.659	H5Ø.69	K9Ø.829	K9Ø.89	NØ4.29	NØ4.2	R92.341	R92.2	TT TT TANK	Categories
G2Ø.A1	G2Ø G2Ø	H5Ø.661	H5Ø.69	K9Ø.83	K9Ø.89	NØ6.2Ø	NØ6.2	R92.342	R92.2		T15-T19
G2Ø.A2 G2Ø.B1	G2Ø G2Ø	H5Ø.662	H5Ø.69	M8Ø.ØB1A	M8Ø.ØAXA	NØ6.21	NØ6.2	R92.343	R92.2	W44.A1XD	Codes in Categories
G2Ø.B1	G2Ø G2Ø	H5Ø.669	H5Ø.69	M8Ø.ØB1D	M8Ø.ØAXD	NØ6.22	NØ6.2	T56.821A	T56.891A		T15-T19
עבשוטב	UZW						_		_	_	

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Disorder — continued	Disorder — continued	Disorder — continued
binocular — continued	bone — continued	cannabis use
movement — continued	continuity — continued	mild F12.10
convergence	specified type — continued	with
excess H51.12	vertebra M84.88	cannabis intoxication
insufficiency H51.11	density and structure M85.9	with perceptual d
internuclear ophthalmoplegia — see Ophthalmo-	cyst — see also Cyst, bone, specified type NEC	without perceptual a
plegia, internuclear	aneurysmal — see Cyst, bone, aneurysmal	cannabis-induced
palsy of conjugate gaze H51.0	solitary — see Cyst, bone, solitary	anxiety disorder F
specified type NEC H51.8	diffuse idiopathic skeletal hyperostosis — see	psychotic disorder
vision NEC — see Disorder, vision, binocular		sleep disorder F12
bipolar (I) seasonal) (type I) F31.9	Hyperostosis, ankylosing	in remission (early) (sust
and related due to a known physiological condition	fibrous dysplasia (monostotic) — see Dysplasia,	moderate or severe F12.20
with	fibrous, bone	with
manic features FØ6.33	fluorosis — see Fluorosis, skeletal	cannabis intoxication
manic- or hypomanic-like episodes FØ6.33	hyperostosis of skull M85.2	with perceptual d
mixed features FØ6.34	osteitis condensans — <i>see</i> Osteitis, condensans	with perceptual of
current (or most recent) episode	specified type NEC M85.8-   ✓	cannabis-induced
	ankle M85.87- <b>☑</b>	anxiety disorder F
depressed F31.9 with psychotic features F31.5	foot M85.87- <b>☑</b>	psychotic disorder
with psychotic features F31.30	forearm M85.83- ✓	sleep disorder F12
mild F31.31	hand M85.84- <b>☑</b>	delirium F12.221
moderate F31.32	lower leg M85.86- <b>☑</b>	in remission (early) (sust
severe (without psychotic features) F31.4	multiple sites M85.89	carbohydrate
with psychotic features F31.5	neck M85.88	absorption, intestinal NEC E
hypomanic F31.0	rib M85.88	metabolism (congenital) E7
manic F31.9	shoulder M85.81- <b>☑</b>	specified NEC E74.89
with psychotic features F31.2	skull M85.88	cardiac, functional I51.89
without psychotic features F31.10	thigh M85.85- <b>☑</b>	carnitine metabolism E71.40
mild F31.11	upper arm M85.82-   ✓	cartilage M94.9
moderate F31.12	vertebra M85.88	articular NEC — see Derand
severe (without psychotic features)	development and growth NEC M89.20	cartilage
F31.13	carpus M89.24- <b>▼</b>	chondrocalcinosis — sec
with psychotic features F31.2	clavicle M89.21- <b>☑</b>	specified type NEC M94.8X-
mixed F31.60	femur M89.25- <b>☑</b>	
mild F31.61	fibula M89.26- <b>☑</b>	articular — see Derange cartilage
moderate F31.62	finger M89.24-	multiple sites M94.8XØ
severe (without psychotic features) F31.63	humerus M89.22-	catatonia (due to known physic
with psychotic features F31.64	ilium M89.28	another mental disorder
severe depression (without psychotic features)	ischium M89.28	catatonic
F31.4	metacarpus M89.24- ✓	due to (secondary to) knowr
with psychotic features F31.5	metatarsus M89.27- ✓	FØ6.1
II (type 2) F31.81	multiple sites M89.29	organic FØ6.1
in remission (currently) F31.70	neck M89.28	central auditory processing HS
in full remission	radius M89.23- ✓	cervical
most recent episode	rib M89.28	region NEC M53.82
depressed F31.76	scapula M89.21- ✓	root (nerve) NEC G54.2
hypomanic F31.72	skull M89.28	character NOS F6Ø.9
manic F31.74	tarsus M89.27- <b>∑</b>	childhood disintegrative NEC I
mixed F31.78	tibia M89.26- ☑	cholesterol and bile acid meta
in partial remission	toe M89.27- 🖸	Barth syndrome E78.71
most recent episode	ulna M89.23- ☑	other specified E78.79
depressed F31.75	vertebra M89.28	Smith-Lemli-Opitz syndrom
hypomanic F31.71	specified type NEC M89.8X- ■	choroid H31.9
manic F31.73	brachial plexus G54.0	atrophy — see Atrophy, ch
mixed F31.77	branched-chain amino-acid metabolism E71.2	degeneration — see Deger
organic FØ6.3Ø	specified NEC E71.19	detachment — see Detach
single manic episode F30.9	breast N64.9	dystrophy — see Dystrophy
mild F3Ø.11	agalactia — see Agalactia	hemorrhage — see Hemorr
moderate F3Ø.12	associated with	rupture — see Rupture, cho
severe (without psychotic symptoms) F3Ø.13	lactation O92.7Ø	scar — see Scar, chorioretir
with psychotic symptoms F30.2	specified NEC 092.79	solar retinopathy — see Re
specified NEC F31.89	pregnancy 092.20	specified type NEC H31.8
bladder N32.9	specified NEC 092.29	ciliary body — see Disorder, iri
functional NEC N31.9	puerperium 092.20	degeneration — see Deger
in schistosomiasis B65.0 [N33]	specified NEC 092.29	coagulation (factor) — see also
specified NEC N32.89	cracked nipple — see Cracked nipple	D68.9
bleeding D68.9	galactorrhea — see Galactorrhea	newborn, transient P61.6
blood D75.9	hypogalactia 092.4	cocaine use
in congenital early syphilis A5Ø.Ø9 [D77]	lactation disorder NEC 092.79	mild F14.10
body dysmorphic F45.22	mastitis — see Mastitis	with
bone M89.9	nipple infection — see Infection, nipple	amphetamine, cocair
continuity M84.9	retracted nipple — see Retraction, nipple	intoxication
specified type NEC M84.80	specified type NEC N64.89	with perceptual d
ankle M84.87- <b>▼</b>	Briguet's F45.0	with perceptual d
fibula M84.86- <b>☑</b>	bullous, in diseases classified elsewhere L14	cocaine intoxication
foot M84.87- <b>☑</b>	caffeine use	cocaine-induced
hand M84.84- <b>▼</b>	mild	anxiety disorder F
humerus M84.82- ▼	with	bipolar and relate
neck M84.88	caffeine-induced	depressive disorde
pelvis M84.859	anxiety disorder F15.180	obsessive-compul
radius M84.83- <b>▼</b>	sleep disorder F15.182	der F14.188
rib M84.88	moderate or severe	psychotic disorder
shoulder M84.81- <b>☑</b>	with	sexual dysfunction
skull M84.88	caffeine-induced	sleep disorder F14
thigh M84.85- ✓	anxiety disorder F15.280	in remission (early) (sust
tibia M84.86- ✓	sleep disorder F15.282	moderate or severe F14.20
ulna M84.83- <b>☑</b>	· · · · · · · · · · · · · · · · · · ·	

```
delirium F12.121
isturbances F12.122
al disturbances F12.129
12.180
r F12.159
2.188
tained) F12.11
..
listurbances F12.222
al disturbances F12.229
12.280
r F12.259
2.288
tained) F12.21
E74.39
74.9
gement, joint, articular
e Chondrocalcinosis
ement, joint, articular
ological condition) (with r) FØ6.1
physiological condition
93.25
F84.3
abolism E78.7Ø
ne E78.72
oroid
neration, choroid
ment, choroid
y, choroid
rhage, choroid
oroid
nal
tinopathy, solar
neration, ciliary body
o Defect, coagulation
ne, or other stimulant
isturbances F14.122
al disturbances F14.129
delirium F14.121
14.180
d disorder F14.14
er F14.14
Isive and related disor-
r F14.159
in F14.139
in F14.181
4.182
tained) F14.11
```

