ICD-10-CM 2021 Physician Update

In October 2020, the Centers for Medicare and Medicaid Services (CMS) issued its final list of hierarchical condition category (HGC) ICD-10-CM mappings for CY2021. As these final mappings were not available at the time of printing your ICD-10-CM physician code book, for your convenience, the following list identifies those ICD-10-CM codes that have been added as an HCC in the final mappings. Although changes to the ICD-10-CM classification became effective October 1, 2020, these HCC designation changes are not effective until January 1, 2021.

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Code	Description
D57.Ø3	Hb-SS disease with cerebral vascular involvement
D57.Ø9	Hb-SS disease with crisis with other specified complication
D57.213	Sickle-cell/Hb-C disease with cerebral vascular involvement
D57.218	Sickle-cell/Hb-C disease with crisis with other specified complication
D57.413	Sickle-cell thalassemia, unspecified, with cerebral vascular involvement
D57.418	Sickle-cell thalassemia, unspecified, with crisis with other specified complication
D57.42	Sickle-cell thalassemia beta zero without crisis
D57.431	Sickle-cell thalassemia beta zero with acute chest syndrome
D57.432	Sickle-cell thalassemia beta zero with splenic sequestration
D57.433	Sickle-cell thalassemia beta zero with cerebral vascular involvement
D57.438	Sickle-cell thalassemia beta zero with crisis with other specified complication
D57.439	Sickle-cell thalassemia beta zero with crisis, unspecified
D57.44	Sickle-cell thalassemia beta plus without crisis
D57.451	Sickle-cell thalassemia beta plus with acute chest syndrome
D57.452	Sickle-cell thalassemia beta plus with splenic sequestration
D57.453	Sickle-cell thalassemia beta plus with cerebral vascular involvement
D57.458	Sickle-cell thalassemia beta plus with crisis with other specified complication
D57.459	Sickle-cell thalassemia beta plus with crisis, unspecified
D57.813	Other sickle-cell disorders with cerebral vascular involvement
D57.818	Other sickle-cell disorders with crisis with other specified complication
D59.10	Autoimmune hemolytic anemia, unspecified
D59.11	Warm autoimmune hemolytic anemia
D59.12	Cold autoimmune hemolytic anemia
D59.13	Mixed type autoimmune hemolytic anemia
D59.19	Other autoimmune hemolytic anemia
D84.81	Immunodeficiency due to conditions classified elsewhere
D84.821	Immunodeficiency due to drugs
D84.822	Immunodeficiency due to external causes
D84.89	Other immunodeficiencies

E76.81 Aromatic L-amino acid decarboxylase deficiency E76.89 Other disorders of aromatic amino-acid metabolism E74.810 Glucose transporter protein type 1 deficiency E74.818 Other disorders of glucose transport E74.819 Disorders of glucose transport, unspecified E74.89 Other specified disorders of carbohydrate metabolism F10.130 Alcohol abuse with withdrawal, uncomplicated F10.131 Alcohol abuse with withdrawal delirium F10.132 Alcohol abuse with withdrawal with perceptual disturbance F10.933 Alcohol use, unspecified with withdrawal, unspecified F10.934 Alcohol use, unspecified with withdrawal delirium F10.935 Alcohol use, unspecified with withdrawal with perceptual disturbance F10.939 Alcohol use, unspecified with withdrawal, unspecified F11.13 Opioid abuse with withdrawal F12.13 Cannabis abuse with withdrawal F13.130 Sedative, hypnotic or anxiolytic abuse with withdrawal delirium F13.131 Sedative, hypnotic or anxiolytic abuse with withdrawal with perceptual disturbance F13.132 Sedative, hypnotic or anxiolytic abuse with withdrawal with perceptual disturbance	Code	Description
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uncomplicated F19.131 Other psychoactive substance abuse with withdrawal delirium F19.132 Other psychoactive substance abuse with withdrawal with perceptual disturbance F19.139 Other psychoactive substance abuse with withdrawal, unspecified G11.10 Early-onset cerebellar ataxia, unspecified G11.11 Friedreich ataxia G11.19 Other early-onset cerebellar ataxia G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.228 Other centronuclear myopathy	F15.13	Other stimulant abuse with withdrawal
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with perceptual disturbance F19.139 Other psychoactive substance abuse with withdrawal, unspecified G11.10 Early-onset cerebellar ataxia, unspecified G11.11 Friedreich ataxia G11.19 Other early-onset cerebellar ataxia G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	F19.131	
unspecified G11.10 Early-onset cerebellar ataxia, unspecified G11.11 Friedreich ataxia G11.19 Other early-onset cerebellar ataxia G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	F19.132	Other psychoactive substance abuse with withdrawal with perceptual disturbance
G11.11 Friedreich ataxia G11.19 Other early-onset cerebellar ataxia G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	F19.139	
G11.19 Other early-onset cerebellar ataxia G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	G11.1Ø	Early-onset cerebellar ataxia, unspecified
G40.42 Cyclin-Dependent Kinase-Like 5 Deficiency Disorder G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	G11.11	Friedreich ataxia
G40.833 Dravet syndrome, intractable, with status epilepticus G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	G11.19	Other early-onset cerebellar ataxia
G40.834 Dravet syndrome, intractable, without status epilepticus G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	G4Ø.42	Cyclin-Dependent Kinase-Like 5 Deficiency Disorder
G71.20 Congenital myopathy, unspecifed G71.21 Nemaline myopathy G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy	G4Ø.833	
G71.21 Nemaline myopathy G71.22Ø X-linked myotubular myopathy G71.228 Other centronuclear myopathy		· · ·
G71.220 X-linked myotubular myopathy G71.228 Other centronuclear myopathy		
G71.228 Other centronuclear myopathy		
	G71.22Ø	
G71.29 Other congenital myopathy		
	G71.29	Other congenital myopathy

