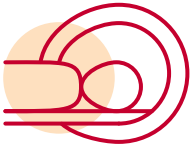


## Medicare-Covered Diagnoses

# 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

### Infectious / Immune

- A15.0 Tuberculosis of lung  
D84.1 Defects in the complement system

### Disorders of Diaphragm

- J98.6 Disorders of diaphragm

### Late Effects of Poliomyelitis

- B91 Sequelae of poliomyelitis  
G14 Postpolio syndrome

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

— Latecia, SmartVest user

*continued on other side...*

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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
- D81.82 Activated Phosphoinositide 3-kinase Delta Syndrome [APDS]
- E74.02 Pompe disease
- E74.05 Lysosome-associated membrane protein 2 [LAMP2] 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
- G72.41 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
- 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

# ELECTROMED, INC.

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500 Sixth Avenue NW Tel: 800.462.1045  
New Prague, MN 56071 Fax: 866.758.5077



info@electromed.com  
www.smartvest.com

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