



COUGH ASSIST COVERAGE CRITERIA

Coverage of cough assists should be reasonable and necessary. Documentation to support that should include:

- Face-to-Face Notes
- Standard Written Order

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.

- Patient has a neuromuscular disease
- Patient's condition is causing a significant impairment of chest and/or diaphragmatic movement, such that is results in an inability to clear retained secretions

STANDARD WRITTEN ORDER: A valid order must be obtained prior to delivery and include a qualifying diagnosis code in order to qualify for a cough assist.

B91	Sequelae of poliomyelitis	G70.01	Myasthenia gravis with (acute) exacerbation	G71.13	Myotonic chondrodystrophy	G72.9	Myopathy, unspecified
E74.02	Pompe disease	G71.0	Muscular dystrophy	G71.14	Drug induced myotonia	G73.7	Myopathy in diseases classified
E74.05	Lysosome-associated membrane protein 2[LAMP2] deficiency	G71.01	Duchenne or Becker muscular dystrophy	G71.19	Other specified myotonic disorder	G80.0	Spastic quadriplegic cerebral palsy
G12.0	Infantile spinal muscular atrophy, type I	G71.02	Facioscapulohumeral muscular dystrophy	G71.20	Congenital myopathy, unspecified	G82.50	Quadriplegia, unspecified
G12.1	Other inherited spinal muscular atrophy	G71.031	Autosomal dominant limb girdle muscular dystrophy	G71.21	Nemaline myopathy	G82.51	Quadriplegia, C1-C4 complete
G12.20	Motor neuron disease, unspecified	G71.032	Autosomal recessive limb girdle muscular dystrophy due to calpain-3 dysfunction	G71.220	X-linked myotubular myopathy	G82.52	Quadriplegia, C1-C4 incomplete
G12.21	Amyotrophic lateral sclerosis	G71.033	Limb girdle muscular dystrophy due to dysferlin dysfunction	G71.228	Other centronuclear myopathy	G82.53	Quadriplegia, C5-C7 complete
G12.22	Progressive bulbar palsy	G71.0340	Limb girdle muscular dystrophy due to sarcoglycan dysfunction, unspecified	G71.29	Other congenital myopathy	G82.54	Quadriplegia, C5-C7 incomplete
G12.23	Primary lateral sclerosis	G71.0341	Limb girdle muscular dystrophy due to alpha sarcoglycan dysfunction	G71.3	Mitochondrial myopathy, not elsewhere classified	J98.6	Disorders of diaphragm
G12.24	Familial motor neuron disease	G71.0342	Limb girdle muscular dystrophy due to beta sarcoglycan dysfunction	G71.8	Other primary disorders of muscles	M33.02	Juvenile dermatomyositis with myopathy
G12.25	Progressive spinal muscle atrophy	G71.0349	Limb girdle muscular dystrophy due to other sarcoglycan dysfunction	G72.0	Drug-induced myopathy	M33.12	Other dermatomyositis with myopathy
G12.29	Other motor neuron disease	G71.035	Limb girdle muscular dystrophy due to anoctamin-5 dysfunction	G72.1	Alcoholic myopathy	M33.22	Polymyositis with myopathy
G12.8	Other spinal muscular atrophies & related syndromes	G71.038	Other limb girdle muscular dystrophy	G72.2	Myopathy due to other toxic agents	M33.92	Dermatopolyomyositis, unspecified with myopathy
G12.9	Spinal muscular atrophy, unspecified	G71.039	Limb girdle muscular dystrophy, unspecified	G72.41	Inclusion body myositis [IBM]	M34.82	Systemic sclerosis with myopathy
G14	Postpolio syndrome	G71.09	Other specified muscular dystrophies	G72.49	Other inflammatory and immune myopathies, not elsewhere classified	M35.03	Sicca syndrome with myopathy
G35	Multiple sclerosis	G71.11	Myotonic muscular dystrophy	G72.89	Other specific myopathies		
G70.00	Myasthenia gravis without (acute) exacerbation	G71.12	Myotonia congenita				

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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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