



# **MSC Quick Assist**

Criteria and Code Guide in a Swipe



#### **Table of Contents**

| Affloyest Criteria ( | ^ard |
|----------------------|------|

| All way clearance Therapy Theu and Falleu      |  |
|--|--|
| Medicare approved ICD-10 Codes                 | 38   |
| Respiratory Assist Device (BIPAP) Criteria     | Card   |
| Restrictive Thoracic Disorder/Neuromuscular Di |  |
| Syndrome (OHS), and Chronic Obstructive Pulme  | onary Disease (COPD) · · · · · · · · · · · · · · · |
| Central Sleep Apnea, Complex (Treatment Emer   | gent Central) Sleep Apnea, and Prescription        |
| Requirements                                   |  |
|  |  |

Airway Claarance Thorany Tried and Failed

Non-Invasive Ventilation Criteria Card Restrictive Thoracic Disorder/Neuromuscular Disease and Obesity Hypoventilation

(Pickwickian) Syndrome (OHS)

**Helpful Links** 

Chronic Obstructive Pulmonary Disease (COPD) and Prescription Requirements · · · · · · · 

10





#### Airway Clearance Therapy Tried and Failed. This must be documented in the patients progress notes.

- 1. Airway clearance techniques **failed** by patient? Examples of methods:
  - CPT (manual or percussor)
  - Huff Coughing
  - Hypertonic Saline

- Oscillating PEP (Flutter, Acapella®, Aerobika®, Pep Valve, Pep Mask)
  - · Breathing Techniques
  - Suctioning

 Mucomyst\* (\*Notes must document the medication was prescribed for secretion mobilization)

- 2. Examples of reasons why therapy failed and/or is contraindicated/inappropriate for this patient:
- Cannot tolerate positioning/ hand CPT
  - Physical limitations of caregiver
  - Gastroesophageal reflux (GERD)
  - · Cognitive level

- · Too fragile for hand CPT Caregiver unable to perform adequate CPT
- Severe arthritis, osteoporosis
- Unable to form mouth seal

- Did not mobilize secretions.
- · Insufficient expiratory force
- Resistance to therapy
- Artificial airway
- **3.** Documentation for bronchiectasis patients should include the following:
  - CT scan

#### AND

Frequent (i.e. more than 2/year) exacerbations requiring antibiotic therapy

#### OR

Daily productive cough for at least 6 continuous months



#### Medicare approved ICD-10 Codes

Page 3

#### **ICD-10 Codes and Descriptions**

| J47.0     | Bronchiectasis with acute lower                                | G12.24  | Familial motor neuron disease                            |
|-----------|--|---------|--|
| J47.1     | respiratory infection Bronchiectasis with (acute) exacerbation | G12.25  | Progressive spinal muscle atrophy                        |
| J47.9     | Bronchiectasis, uncomplicated                                  | G12.29  | Other motor neuron disease                               |
| 033.4     | Congenital bronchiectasis                                      | G12.8   | Other spinal muscular atrophies and related syndromes    |
| A15.0     | Tuberculosis of lung   | G12.9   | Spinal muscular atrophy, unspecified                     |
| B91       | Sequelae of poliomyelitis                                      | G14     | Postpolio syndrome                                       |
| D81.810** | Biotinidase deficiency   | G35A*   | Relapsing-remitting multiple sclerosis                   |
| D84.1**   | Defects in the complement system                               | G35.B0* | Primary progressive multiple sclerosis, unspecified      |
| E74.02*   | Pompe Disease  | G35.B1* | Active primary progressive multiple sclerosis            |
| E74.05*   | Lysosome-associated membrane protein 2                         | G35.B2* | Non-active primary progressive multiple sclerosis        |
|           | [ĹAMP2] deficiency   | G35.C0* | Secondary progressive multiple sclerosis,                |
| E84.0     | Cystic fibrosis with pulmonary manifestations                  |         | unspecified  |
| E84.9     | Cystic fibrosis, unspecified                                   | G35.C1* | Active secondary progressive multiple sclerosis          |
| G12.0     | Infantile spinal muscular atrophy,<br>type I (Werdnig-Hoffman] | G35.C2* | Non-active secondary progressive multiple sclerosis      |
|           | type I (Werdnig-Hoffman)                                       | G35.D*  | Multiple sclerosis, unspecified                          |
| G12.1     | Other inherited spinal muscular atrophy                        | G70.00* | Myasthenia gravis without (acute) exacerbation           |
| G12.20    | Motor neuron disease, unspecified                              | G71.00  | Muscular dystrophy, unspecified                          |
| G12.21    | Amyotrophic lateral sclerosis                                  | G71.01  | Duchenne or Becker muscular dystrophy                    |
| G12.22    | Progressive bulbar palsy                                       | G71.02  | Facioscapulohumeral muscular dystrophy                   |
| G12.23    | Primary lateral sclerosis                                      |         | *Specific to Medicare Criteria **Not covered by Medicare |