ICD-10-CM 2025 Chapter 2. Neoplasms Guidelines and Examples

#### Chapter 2. Neoplasms (CØØ-D49)

#### **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.

#### **General guidelines**

Chapter 2 of the ICD-10-CM contains the codes for most benign and all malignant neoplasms. Certain benign neoplasms, such as prostatic adenomas, may be found in the specific body system chapters. To properly code a neoplasm, it is necessary to determine from the record if the neoplasm is benign, in-situ, malignant, or of uncertain histologic behavior. If malignant, any secondary (metastatic) sites should also be determined.

Primary malignant neoplasms overlapping site boundaries

A primary malignant neoplasm that overlaps two or more contiguous (next to each other) sites should be classified to the subcategory/code .8 ('overlapping lesion'), unless the combination is specifically indexed elsewhere. For multiple neoplasms of the same site that are not contiguous such as tumors in different quadrants of the same breast, codes for each site should be assigned.

A 73-year-old white female with a large rapidly growing malignant tumor in the left breast extending from the upper outer quadrant into the axillary tail

## C5Ø.812 Malignant neoplasm of overlapping sites of left female

*Explanation*: Because this is a single large tumor that overlaps two contiguous sites, a single code for overlapping sites is assigned.

A 52-year old white female with two distinct lesions of the right breast, one (0.5 cm) in the upper outer quadrant and a second (1.5 cm) in the lower outer quadrant; path report indicates both lesions are malignant.

## C50.411 Malignant neoplasm of upper-outer quadrant of right

## C50.511 Malignant neoplasm of lower-outer quadrant of right

Explanation: This patient has two distinct malignant lesions of right breast in adjacent quadrants. Because the lesions are not contiguous, two codes are reported.

#### Malignant neoplasm of ectopic tissue

Malignant neoplasms of ectopic tissue are to be coded to the site of origin mentioned, e.g., ectopic pancreatic malignant neoplasms involving the stomach are coded to malignant neoplasm of pancreas, unspecified (C25.9).

The neoplasm table in the Alphabetic Index should be referenced first. However, if the histological term is documented, that term should be referenced first, rather than going immediately to the Neoplasm Table, in order to determine which column in the Neoplasm Table is appropriate. For example, if the documentation indicates "adenoma," refer to the term in the Alphabetic Index to review the entries under this term and the instructional note to "see also neoplasm, by site, benign." The table provides the proper code based on the type of neoplasm and the site. It is important to select the proper column in the table that corresponds to the type of neoplasm. The Tabular List should then be referenced to verify that the correct code has been selected from the table and that a more specific site code does not exist.

See Section I.C.21. Factors influencing health status and contact with health services, Status, for information regarding Z15.0, codes for genetic susceptibility to cancer.

#### a. Admission/Encounter for treatment of primary site

If the malignancy is chiefly responsible for occasioning the patient admission/encounter and treatment is directed at the primary site, designate the primary malignancy as the principal/first-listed diagnosis.

The only exception to this guideline is if the administration of chemotherapy, immunotherapy or external beam radiation therapy is chiefly responsible for occasioning the admission/encounter. In that case, assign the appropriate Z51.— code as the first-listed or principal diagnosis, and the underlying diagnosis or problem for which the service is being performed as a secondary diagnosis.

#### b. Admission/Encounter for treatment of secondary site

When a patient is admitted because of a primary neoplasm with metastasis and treatment is directed toward the secondary site only, the secondary neoplasm is designated as the principal diagnosis even though the primary malignancy is still present.

Patient with primary prostate cancer with metastasis to lungs admitted for wedge resection of mass in right lung

#### C78.Ø1 Secondary malignant neoplasm of right lung

#### C61 Malignant neoplasm of prostate

Explanation: Since the admission is for treatment of the lung metastasis, the secondary lung metastasis is sequenced before the primary prostate cancer.

#### c. Coding and sequencing of complications

Coding and sequencing of complications associated with the malignancies or with the therapy thereof are subject to the following guidelines:

#### 1) Anemia associated with malignancy

When admission/encounter is for management of an anemia associated with the malignancy, and the treatment is only for anemia, the appropriate code for the malignancy is sequenced as the principal or first-listed diagnosis followed by the appropriate code for the anemia (such as code D63.0, Anemia in neoplastic disease).

Patient is admitted for treatment of anemia in advanced colon cancer

#### C18.9 Malignant neoplasm of colon, unspecified

#### D63.0 Anemia in neoplastic disease

Explanation: Even though the admission was solely to treat the anemia, this guideline indicates that the code for the malignancy is sequenced

# 2) Anemia associated with chemotherapy, immunotherapy and radiation therapy

When the admission/encounter is for management of an anemia associated with an adverse effect of the administration of chemotherapy or immunotherapy and the only treatment is for the anemia, the anemia code is sequenced first followed by the appropriate codes for the neoplasm and the adverse effect (T45.1X5, Adverse effect of antineoplastic and immunosuppressive drugs).

A 56-year-old Hispanic male with grade II follicular lymphoma involving multiple lymph node sites referred for blood transfusion to treat anemia due to chemotherapy

#### D64.81 Anemia due to antineoplastic chemotherapy

C82.18 Follicular lymphoma grade II, lymph nodes of multiple sites

# T45.1X5A Adverse effect of antineoplastic and immunosuppressive drugs, initial encounter

Explanation: The code for the anemia is sequenced first followed by the code for the malignant neoplasm and lastly the code for the adverse

When the admission/encounter is for management of an anemia associated with an adverse effect of radiotherapy, the anemia code should be sequenced first, followed by the appropriate neoplasm code and code Y84.2, Radiological procedure and radiotherapy as the cause of abnormal reaction of the patient, or of later complication, without mention of misadventure at the time of the procedure.