Physician Update ICD-10-CM 2021

Code	Description
J82.81	Chronic eosinophilic pneumonia
J82.89	Other pulmonary eosinophilia, not elsewhere classified
J84.17Ø	Interstitial lung disease with progressive fibrotic phenotype in diseases classified elsewhere
J84.178	Other interstitial pulmonary diseases with fibrosis in diseases classified elsewhere
MØ5.7A	Rheumatoid arthritis with rheumatoid factor of other specified site without organ or systems involvement
MØ5.8A	Other rheumatoid arthritis with rheumatoid factor of other specified site
MØ6.ØA	Rheumatoid arthritis without rheumatoid factor, other specified site
MØ6.8A	Other specified rheumatoid arthritis, other specified site
MØ8.ØA	Unspecified juvenile rheumatoid arthritis, other specified site
MØ8.2A	Juvenile rheumatoid arthritis with systemic onset, other specified site
MØ8.4A	Pauciarticular juvenile rheumatoid arthritis, other specified site
MØ8.9A	Juvenile arthritis, unspecified, other specified site
N18.3Ø	Chronic kidney disease, stage 3 unspecified
N18.31	Chronic kidney disease, stage 3a
N18.32	Chronic kidney disease, stage 3b
P91.821	Neonatal cerebral infarction, right side of brain
P91.822	Neonatal cerebral infarction, left side of brain
P91.823	Neonatal cerebral infarction, bilateral
P91.829	Neonatal cerebral infarction, unspecified side
T4Ø.411A	Poisoning by fentanyl or fentanyl analogs, accidental (unintentional), initial encounter

Code	Description
T4Ø.412A	Poisoning by fentanyl or fentanyl analogs, intentional self-harm, initial encounter
T4Ø.412S	Poisoning by fentanyl or fentanyl analogs, intentional self-harm, sequela
T4Ø.414A	Poisoning by fentanyl or fentanyl analogs, undetermined, initial encounter
T4Ø.421A	Poisoning by tramadol, accidental (unintentional), initial encounter
T4Ø.422A	Poisoning by tramadol, intentional self-harm, initial encounter
T4Ø.422S	Poisoning by tramadol, intentional self-harm, sequela
T4Ø.424A	Poisoning by tramadol, undetermined, initial encounter
T4Ø.491A	Poisoning by other synthetic narcotics, accidental (unintentional), initial encounter
T4Ø.492A	Poisoning by other synthetic narcotics, intentional self-harm, initial encounter
T4Ø.492S	Poisoning by other synthetic narcotics, intentional self-harm, sequela
T4Ø.494A	Poisoning by other synthetic narcotics, undetermined, initial encounter
T86.8421	Corneal transplant infection, right eye
T86.8422	Corneal transplant infection, left eye
T86.8423	Corneal transplant infection, bilateral
T86.8429	Corneal transplant infection, unspecified eye