#### Medicare approved ICD-10 Codes

\*\*Not covered by Medicare

Page 4

#### **ICD-10 Codes and Descriptions**

| 100-10   | codes and Descriptions                            |        |  |
|----------|---|--------|--|
| G71.031* | Autosomal dominant limb girdle muscular dystrophy | G72.41 | Inclusion body myositis  |
| G71.039* | Limb girdle muscular dystrophy, unspecified       | G72.49 | Other inflammatory and immune myopathies, not elsewhere classified |
| G71.09   | Other specified muscular dystrophies              | G72.89 | Other specified myopathies   |
| G71.11   | Mytonic muscular dystrophy                        | G72.9  | Myopathy, unspecified  |
| G71.12   | Myotonia congenita                                | G73.7  | Myopathy in diseases classified elsewhere                          |
| G71.13   | Myotonic chondrodystrophy                         | G80.0  | Spastic quadriplegic cerebral palsy                                |
| G71.14   | Drug induced myotonia                             | G82.50 | Quadriplegia, unspecified  |
| G71.19   | Other specified myotonic disorders                | G82.51 | Quadriplegia, C1-C4 complete                                       |
| G71.2    | Congenital myopathies                             | G82.52 | Quadriplegia, C1-C4 incomplete                                     |
| G71.20*  | Congenital myopathies, unspecified                | G82.53 | Quadriplegia, C5-C7 complete                                       |
| G71.21   | Nemaline myopathy                                 | G82.54 | Quadriplegia, C5-C7 incomplete                                     |
| G71.220  | X-linked myotubular myopathy                      | M33.02 | Juvenile dermatomyositis with myopathy                             |
| G71.228  | Other centronuclear myopathy                      | M33.12 | Other dermatomyositis with myopathy                                |
| G71.29   | Other congenital myopathy                         | M33.22 | Polymyositis with myopathy   |
| G71.3    | Mitochondrial myopathy, not elsewhere classified  | M33.92 | Dermatopolymyositis, unspecified with myopathy                     |
| G71.8    | Other primary disorders of muscles                | M34.82 | Systemic sclerosis with myopathy                                   |
| G72.0    | Drug-induced myopathy                             | M35.03 | Sicca syndrome with myopathy                                       |
| G72.1    | Alcoholic myopathy                                | J98.6  | Disorders of diaphragm   |
| G72.2    | Myopathy due to other toxic agents                |        |  |
|          |   |        | *Specific to Medicare Criteria                                     |



### Respiratory Assist Device (BIPAP) Criteria Card

Page 5

#### Restrictive Thoracic Disorder/ Neuromuscular Disease Qualifying Testing (at least one of the following):

- 1.PaCO2 ≥ 45 mmHg while awake and breathing prescribed FI02
- 2. Overnight pulse oximetry showing saturations < 88% for ≥ 5 minutes 3. Maximal inspiratory pressures < 60 cmH2O \*neuromuscular disease only
- 4. Forced vital capacity < 50% predicted \*neuromuscular disease only

#### Progress Note Documentation

- 1. Specific diagnosis (examples): · Amyotrophic Lateral Sclerosis
  - (ALŚ)
  - Kvphoscoliosis
  - Diaphragmatic paralysis Multiple sclerosis (MS)
  - Interstitial lung disease
  - Thoracic cage abnormality
- 2. Symptoms of respiratory failure (examples)
  - Daytime hypersomnolence Morning headache
  - Dyspnea
  - Cognitive dysfunction
- 3. COPD ruled out as the primary cause of respiratory failure

#### **Obesity Hypoventilation** (Pickwickian) Syndrome (OHS)

#### **Qualifying Testing**

- 1.PaCO2 ≥ 45 mmHg while awake and breathing prescribed FI02 AND
- 2. FEV1/FVC ratio ≥ 70% AND
- 3.2nd PaCO2, done during sleep or immediately upon awakening, breathing prescribed FI02 shows PaCO2 worsened > 7 mm Hr compared to original PaCO2. OR
- 4. Facility based PSG or HST demonstrates oxygen saturation ≤ 88% for > 5 minutes

