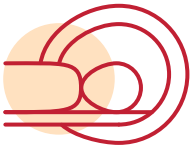
Medicare-Covered Diagnoses

ICD-10 Codes Covered for HFCWO

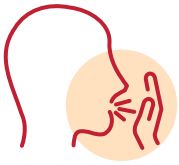
To qualify for reimbursement, a bronchiectasis patient must have documented:

1



HRCT scan confirming bronchiectasis diagnosis

2



Daily productive cough for at least six continuous months

OR

Frequent exacerbations requiring antibiotic therapy (i.e. three or more annually)

3

Failure of standard treatments such as OPEP or CPT to mobilize retained secretions

Bronchiectasis Diagnoses Covered by Medicare

Bronchiectasis

J47.0 Bronchiectasis with acute lower respiratory infection

J47.1 Bronchiectasis with (acute) exacerbation

J47.9 Bronchiectasis, uncomplicated

Q33.4 Congenital bronchiectasis

Other Diagnoses Covered by Medicare

Cystic Fibrosis

E84.0 Cystic fibrosis with pulmonary manifestations

E84.9 Cystic fibrosis, unspecified

Disorders of Diaphragm

J98.6 Disorders of diaphragm

Infectious / Immune

A15.0 Tuberculosis of lung

D84.1 Defects in the complement system

Late Effects of Poliomyelitis

B91 Sequelae of poliomyelitis

G14 Postpolio syndrome

continued on other side…

"Amazing for bronchiectasis, I've seen fewer pneumonias"

— Latecia, SmartVest user

smartvest®

AIRWAY CLEARANCE SYSTEM

Motor Neuron / Neuromuscular / Anterior Horn Cell Disease

G12.0	Infantile spinal muscular atrophy, type I [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

Multiple Sclerosis

G35	Multiple sclerosis

Muscular Dystrophy

G71.00	Muscular dystrophy, unspecified
G71.01	Duchenne or Becker muscular dystrophy
G71.02	Fascioscapulohumeral muscular dystrophy
G71.031	Autosomal dominant limb girdle muscular dystrophy
G71.032	Autosomal recessive limb girdle muscular dystrophy due to calpain-3 dysfunction
G71.09	Other specified muscular dystrophies

Limb Girdle Muscular Dystrophy

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

Myotonic and Metabolic Disorders

D81.810	Biotinidase deficiency
G71.11	Myotonic muscular dystrophy
G71.12	Myotonia congenita
G71.13	Myotonic chondrodystrophy
G71.14	Drug induced myotonia
G71.19	Other specified myotonic disorders

Other Myopathies

G71.2	Congenital myopathies
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
G72.89	Other specified myopathies
G73.7	Myopathy in diseases classified elsewhere
M33.02	Juvenile dermatomyositis with myopathy
M33.12	Other dermatomyositis with myopathy
M33.22	Polymyositis with myopathy
M33.92	Dermatopolymyositis, unspecified with myopathy
M34.82	Systemic sclerosis with myopathy
M35.03	Sicca syndrome with myopathy

Quadriplegia

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

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