ICD-10-CM 2025 475

CD-1	0-CM	2025		Chapter 3. Diseases	of the Blo	od and	d Bloo	d-forn	ning O	rgans		D82.8-D89.81
		D82.8		deficiency associated with other specified					D86.8		oid arthropathy	
		D82.9	defects Immuno unspeci	deficiency associated with major defect,	нсс					7 Sarc	lyarthritis in sarcoidosis  oid myositis  oidosis of other sites	
√4th	D83	Comm	-	ole immunodeficiency					D00.0		patic granuloma	
				n variable immunodeficiency with predom	inant			D06.0			eoparotid fever [Heerfordt]	
			abnorm	alities of B-cell numbers and function	CC HCC	(1)	D00				Inspecified	
		D83.1		n variable immunodeficiency with predom regulatory T-cell disorders	inant cc Hcc	√ 4 <sup>m</sup>	D89	classif		ers invo	lving the immune mechanisi	n, not eisewner
		D83.2		n variable immunodeficiency with autoant				EXCL	UDES 1	,, ,	obulinemia NOS (R77.1) onal gammopathy (of undetermi	ned significance)
		D83.8		mmon variable immunodeficiencies	CC HCC					(D4	7.2)	,
				n variable immunodeficiency, unspecified	CC HCC				JDES 2	,	nt failure and rejection (T86)	
√4 <sup>th</sup>	D84	Other	immuno	deficiencies				D89.0			pergammaglobulinemia rgammaglobulinemic purpura	
		D84.Ø	Lympho	cyte function antigen-1 [LFA-1] defect	HCC						ammopathy NOS	
		D84.1		in the complement system	HCC			D89.1		lobulin	<b>emia</b> nemic purpura	нсс
	√5 <sup>th</sup>	D84.8		erase inhibitor [C1-INH] deficiency recified immunodeficiencies							nemic vasculitis	
		200	•	020,4Q,10-12					Esse	ential cry	oglobulinemia	
			D84.81	Immunodeficiency due to conditions classis							yoglobulinemia lobulinemia	
				elsewhere	CC HCC						globulinemia	
				Code first underlying condition, such as: chromosomal abnormalities (Q90-Q99)							ryoglobulinemia	
				diabetes mellitus (EØ8-E13)							globulinemia, unspecified	
				malignant neoplasms (CØØ-C96)				D89.3			nstitution syndrome onstitution inflammatory syndrometry	нсо ome [IRIS]
				<b>EXCLUDES 1</b> certain disorders involving the inmechanism (D80-D83, D84							al code for adverse effect, if app	
				D84.9)	.0, 004.1,					_	36-T5Ø with fifth or sixth charac	
				human immunodeficiency virus	[HIV]		√5 <sup>th</sup>	D89.4			vation syndrome and related	
				disease (B2Ø) <b>AHA:</b> 2021,1Q,52			$\neg$		EXC		aggressive systemic mastocytosi congenital cutaneous mastocyto	
		√6 <sup>th</sup>	D84.82	Immunodeficiency due to drugs and extern	al causes						(non-congenital) cutaneous mas	
				D84.821 Immunodeficiency due to							(indolent) systemic mastocytosis	
				drugs	CC HCC						malignant mast cell neoplasm (C	
				Immunodeficiency due to (currer medication	it or past)						malignant mastocytoma (C96.29 mast cell leukemia (C94.3-)	9)
				Use additional code for adverse	effect if						mast cell sarcoma (C96.22)	
				applicable, to identify adve							mastocytoma NOS (D47.09)	
				of drug (T36-T50 with fifth character 5)	or six						other mast cell neoplasms of und (D47.09)	ertain behavior
				Use additional code, if applicable	e, for						systemic mastocytosis associated	d with a clonal
				associated long term (curre	nt) drug	`					hematologic non-mast cell	lineage disease
				therapy drug or medication long term (current) drug thera					ΔН	<b>\:</b> 2016,4	(SM-AHNMD) (D47.Ø2) ∩ 11	
				systemic steroids (Z79.52							t cell activation, unspecified	нес
				other long term (current) drug						Ma	st cell activation disorder, unsp	
				(Z79.899)					D00 /		st cell activation syndrome, NO	
				D84.822 Immunodeficiency due to exter	cc HCC						oclonal mast cell activation s pathic mast cell activation sy	,
				Code also, if applicable, radiolog							ndary mast cell activation	нсс
				procedure and radiotherap						Se	condary mast cell activation syr	
				Use additional code for external c as:	ause such				D00 4		de also underlying etiology, if k	
				exposure to ionizing radiation	(W88)				D09.4		editary alpha tryptasemia e additional code, if applicable,	for:
			D84.89	Other immunodeficiencies	CC HCC						allergy status, other than to dru	
		D84.9		deficiency, unspecified	CC HCC						substances (Z91.Ø-)	(707.000)
				ocompromised NOS odeficient NOS							personal history of anaphylaxis (IA: 2021,4Q,8	(287.892)
				osuppressed NOS					D89.4		er mast cell activation disorde	er HCC
			AHA: 2	020,4Q,10							her mast cell activation syndror	
√4 <sup>th</sup>	D86	Sarcoi DEF:		of immune cells resulting in granuloma formati	on. Often		√5 <sup>th</sup>		not e	sewher	ed disorders involving the imm e classified	nune mechanism
				s and lymphatic system but can occur in other b				√6 <sup>th</sup>	D89.8		t-versus-host disease	
				osis of lung	HCC						de first underlying cause, such a complications of blood transfus	
				osis of lymph nodes Osis of lung with sarcoidosis of lymph node	s HCC						complications of transplanted o	
				osis of rung with sarcoldosis of lymph hode							(T86)	
	$\sqrt{5}$ th			osis of other sites						Us	e additional code to identify ass	sociated
			D86.81	Sarcoid meningitis						,	manifestations, such as: desquamative dermatitis (L3Ø.8)	)
				Multiple cranial nerve palsies in sarcoidos	is HCC						diarrhea (R19.7)	,
				Sarcoid iridocyclitis Sarcoid pyelonephritis							elevated bilirubin (R17)	
			JUU.04	Tubulo-interstitial nephropathy in sarcoido	sis						nair loss (L65.9) .810   Acute graft-versus-host	
			D86.85	Sarcoid myocarditis						ניסט.	disease	CC UPD HCC
										D89.	.811 Chronic graft-versus-ho	
											disease	CC UPD HCC

