ICD-10- Clinical Modification (CM) Codes Effective October 1, 2020 to September 20, 2021

490 code additions, 47 revisions, 58 deletions

	REVISED CODES		
From	То	Tabular Description	
A84.8	A84.41	Disease, diseased	
		Viral, Powassan	
A84.8	A84.81	Encephalitis	
		Powassan	
A84.8	A84.89	Encephalitis	
	1	Louping ill	
		Viral, arthropod	
		Louping ill (encephalitis)	
B60.0	B60.00	Babesiosis	
		Disease	
		Protozoal, babesiosis	
		Piroplasmosis (see also Babesiosis)	
D59.1	D59.10	Disease, diseased	
		Autoimmune	
**************************************		Hemolytic (cold type) (warm type)	
D59.1	D59.11	Anemia	
1		Warm-type (primary) (secondary) (symptomatic)	
		Disease, diseased	
		Friedreich's, combined systemic or ataxia	
059.1	D59.12	Anemia	
		Cold-type (primary) (secondary) (symptomatic)	
		Cold, agglutinin disease or hemoglobinuria (chronic)	
		Cold, sensitivity, autoimmune	
		Disease, diseased	
		Agglutinin or hemoglobinuria	
		Hemagglutinin	
		Sensitive, sensitivity	
		Cold, autoimmune	
59.1	D59.13	Anemia	
		Mixed-type	

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D59.1	D59.19	Anemia
		Acute
		Autoimmune
		Dyke-Young type (secondary)
		Hemolytic
		Autoimmune NEC
		Acute
		Lederer's
		Hemolytic
		Lederer's
		Dyke-Young anemia (secondary) (symptomatic)
		Lederer's anemia
D72.1	D72.10	Acidocytosis
		Eosinophilia (allergic) (idiopathic) (secondary)
D72.1	D72.119	Syndrome
		Hypereosinophilic (HES)
D72.1	D72.19	Leukocytosis
		Eosinophilic
D84.8	D84.89	Deficiency
		Immunity, cell-mediated
		Immunodeficiency
		Specified type NEC
E41	E43	Emaciation (due to malnutrition)
E70.8	E70.89	Aromatic amino-acid metabolism
		Specified NEC
		Disorder (of)
		Aromatic amino-acid metabolism, specified NEC
		Metabolism, amino-acid, aromatic, other specified
		Hydroxykynureninuria
E74.8	E74.818	Deficiency, deficient
		Phosphomannomutase
		Phosphomannose isomerase
		Phosphomannosyl mutase
E74.8	E74.818	Glycosuria
		Renal (familial)
E74.8	E74.89	Disorder (of)
		Carbohydrate
		Metabolism, specified NEC
		Congenital glycosylation (CDG)
		Pentosuria (essential)

	REVISED CODES			
F20.5	F20.9	Schizophrenia, schizophrenic		
		Chronic undifferentiated		
G11.1	G11.10	Ataxia		
		Early-onset		
G11.1	G11.11	Ataxia		
		Friedreich's (heredofamilial) (cerebellar) (spinal) (with retained reflexes)		
		Spinal (Friedreich's)		
		Hereditary (Friedreich's)		
		Cardiomyopathy (familial) (idiopathic)		
		Friedreich's		
		Friedreich's		
		Ataxia		
		Combined systemic disease		
		Sclerosis		
		Sclerosis, sclerotic		
		Friedreich's (spinal cord)		
		Hereditary		
		Spinal (Friedreich's ataxia)		
		Spinal (cord) (progressive)		
		Hereditary (Friedreich's ataxia)		
G11.1	G11.19	Ataxia		
		Hunt's		
		Spinocerebellar, X-linked recessive		
		Dyssynergia		
		Cerebellaris myoclonica (Hunt's ataxia)		
		Hunt's		
		Dyssynergia cerebellaris myoclonica		
		Ramsay-Hunt disease or syndrome (see also Hunt's disease)		
		Meaning dyssynergia cerebellaris myoclonica		
G71.2	G71.20	Disproportion		
		Fiber-type		
		Myopathy		
		Benign congenital		
		Congenital		
G71.2	G71.21	Disease, diseased		
	•	Lung, nemaline body		
		Myopathy		
		Nemaline		
		Rod (body)		

