Medicare requirements for bronchiectasis:

1. Required: CT scan confirming diagnosis of bronchiectasis.

   AND

2. Required: Daily productive cough for more than six continuous months.

   OR

   Frequent (i.e., more than two/year) exacerbations requiring antibiotic therapy in the last 12 months.

   AND

3. Required: Documentation (chart notes) of another treatment tried to mobilize secretions and clearly indicating the other technique or device has failed.

BRONCHIECTASIS ICD-10 CODE/DESCRIPTION

J47.0	Bronchiectasis with acute lower respiratory infection
J47.1	Bronchiectasis with acute exacerbation
J47.9	Bronchiectasis, uncomplicated
Q33.4	Congenital bronchiectasis

XXS	XS	S	M	L	XL	XXL	XXXL
18"–23" (46–58 cm)	23"–29" (58–74 cm)	29"–35" (74–89 cm)	35"–41" (89–104 cm)	41"–48" (104–122 cm)	48"–55" (122–140 cm)	55"–65" (140–165 cm)	65"–75" (165–190.5 cm)

Sizing determined by the larger of two circumferential measurements: nipple line and navel line.

AFFLOVEST®
by Tactile Medical

AffloVest®

MOBILE AIRWAY CLEARANCE THERAPY

Medicare approved
ICD-10 codes for AffloVest
HFCWO therapy
(HCPCS E0483)

Medicare requirements for cystic fibrosis and neuromuscular conditions:

Physician's order that includes: AffloVest prescription, qualifying diagnosis, chart notes to support the diagnosis, and well-documented failure of standard treatments to adequately mobilize retained secretions.

ICD-10 CODE/DESCRIPTION

Cystic Fibrosis

E84.0	Cystic fibrosis with pulmonary manifestations
E84.9	Cystic fibrosis, unspecified

Infectious/Immune

A15.0	Tuberculosis of lung

Polyiomyelitis

B91	Sequelae of poliomyelitis
G14	Postpolio syndrome

Myotonic/Metabolic Disorders

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

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.A	Relapsing remitting MS
G35.B0	Primary progressive MS
G35.B1	Active primary progressive MS
G35.B2	Non-active primary progressive MS
G35.C0	Secondary progressive MS
G35.C1	Active secondary progressive MS
G35.C2	Non-active secondary progressive MS
G35.D	Multiple sclerosis, unspecified

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
G71.033	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
G71.036	Limb girdle muscular dystrophy due to fukutin related protein dysfunction
G71.038	Other limb girdle muscular dystrophy
G71.039	Limb girdle muscular dystrophy, unspecified
G71.09	Other specified muscular dystrophies

Disorders of the Diaphragm

J98.6	Disorders of diaphragm

Myopathies

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
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

Myasthenia Gravis

G70.00	Myasthenia gravis without acute exacerbation
G70.01	Myasthenia gravis with acute exacerbation

Feel the difference.

For more information, please visit afflovest.com