Unspecified

CC

CC

CC HCC

# Chapter 9. Diseases of the Circulatory System (IØØ-I99)

certain conditions originating in the perinatal period (PØ4-P96) certain infectious and parasitic diseases (AØØ-B99)

complications of pregnancy, childbirth and the puerperium (OØØ-O9A) congenital malformations, deformations, and chromosomal abnormalities (000-099)

endocrine, nutritional and metabolic diseases (EØØ-E88) injury, poisoning and certain other consequences of external causes (SØØ-T88)

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.-)

This chapter contains the following blocks:

IØØ-IØ2 Acute rheumatic fever Chronic rheumatic heart diseases 105-109 11Ø-I1A Hypertensive diseases 120-125 Ischemic heart diseases

126-128 Pulmonary heart disease and diseases of pulmonary circulation

13Ø-15A Other forms of heart disease 160-169 Cerebrovascular diseases

170-179 Diseases of arteries, arterioles and capillaries

180-189 Diseases of veins, lymphatic vessels and lymph nodes, not elsewhere

195-199 Other and unspecified disorders of the circulatory system

#### Acute rheumatic fever (IØØ-IØ2)

**DEF:** Rheumatic fever: Inflammatory disease that can follow a throat infection by group A streptococci. Complications can involve the joints (arthritis), subcutaneous tissue (nodules), skin (erythema marginatum), heart (carditis), or brain (chorea).

#### IØØ Rheumatic fever without heart involvement

**INCLUDES** arthritis, rheumatic, acute or subacute

**EXCLUDES 1** rheumatic fever with heart involvement (IØ1.Ø-IØ1.9)

#### IØ1 Rheumatic fever with heart involvement

**EXCLUDES 1** chronic diseases of rheumatic origin (105-109) unless rheumatic fever is also present or there is evidence of reactivation or activity of the rheumatic process

#### **IØ1.0** Acute rheumatic pericarditis

Any condition in 100 with pericarditis

Rheumatic pericarditis (acute)

**EXCLUDES 1** acute pericarditis not specified as rheumatic (130.-)

#### **IØ1.1** Acute rheumatic endocarditis

Any condition in IØØ with endocarditis or valvulitis Acute rheumatic valvulitis

#### **IØ1.2** Acute rheumatic myocarditis

Any condition in 100 with myocarditis

#### IØ1.8 Other acute rheumatic heart disease

Any condition in 100 with other or multiple types of heart involvement

Acute rheumatic pancarditis

#### 101.9 Acute rheumatic heart disease, unspecified

ny condition in IØØ with unspecified type of heart involvement Rheumatic carditis, acute

Rheumatic heart disease, active or acute

#### ✓4 IØ2 Rheumatic chorea

INCLUDES Sydenham's chorea

chorea NOS (G25.5) EXCLUDES 1 Huntington's chorea (G1Ø)

#### IØ2.Ø Rheumatic chorea with heart involvement

Chorea NOS with heart involvement

Rheumatic chorea with heart involvement of any type classifiable under IØ1.-

**IØ2.9** Rheumatic chorea without heart involvement

Rheumatic chorea NOS

#### Chronic rheumatic heart diseases (105-109)

#### IØ5 Rheumatic mitral valve diseases

conditions classifiable to both IØ5.Ø and IØ5.2-IØ5.9. INCLUDES whether specified as rheumatic or not

EXCLUDES 1 mitral valve disease specified as nonrheumatic (134.-) mitral valve disease with aortic and/or tricuspid valve involvement (IØ8.-)

#### 105.0 Rheumatic mitral stenosis

Mitral (valve) obstruction (rheumatic)

#### IØ5.1 Rheumatic mitral insufficiency

Rheumatic mitral incompetence Rheumatic mitral regurgitation

**EXCLUDES 1** mitral insufficiency not specified as rheumatic (134.0)

#### **IØ5.2** Rheumatic mitral stenosis with insufficiency Rheumatic mitral stenosis with incompetence or regurgitation

105.8 Other rheumatic mitral valve diseases

#### Rheumatic mitral (valve) failure

Rheumatic mitral valve disease, unspecified 105.9 Rheumatic mitral (valve) disorder (chronic) NOS