		REVISED CODES
G71.2	G71.220	Myopathy
		Myotubular (centronuclear)
G71.2	G71.228	Myopathy, centronuclear
G71.2	G71.29	Disease, diseased
		Central core
		Lung
		Minicore
		Multicore
		Multiminicore
		Myopathy
		Central core
G96.0	G96.00	Leak, leakage
		Cerebrospinal fluid
G96.0	G96.01	Otorrhea
		Cerebrospinal (fluid)
		Rhinorrhea
		Cerebrospinal (fluid)
G96.0	G96.08	Fistula (cutaneous), cerebrospinal (fluid)
G96.19	G96.198	Calcification
		Meninges (cerebral) (spinal)
		Cyst (colloid) (mucous) (simple) (retention)
		Spinal
		Spinal meninges
		Subdural
		Spinal cord
		Deformity
		Choroid plexus, acquired
		Meninges, acquired
		Spinal cord, acquired
		Fibrosis, fibrotic
		Meninges
		Leptomeningopathy
		Meningitis(basal) (basic) (brain) (cerebral) (cervical) (congestive) (diffuse) (hemorrhagic) (infantile)(membranous) (metastatic) (nonspecific) (pontine) (progressive) (simple) (spinal) (subacute)(sympathetic) (toxic)
		Ossificans
		Meningocele (spinal)
		Acquired
		Ossification
		Falx cerebri
		Meninges (cerebral) (spinal)

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G96.8	G96.89	Cyst (colloid) (mucous) (simple) (retention) Nervous system NEC
		Deficiency, deficient
		Central nervous system
		Disease, diseased
		Nervous system, specified NEC
		Disorder (of)
		Nervous system, central, specified NEC
		Trophoneurosis NEC
149.9	149.8	Arrhythmia (auricle) (cardiac) (juvenile) (nodal) (reflex) (supraventricular) (transitory) (ventricle)
J43.9	J44.9	Disease, diseased
		Lung, obstructive, chronic
J68.0	J69.0	Bronchitis, (diffuse) (fibrinous) (hypostatic) (infective) (membranous)
		Aspiration (due to food and vomit)
J70.5	T59.81-	Inhalation
		Smoke
J82	J82.81	Pneumonia
		Eosinophilic
	90	Pneumonitis (acute) (primary)
		Eosinophilic
		Acute
		Chronic
J82	J82.83	Asthma, asthmatic (bronchial) (catarrh) (spasmodic)
		Eosinophilic
		Pulmonary eosinophilic

		REVISED CODES
J82	J82.89	Allergy, allergic (reaction)
		Pneumonia
		Eosinophilia (allergic) (idiopathic) (secondary)
		Löffler's
		Pulmonary
		Tropical (pulmonary)
		Löffler's
		Eosinophilia
		Pneumonia
		Syndrome
		Pneumonia(acute) (double) (migratory) (purulent) (septic) (unresolved)
		Allergic (see also Pneumonitis, hypersensitivity)
		Broncho-, bronchial
		Allergic
		Syndrome
		Löffler's
		PIE (pulmonary infiltration with eosinophilia)
		Weingarten's (tropical eosinophilia)
		Weingarten's syndrome
J84.17	J84.718	Pneumonia
		Interstitial
		Due to, collagen vascular disease
		Known underlying cause
		In disease classified elsewhere
		Lymphocytic (due to collagen vascular disease) (in diseases classified elsewhere)
		Usual
		Due to collagen
		In diseases classified elsewhere
K20.8	K20.80	Abscess (connective tissue) (embolic) (fistulous) (infective) (metastatic) (multiple) (pernicious)(pyogenic) (septic)
		Esophagus
		Esophagitis (acute) (alkaline) (chemical) (chronic) (infectional) (necrotic) (peptic) (postoperative) (without bleeding)
		Specified NEC
K20.9	K20.90	Esophagitis (acute) (alkaline) (chemical) (chronic) (infectional) (necrotic) (peptic) (postoperative) (without bleeding)

