

HIGH FREQUENCY CHEST WALL OSCILLATION [HFCWO] MEDICARE COVERED ICD10 LIST

DX CODE	DIAGNOSIS DESCRIPTION
A15.0	Tuberculosis of Lung
B91	Sequelae of Poliomyelitis
E74.02	Pompe disease
E74.05	Lysosome-associated membrane protein 2 (LAMP2) deficiency
E84.0	Cystic Fibrosis with Pulmonary Manifestations
E84.9	Cystic Fibrosis, Unspecified
G12.0	Infantile Spinal Muscular Atrophy, Type 1 [Werdnig-Hoffman]
G12.1	Other Inherited Spinal Muscular Atrophy
G12.20	Motor Neuron Disease, Unspecified
G12.21	Amyotrophic Lateral Sclerosis
G12.22	Progressive Bulbar Palsy
G12.23	Primary Lateral Sclerosis
G12.24	Familial Motor Neuron Disease
G12.25	Progressive Spinal Muscle Atrophy
G12.29	Other Motor Neuron Disease
G12.8	Other Spinal Muscular Atrophies and Related Syndromes
G12.9	Spinal Muscular Atrophy, Unspecified
G14	Postpolio Syndrome
G35	Multiple Sclerosis
G70.00	Myasthenia Gravis without (acute) exacerbation
G70.01	Myasthenia Gravis with acute exacerbation
G71.00	Muscular Dystrophy, Unspecified
G71.01	Duchenne or Becker Muscular Dystrophy
G71.02	Facioscapulohumeral Muscular Dystrophy
G71.031	Autosomal Dominant Limb Girdle Muscular Dystrophy
G71.032	Autosomal Recessive Limb Girdle Muscular Dystrophy Due to Calpain-3 Dysfunction

DX CODE	DIAGNOSIS DESCRIPTION
G71.033	Limb Girdle Muscular Dystrophy Due to Dysferlin Dysfunction
G71.0340	Limb Girdle Muscular Dystrophy Due to Sarcoglycan Dysfunction, Unspecified
G71.0341	Limb Girdle Muscular Dystrophy Due to Alpha Sarcoglycan Dysfunction
G71.0342	Limb Girdle Muscular Dystrophy Due to Beta Sarcoglycan Dysfunction
G71.0349	Limb Girdle Muscular Dystrophy Due to Other Sarcoglycan Dysfunction
G71.035	Limb Girdle Muscular Dystrophy Due to Anoctamin-5 Dysfunction
G71.038	Other Limb Girdle Muscular Dystrophy
G71.039	Limb Girdle Muscular Dystrophy, Unspecified
G71.09	Other Specified Muscular Dystrophy
G71.11	Myotonic Muscular Dystrophy
G71.12	Myotonia Congenita
G71.13	Myotonic Chondrodystrophy
G71.14	Drug Induced Myotonia
G71.19	Other Specified Myotonic Disorders
G71.20	Congenital Myopathy, Unspecified
G71.21	Nemaline Myopathy
G71.220	X-linked Myotubular Myopathy
G71.228	Other Centronuclear Myopathy
G71.29	Other Congenital Myopathy
G71.3	Mitochondrial Myopathy, Not Elsewhere Classified
G71.8	Other Primary Disorders of Muscles
G72.0	Drug-induced Myopathy
G72.1	Alcoholic Myopathy
G72.2	Myopathy Due to Other Toxic Agents

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DX CODE	DIAGNOSIS DESCRIPTION
G72.41	Inclusion Body Myositis (IBM)
G72.49	Other inflammatory and Immune Myopathies, not elsewhere classified
G72.89	Other Specified Myopathies
G72.9	Myopathy, unspecified
G73.7	Myopathy in Diseases Classified Elsewhere
G80.0	Spastic Quadriplegic Cerebral Palsy
G82.50	Quadriplegia, Unspecified
G82.51	Quadriplegia, C1-C4 Complete
G82.52	Quadriplegia, C1-C4 Incomplete
G82.53	Quadriplegia, C5-C7 Complete
G82.54	Quadriplegia, C5-C7 Incomplete
J47.0	Bronchiectasis with Acute Lower Respiratory Infection
J47.1	Bronchiectasis with (Acute) Exacerbation

DX CODE	DIAGNOSIS DESCRIPTION
J47.9	Bronchiectasis, Uncomplicated
J98.6	Disorders of Diaphragm*
M33.02	Juvenile Dermatopolymyositis with Myopathy
M33.12	Other Dermatopolymyositis with Myopathy
M33.22	Polymyositis with Myopathy
M33.92	Dermatopolymyositis, Unspecified with Myopathy
M34.82	Systemic Sclerosis with Myopathy
M35.03	Sjogren Syndrome with Myopathy
Q33.4	Congenital Bronchiectasis

*CMS guidance states Disorders of the Diaphragm diagnosis must include documentation to support one of the following:

- Diaphragmatitis
- Paralysis of Diaphragm
- Relaxation of Diaphragm

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