#### IØ6 Rheumatic aortic valve diseases

EXCLUDES 1 aortic valve disease not specified as rheumatic (135.-) aortic valve disease with mitral and/or tricuspid valve involvement (IØ8.-)

#### 106.0 Rheumatic aortic stenosis

Rheumatic aortic (valve) obstruction

#### 106.1 Rheumatic aortic insufficiency

Rheumatic aortic incompetence Rheumatic aortic regurgitation

106.2 Rheumatic aortic stenosis with insufficiency

Rheumatic aortic stenosis with incompetence or regurgitation

#### 106.8 Other rheumatic aortic valve diseases

#### Rheumatic aortic valve disease, unspecified

Rheumatic aortic (valve) disease NOS

#### IØ7 Rheumatic tricuspid valve diseases

**EXCLUDES 1** 

INCLUDES rheumatic tricuspid valve diseases specified as rheumatic or unspecified

> tricuspid valve disease specified as nonrheumatic (136.-) tricuspid valve disease with aortic and/or mitral valve involvement (IØ8.-)

#### 107.0 Rheumatic tricuspid stenosis

Tricuspid (valve) stenosis (rheumatic)

#### Rheumatic tricuspid insufficiency

Tricuspid (valve) insufficiency (rheumatic)

#### 107.2 Rheumatic tricuspid stenosis and insufficiency

107.8 Other rheumatic tricuspid valve diseases

107.9 Rheumatic tricuspid valve disease, unspecified Rheumatic tricuspid valve disorder NOS

#### 108 Multiple valve diseases

**√**40

CC

CC

CC

multiple valve diseases specified as rheumatic or INCLUDES unspecified

endocarditis, valve unspecified (138) EXCLUDES 1

multiple valve disease specified a nonrheumatic (134.-, 135.-, . 136.-, 137.-, 138.-, Q22.-, Q23.-, Q24.8-) rheumatic valve disease NOS (109.1)

#### IØ8.Ø Rheumatic disorders of both mitral and aortic valves

Involvement of both mitral and aortic valves specified as rheumatic or unspecified AHA: 2019,2Q,5

- IØ8.1 Rheumatic disorders of both mitral and tricuspid valves
- IØ8.2 Rheumatic disorders of both aortic and tricuspid valves
- Combined rheumatic disorders of mitral, aortic and tricuspid valves
- **IØ8.8** Other rheumatic multiple valve diseases
- 108.9 Rheumatic multiple valve disease, unspecified

#### IØ9 Other rheumatic heart diseases

109.0 Rheumatic myocarditis

**EXCLUDES 1** myocarditis not specified as rheumatic (151.4)

#### Rheumatic diseases of endocardium, valve unspecified

Rheumatic endocarditis (chronic) Rheumatic valvulitis (chronic)

**EXCLUDES 1** endocarditis, valve unspecified (138)

#### 109.2 Chronic rheumatic pericarditis

Adherent pericardium, rheumatic Chronic rheumatic mediastinopericarditis Chronic rheumatic myopericarditis