		REVISED CODES
K21.0	K21.00	Allergy, allergic (reaction)
		Gastroesophageal reflux with esophagitis (without bleeding)
	Ц	Gastroesophageal reflux
		With esophagitis (w/o bleeding)
		Esophagitis (acute) (alkaline) (chemical) (chronic) (infectional) (necrotic) (peptic (postoperative)(without bleeding)
		Due to gastrointestinal reflux disease
		Reflux
		Reflux
		Esophageal
		With esophagitis (without bleeding)
		Gastroesophageal
		With esophagitis (without bleeding)
		Ulcer, ulcerated, ulcerating, ulceration, ulcerative
		Esophagus
		Due to
		Gastroesophageal reflux disease

		REVISED CODES
K59.8	K59.89	Achalasia (cardia) (esophagus)
		Sphincteral NEC
		Atonia (atony, atonic) of
		Cecum
		Intestine
		Contraction (contracture)
		Anus
		Rectum, rectal sphincter
		Dilatation
		Anus
		Duodenum
		lleum
		Jejunum
		Disease, diseased
		Intestine, functional, specified NEC
		Dyskinesia
		Intestinal
		Dyspepsia
		Intestinal
		Hyperemia (acute) (passive)
		Enteric
		Intestine
		Hypofunction
		Intestinal
		Hypomotility
		Intestine
		Ileus (bowel) (colon) (inhibitory) (intestine)
		Myxedema
		Immobile, immobility
		Intestine
		Irritable, irritability
		Duodenum
		lleum
		Jejunum
		Rectum
		Paralysis (paralytic)
<74.0	K74.00	Fibrosis, fibrotic
		Liver

		REVISED CODES
M77.9	M77.8	Enthesopathy (peripheral)
DO PODOSON STORANDON	50 FEBRUAR - 1000 FEB	Foot
		Forearm
		Hand
		Multiple sites
		Pelvis
		Shoulder
		Upper arm
M92.5	M92.50-	Osteochondrosis
		Fibula
		Tibia
N18.3	N18.30	Disease, diseased
		Kidney, chronic, stage 3 (moderate)
N92.3	A77.9	Spotted fever
O99.89	099.891	Pregnancy (single) (uterine)
		Complicated by
		Connective system disorders
		Diseases of
		Specified NEC
		Disorders of
		Ear and mastoid process
		Eye and adnexa
		Specified NEC
		Musculoskeletal condition
		Pelvic inflammatory disease
		Peritoneal (pelvic) adhesions
099.89	099.892	Delivery (childbirth) (labor)
		Complicated by condition NEC
O99.89	099.893	Prediabetes, prediabetic
		Complicating
		Puerperium
		Puerperal, puerperium (complicated by, complications)
		Disease
		Nonobstetric NEC
		Disorder
		Nonobstetric NEC
		Specified condition
251.20	Q51.28	Septum, septate (congenital)
		Uterus

		REVISED CODES
R51	R51.9	Headache
		Chronic daily
		Daily chronic
		Nasal septum
		Pain(s) (see also Painful)
		Face, facial
R74.0	R74.01	Elevated, elevation
		Liver function, aminotransferase
		SGOT
		SGPT
		Transaminase level
		Findings, abnormal, inconclusive, without diagnosis
		SGOT
		SGPT
		Transaminase
		Transaminasemia
R74.0	R74.02	Elevated, elevation
		Lactic acid dehydrogenase (LDH)
		Liver function lactate dehydrogenase
		Findings, abnormal, inconclusive, without diagnosis
***************************************		Lactic acid dehydrogenase (LDH)
R93.09	R93.0	Abnormal, abnormality, abnormalities, skull
T81.49	T81.43	Abscess (connective tissue) (embolic) (fistulous) (infective) (metastatic) (multiple) (pernicious)(pyogenic) (septic)
		Postprocedural

	ADDED CODES
Add	Tabular Description
CHAPTER 1. CERTA	AIN INFECTIOUS AND PARASITIC DISEASES (A00-B99)
A84.81	Powassan virus disease
A84.89	Other tick-borne viral encephalitis
	Louping ill
B60.00-B60.03	Babesiosis
B60.09	
	SES OF THE BLOOD AND BLOOD-FORMING ORGANS AND CERTAIN DISORDERS IMUNE MECHANISM (D50-D89)
D57.00	Sickle-cell disorders
D57.03	Hb-SS disease with crisis
D57.09	
D57.213	Sickle-cell/Hb-C disease
D57.218	
D57.219	

