



AFFLOVEST COVERAGE CRITERIA

The AffloVest requires a standard written order and an in-person, face-to-face examination that documents the need for the AffloVest.

STANDARD WRITTEN ORDER: A valid order must be signed prior to delivery of the equipment. The treating physician who conducted the face-to-face examination does not need to be the prescribing practitioner who writes the written order. However, the practitioner must have knowledge about what was covered during the face-to-face exam.

FACE-TO-FACE NOTES: The patient must have a face-to-face clinical evaluation by the treating physician within 6 months prior to the date of the order. The physician must document that the patient was evaluated and treated for a condition that supports the need for the AffloVest.

High frequency chest wall oscillation devices are covered for patient who meet Criteria 1, 2, or 3 and Criteria 4.

1. Patient has a diagnosis of cystic fibrosis
2. Patient has a diagnosis of bronchiectasis which has been confirmed by a high resolution, spiral, or standard CT scan which is characterized by:
 - a. Daily productive cough for at least 6 continuous months or
 - b. Frequent (more than 2 per year) exacerbations requiring antibiotic treatment
3. Patient has one of the following neuromuscular diseases diagnosis (see diagnosis codes that support medical necessity below): post-polio, acid maltase deficiency, anterior horn cell diseases, multiple sclerosis, quadriplegia, hereditary muscular dystrophy, myotonic disorders, other myopathies, paralysis of the diaphragm.
4. Patient must have tried and failed standard treatments for adequately mobilize retained secretions.

A15.0	Tuberculosis of lung	G12.8	Other spinal muscular atrophies & related syndromes	G71.8	Other primary disorders of muscles	J47.0	Bronchiectasis with acute lower respiratory infection
B91	Sequelae of poliomyelitis	G12.9	Spinal muscular atrophy, unspecified	G72.0	Drug-induced myopathy	J47.1	Bronchiectasis with (acute) exacerbation
D81.810	Biotinidase deficiency	G14	Postpolio syndrome	G72.1	Alcoholic myopathy	J47.9	Bronchiectasis, uncomplicated
D84.1	Defects in the complement system	G35	Multiple sclerosis	G72.2	Myopathy due to other toxic agents	J98.6	Disorders of diaphragm
E84.0	Cystic fibrosis with pulmonary manifestations	G71.0	Muscular dystrophy	G72.89	Other specific myopathies	M33.02	Juvenile dermatomyositis with myopathy
E84.9	Cystic fibrosis, unspecified	G71.11	Myotonic muscular dystrophy	G73.7	Myopathy in diseases classified elsewhere	M33.12	Other dermatomyositis with myopathy
G12.0	Infantile spinal muscular atrophy, type I	G71.12	Myotonic congenita	G82.50	Quadriplegia, unspecified	M33.22	Polymyositis with myopathy
G12.1	Other inherited spinal muscular atrophy	G71.13	Myotonic chondrodystrophy	G82.51	Quadriplegia, C1-C4 complete	M33.92	Dermatomyositis, unspecified with myopathy
G12.20	Motor neuron disease, unspecified	G71.14	Drug induced myotonia	G82.52	Quadriplegia, C1-C4 incomplete	M34.82	Systemic sclerosis with myopathy
G12.21	Amyotrophic lateral sclerosis	G71.19	Other specified myotonic disorders	G82.53	Quadriplegia, C5-C7 complete	M35.03	Sicca syndrome with myopathy
G12.22	Progressive bulbar palsy	G71.2	Congenital myopathies	G82.54	Quadriplegia, C5-C7 incomplete	Q33.4	Congenital bronchiectasis
G12.29	Other motor neuron disease	G71.3	Mitochondrial myopathy, not elsewhere classified				

E-PRESCRIBE: Frontier Home Medical makes ordering equipment easier with electronic ordering through Parachute. Visit our website (FrontierHomeMedical.com) to learn more about Parachute and get started with electronic ordering.

FAX PRESCRIPTION & FACE-TO-FACE NOTES TO THE LOCAL FRONTIER BRANCH.

If you have any questions or concerns, please contact Frontier Home Medical. Thank you for your cooperation!

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