**EXCLUDES 1** chronic pericarditis not specified as rheumatic (131.-)

#### 109.8 Other specified rheumatic heart diseases

109.81 Rheumatic heart failure

Use additional code to identify type of heart failure

#### Other specified rheumatic heart diseases Rheumatic disease of pulmonary valve

109.9 Rheumatic heart disease, unspecified

Rheumatic carditis

**EXCLUDES 1** rheumatoid carditis (MØ5.31)

✓ Additional Character Required ICD-10-CM 2025

√x7<sup>th</sup> Placeholder

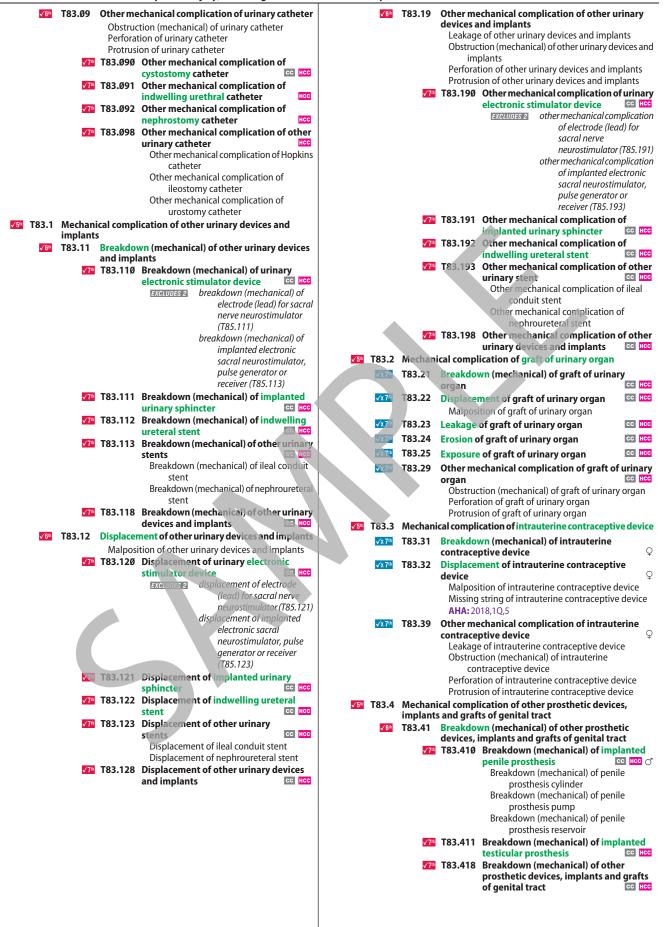
Questionable PDx

Manifestation

Unacceptable PDx H1-H14 HAC HCC CMS-HCC Dx Unspecified

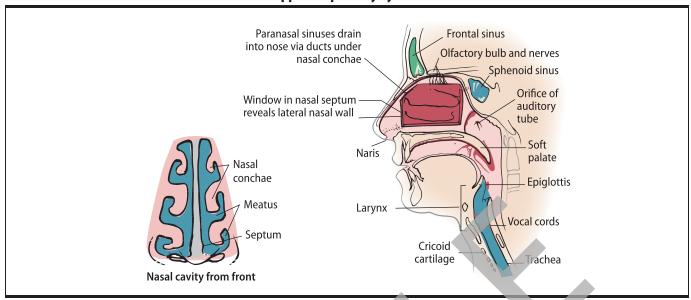
HIV HIV DX

D-10	-CM	2025			Chapter 15. Preg	nancy, Chi	ldbirth and t	he Pu	erperium		009.892-010.912
				009.892	Supervision of other high risk		√5 <sup>th</sup>	01ø.2			tensive chronic kidney disease
					pregnancies, second trimester	UPD M Q					nancy, childbirth and the puerperium 12 specified as a reason for obstetric care
				009.893	Supervision of other high risk				du	ring pregn	ancy, childbirth or the puerperium
				000 000	pregnancies, third trimester Supervision of other high risk	UPD M Q				ditional cod ronic kidne	le from I12 to identify the type of hypertensive
				009.099	pregnancies, unspecified			√6 <sup>th</sup>			ing hypertensive chronic kidney disease
	_				trimester	UPD M Q				complica	ting pregnancy
	√5 <sup>th</sup>	009.9	•	-	h risk pregnancy, unspecified	a aifi a d				010.211	Pre-existing hypertensive chronic kidney disease complicating pregnancy, first
			09.90		ion of high risk pregnancy, unsp ied trimester	UPD M Q					trimester <b>™</b> ♀
			009.91	Supervisi	ion of high risk pregnancy, unsp					010.212	Pre-existing hypertensive chronic kidney disease complicating pregnancy, second
			000 02	first trim	ester ion of high risk pregnancy, unsp						trimester M Q
			000.02	second tr		UPD M Q				010.213	Pre-existing hypertensive chronic kidney disease complicating pregnancy, third
			009.93	Supervisi	ion of high risk pregnancy, unsp	ecified,					trimester M Q
				uma um	iestei	0.5 M ¥				010.219	Pre-existing hypertensive chronic kidney disease complicating pregnancy,
E	der				pertensive disorders in pre	gnancy,					unspecified trimester
				n and th	e puerperium (01Ø-016)				010.22		ing hypertensive chronic kidney disease
		2016,40	,	nortoncio	on complicating pregnancy, chi	ldhisth			010.23		ting childbirth
V 4	שוט		e puerpe	rium						complica	ting the puerperium <b>™</b> ♀
		INCL	JDES pr		hypertension with pre-existing pro ating pregnancy, childbirth and th		√5 <sup>th</sup>	01Ø.3	Pre-exist	ting hyper	tensive heart and chronic kidney disease nancy, childbirth and the puerperium
				puerperi		e			Any co	ndition in I	13 specified as a reason for obstetric care
		EXCLU	DES 2 pre		ypertension with superimposed pre- ating pregnancy, childbirth and the p						ancy, childbirth or the puerperium le from 113 to identify the type of hypertensive
				(O11)	ating pregnancy, childoirth and the p	uerperium					ronic kidney disease
	$\sqrt{5}$ <sup>th</sup>	010.0			tial hypertension complicating p	regnancy,		√6 <sup>th</sup>	010.31		ng hypertensive heart and chronic kidney
					<b>puerperium</b> 10 specified as a reason for obstet	ric care					omplicating pregnancy Pre-existing hypertensive heart and
			dι	ıring pregr	nancy, childbirth or the puerperiun	n					chronic kidney disease complicating