	ADDED CODES
CHAPTER 3 (CONT	TINUED)
D57.413 D57.419 D47.42 D47.431-D47.433 D47.438-D47.439 D57.44-D57.45 D57.451-D57.453 D57.458-D57.459	Sickle-cell thalassemia
D57.813 D57.818-D57.819	Other sickle-cell disorders
D59.11-D59.13 D59.19	Acquired hemolytic anemia
D84.81 D84.821-D84.822	Other immunodeficiencies Immunodeficiency due to
D89.831-D89.835 D89.839	Cytokine release syndrome Grade -
CHAPTER 4. ENDO	CRINE, NUTRITIONAL AND METABOLIC DISEASES (E00-E89)
E70.81 E70.89	Other disorders of aromatic amino-acid metabolism
E74.810 E74.818-E74.819	Disorders of glucose transport
E74.89	Other specified disorders of carbohydrate metabolism
CHAPTER 5. MENTA	AL, BEHAVIORAL AND NEURODEVELOPMENTAL DISORDERS (F01-F99)
F10.130-F10.132 F10.939	Alcohol abuse with withdrawal
F10.930-F10.932 F10.939	Alcohol use, unspecified with withdrawal
F11.13	Opioid abuse with withdrawal
F12.13	Cannabis abuse with withdrawal
F13.130-F13.132 F13.139	Sedative, hypnotic or anxiolytic-related abuse
F14.13	Cocaine abuse, unspecified with withdrawal
F14.93	Cocaine use, unspecified with withdrawal
F15.13	Other stimulant abuse with withdrawal
F19.130-F19.132 F19.139	Other psychoactive substance abuse with withdrawal
CHAPTER 6. DISEAS	SES OF THE NERVOUS SYSTEM (G00-G99)
G11.10-G11.11 G11.19	Early-onset cerebellar ataxia

	ADDED CODES		
CHAPTER 6 (CONTINUED)			
G40.42	Cyclin-Dependent Kinase-Like 5 Deficiency Disorder CDKL5		
G40.833-G40.834	Dravet syndrome		
G71.20-G71.21 G71.220 G71.228 G71.29	Congenital myopathies		
G96.00-G96.02 G96.08-G96.09	Cerebrospinal fluid leak		
G96.191	Perineural cyst		
G96.198	Other disorders of meninges, not elsewhere classified		
G96.810-G96.811 G96.819 G96.89 G97.83-G97.84	Intracranial hypotension		
CHAPTER 7. DISEAS	SES OF THE EYE AND ADNEXA (H00-H59)		
H18.501-H18.503 H18.509	Unspecified hereditary corneal dystrophies		
H18.511-H18.513 H18.519	Endothelial corneal dystrophy		
H18.521-H18.523 H18.529	Epithelial (juvenile) corneal dystrophy		
H18.531-H18.533 H18.539	Granular corneal dystrophy		
H18.541-H18.543 H18.549	Lattice corneal dystrophy		
H18.551-H18.553 H18.559	Macular corneal dystrophy		
H18.591-H18.593 H18.599	Other hereditary corneal dystrophies		
H55.82	Deficient smooth pursuit eye movements		
CHAPTER 10. DISEA	SES OF THE RESPIRATORY SYSTEM (J00-J99)		
J82.81-J82.83 J82.89	Pulmonary eosinophilia		
J84.710 J84.178	Other interstitial pulmonary disease with fibrosis		
CHAPTER 11. DISEA	SES OF THE DIGESTIVE SYSTEM (K00-K95)		
K20.80-K20.81	Other esophagitis		
<20.90-K20.91	Esophagitis, unspecified		
K21.00-K21.01	Gastro-esophageal reflux disease		