		√6 <sup>th</sup>	010.01	Pre-exist pregnance	ing essential hypertension com	plicating				010.312	pregnancy, first trimester $\square \square$ Pre-existing hypertensive heart and
					Pre-existing essential hyperter	nsion				-	chronic kidney disease complicating
					complicating pregnancy, first trimester	сс М Ф				010.313	pregnancy, second trimester    □ ♀  Pre-existing hypertensive heart and
				010.012	Pre-existing essential hyperter	nsion				015.515	chronic kidney disease complicating
					complicating pregnancy, secon trimester	nd cc M ♀				O1Ø 319	pregnancy, third trimester    □ ♀  Pre-existing hypertensive heart and
				010.013	Pre-existing essential hyperter	nsion				010.517	chronic kidney disease complicating
					complicating pregnancy, third trimester	сс М Q			O1Ø 32	Dro-ovicti	pregnancy, unspecified trimester $\square \bigcirc$ ing hypertensive heart and chronic kidney
				010.019	Pre-existing essential hyperter	ision			010.52		omplicating childbirth
					complicating pregnancy, unsp- trimester	ecified M O			010.33		ing hypertensive heart and chronic kidney omplicating the puerperium
			010.02	Pre-exist	ing essential hypertension com		√5th	01ø.4	Pre-exist		dary hypertension complicating
			010 02	childbirt		cc MQ			pregnan	cy, childb	irth and the puerperium
			C 10.03	the puer	ing essential hypertension com perium	piicating <u>M</u> ♀			du	ring pregn	15 specified as a reason for obstetric care ancy, childbirth or the puerperium
	$\sqrt{5}^{\text{th}}$	010.1			rtensive heart disease complica	ting					de from I15 to identify the type of secondary
					irth and the puerperium  111 specified as a reason for obstet	ric care		√6th	,	pertension  Pre-exist	ing secondary hypertension complicating
					nancy, childbirth or the puerperium					pregnand	y
				ditional cod eart disease	de from I11 to identify the type of hy	pertensive				010.411	Pre-existing secondary hypertension complicating pregnancy, first
		√6 <sup>th</sup>	_	Pre-exist	ing hypertensive heart disease						trimester cc M ♀
					ting pregnancy Pre-existing hypertensive hear	t disassa				010.412	Pre-existing secondary hypertension complicating pregnancy, second
				0.5	complicating pregnancy, first						trimester cc M ♀
				010 112	trimester Pre-existing hypertensive hear	Q M azsazih tr				010.413	Pre-existing secondary hypertension complicating pregnancy, third
				0.0	complicating pregnancy, secon	nd					trimester cc M ♀
				O1Ø 113	trimester Pre-existing hypertensive hear	Q M azsazih tr				010.419	Pre-existing secondary hypertension complicating pregnancy, unspecified
				0.55	complicating pregnancy, third						trimester <b>™</b> ♀
				010.119	trimester Pre-existing hypertensive hear	™ ♀ T disease			010.42	Pre-exist	ing secondary hypertension complicating
					complicating pregnancy, unsp	ecified			010.43	Pre-exist	ing secondary hypertension complicating
			O1Ø 12	Pro-ovice	trimester ing hypertensive heart disease	MQ	(Fib.	014 0	Hnenes!	the puer	
				complica	ting childbirth	MQ	√ 0**	פ.שו ט			xisting hypertension complicating irth and the puerperium
			010.13		ing hypertensive heart disease ting the puerperium	MΩ		$\sqrt{6}$ <sup>th</sup>	010.91		ed pre-existing hypertension complicating
				complica	any are puerperium	ш Ұ				pregnand 010.911	Unspecified pre-existing hypertension
											complicating pregnancy, first trimester □ □ □ □ □
										010.912	Unspecified pre-existing hypertension
											complicating pregnancy, second trimester  □□ M ♀
							İ.				unicatei 🔤 🖳 ¥