ADDED CODES		
CHAPTER 11 (CON	TINUED)	
K59.81	Other specified functional intestinal disorders	
K59.89		
K74.00-K74.03	Hepatic fibrosis	
CHAPTER 13. DISE	ASES OF THE MUSCULOSKELETAL SYSTEM AND CONNECTIVE TISSUE (M00-M99)	
M05.7A	Rheumatoid arthritis with rheumatoid factor	
M05.8A		
M06.8A	Other rheumatoid arthritis	
M08.0A	Juvenile arthritis	
M08.2A		
M08.4A		
M08.9A		
M18.09	Other and unspecified osteoarthritis	
M19.19-M19.29		
M24.19-M24.69	Other specific joint derangements	
M24.89		
M25.39	Other joint disorder, not elsewhere classified	
M25.59-M25.69		
M26.641-M26.643	Arthritis of temporomandibular joint	
M26.649		
M26.651-M26.653	Arthropathy of temporomandibular joint	
M26.659		
M80.0A	Osteoporosis with current pathological fracture	
M80.8A		
M92.501-M92.503	Juvenile osteochondrosis	
M92.509		
M92.511-M92.513		
M92.519		
M92.521-M92.523		
M92.529		
M92.591-M92.593 M92.599		
	ASES OF THE GENITOURINARY SYSTEM (N00-N99)	
N00.A-N07.A	Glomerular diseases	
N18.30-N18.32	Chronic kidney disease, stage 3	
N61.20-N61.23	Granulomatous mastitis	
	NANCY, CHILDBIRTH AND THE PUERPERIUM (000-09A)	
034.218	Maternal care due to uterine scar from previous surgery	
034.22		
O99.891-O99.893	Other specified diseases and conditions complicating pregnancy, childbirth and the puerperium	

	ADDED CODES
CHAPTER 16. CER	TAIN CONDITIONS ORIGINATING IN THE PERINATAL PERIOD (P00-P96)
P91.821-P91.823	Neonatal cerebral infarction
P91.829	
CHAPTER 18. SYM ELSEWHERE CLAS	PTOMS, SIGNS AND ABNORMAL CLINICAL AND LABORATORY FINDINGS, NOT SIFIED (R00-R99)
R51.0	Headache
R51.9	
R74.01-R74.02	Nonspecific elevation of levels of transaminase and lactic acid dehydrogenase (LDH)
CHAPTER 19. INJU (S00-T88)	RY, POISONING AND CERTAIN OTHER CONSEQUENCES OF EXTERNAL CAUSES
S20.213-S20.374	Contusion of thorax
T40.411-T40.416	Poisoning by, adverse effect of and underdosing of narcotics and psychodyslep
T40.421-T40.426	tics [hallucinogens]
T40.491-T40.496	
T86.8401-T86.8429	Complications of transplanted organs
T86.8481-T86.8499	Corneal
CHAPTER 20. EXTE	RNAL CAUSES OF MORBIDITY (V00-Y99)
V00.03- thru V06.93	Pedestrian injured in transport accident
Y77.11	Ophthalmic devices associated with adverse incidents
Y77.19	
CHAPTER 21. FACT (Z00-Z99)	ORS INFLUENCING HEALTH STATUS AND CONTACT WITH HEALTH SERVICES
Z03.821-Z03.823	Encounter for medical observation for suspected diseases and conditions ruled out
CHAPTER 22. CODE	S FOR SPECIAL PURPOSES (U00-U85)
U07.0	Vaping-related disorder
	Dabbing related lung damage
	Dabbing related lung injury
	E-cigarette, or vaping, product use associated lung injury [EVALI]
	Electronic cigarette related lung damage
	Electronic cigarette related lung injury
	Use Additional code to identify manifestations, such as:
	abdominal pain (R10.84)
	acute respiratory distress syndrome (J80)
	diarrhea (R19.7)
	drug-induced interstitial lung disorder (J70.4)
	lipoid pneumonia (J69.1)
	weight loss (R63.4)

CHAPTER 22. CODES FOR SPECIAL PURPOSES (U00-U85)		
U07.1	COVID-19	
	Use Additional code to identify pneumonia or other manifestations	
	Excludes:	
	Coronavirus infection, unspecified (B34.2)	
	Coronavirus as the cause of diseases classified elsewhere (B97.2-)	
	Pneumonia due to SARS-associated coronavirus (J12.81)	
	DELETED CODES	
Deleted	Tabular Description	
A84.8	Code replaced with more specificity	
B60.0	Code replaced with more specificity	
D59.1	Code replaced with more specificity	
D72.1	Code replaced with more specificity	
E74.8	Code replaced with more specificity	
G11.1	Code replaced with more specificity	
G71.2	Code replaced with more specificity	
J82	Code replaced with more specificity	
Q51.20	Other doubling of uterus, unspecified	
T40.4X1-T40.4X6	Poisoning by, adverse effect of and underdosing of other synthetic narcotics	