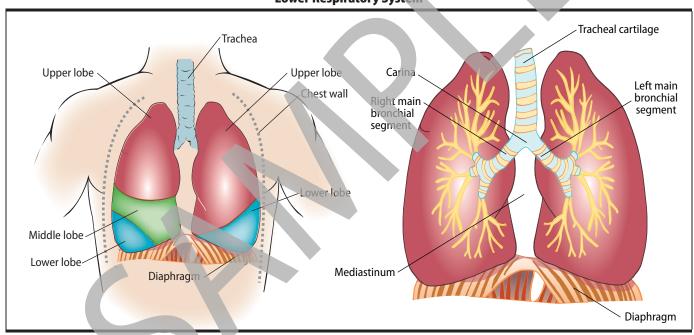


ICD-10-CM 2025 Illustrations

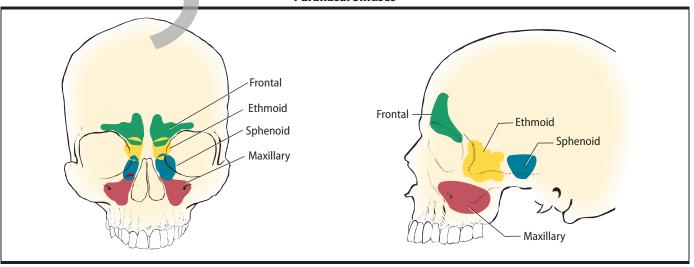
#### **Upper Respiratory System**



#### **Lower Respiratory System**



#### **Paranasal Sinuses**



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