#### **Progress Note Documentation**

- 1. Obstructive sleep apnea ruled out as the primary cause of hypercapnia (examples):
  - PSG/HST AHI < 5</li>
    - Increase in PaCO2 during sleep of > 10 mmHg
- Symptoms of respiratory failure (examples)
  - Daytime hypersomnolence
  - Morning headache
  - Dyspnea
  - Cognitive dysfunction

#### **Chronic Obstructive** Pulmonary Disease (COPD)

#### Qualifying Testing

Advantage Plans

- 1.ABG with PaCO2 ≥ 52 mmHg (on prescribed oxygen level) AND 2. \*Overnight oximetry showing oxygen saturations < 89% for ≥ 5 minutes \*Not required for Medicare FFS and Medicare
- **Progress Note Documentation** 
  - 1. Sleep appearuled out as the primary cause of hypercapnia
- 2. For hospital discharges patient must have used RAD or NIV within 24-hour period prior to discharge and is at risk of symptom exacerbation or PCO2 rise

Please Note: CMS has updated this policy, further guidance is expected late October 2025



### Respiratory Assist Device (BIPAP) Criteria Card

Page 6

#### **Central Sleep Apnea**

#### **Qualifying Testing**

- 1. PSG/HST AHI ≥ 5 AND
- 2. Central events are > 50% of total events AND
- Central AHI ≥ 5 AND
- 4. Significant improvement of signs/symptoms on prescribed settings during an in-lab titration studv

#### **Progress Note Documentation** (Prior to PSG/HST)

- 1. Signs symptoms of sleep disordéred breathing (examples)
  - Difficulty initialing or maintain sleep
  - Frequent awakenings
  - Non-restorative sleep
  - Snoring
  - Witnessed apneas
  - Excessive daytime sleepiness

#### Complex (Treatment Emergent Central) Sleep Apnea

### Qualifying Testing

- 1. PSG/HST AHI ≥ 5 On an in-lab titration study using CPAP or BIPAP without a backup respiratory rate, a pressure where:
  - 2. The obstructive AHI is < 5 AND
  - 3. Central events are > 50% of the total events AND
  - 4. The central AHI is > 5 AND
  - 5. Significant improvement of signs/symptoms on prescribed settings during an in-lab titration study

#### **Progress Note Documentation** (Prior to PSG/HST)

- 1. Signs symptoms of sleep disordéred breathing (examples)
  - Difficulty initialing or maintain
  - sleep
  - Frequent awakenings Non-restorative sleep
  - Snoring
  - Witnessed apneas
  - Excessive daytime sleepiness

#### **Prescription Requirements**

#### Spontaneous BIPAP

IPAP/EPAP | Oxygen bleed-in (if required) Supplies (heated humidifier, mask, filter, heated tubing)

#### Auto BIPAP

IPAP max | EPAP min | Pressure Support (PS) | Oxygen bleed-in (if required) Supplies (heated humidifier, mask, filter, heated tubing)

BIPAP ST (May not be initially ordered for OHS)

IPAP/EPAP | Backup respiratory rate | Oxygen bleed-in (if required) Supplies (heated humidifier, mask, filter,

heated tubing) BIPAP ST with iVAPS (May not be

initially ordered for OHS) EPAP | Pressure Support (PS) min | Pressure Support (PS) max | Backup

respiratory rate | Target tidal volume (Vt) or minute ventilation (Va) | Oxygen bleedin (if required) Supplies (heated humidifier, mask, filter, heated tubing)

**BIPAP Adaptive Servo Ventilation (ASV)** (Central or Complex Sleep Apnea only)

EPAP (set) OR EPAP min | EPAP max Pressure Support (PS) min | Pressure Support (PS) max | Oxygen bleed-in (if reauired)

Supplies (heated humidifier, mask, filter, heated tubing)

#### **Restrictive Thoracic** Disorder/Neuromuscular Disease

#### Qualifying Testing (at least one of the following):

- 1. PaCO2 > 45 mmHg
- 2. Maximal inspiratory pressures < 60 cmH2O 3. Forced vital capacity < 50% predicted

#### **Progress Note Documentation**

- 1. Specific diagnosis (examples):
  - Amyotrophic Lateral Sclerosis (ALS)
  - Kvphoscoliosis
  - Diaphragmatic paralysis
  - Multiple sclerosis (MS)
  - · Interstitial lung disease
  - Thoracic cage abnormality
- 2. COPD ruled out as the primary cause of respiratory failure
- 3. Hospitalizations (if any)
- 4. Reasons BIPAP has been considered and ruled out (examples):
  - Need for volume ventilation and backup respiratory rate
  - · Need for audible alarms
  - Expected use > 8 hours per day
     BIPAP tried and failed

#### Obesity Hypoventilation (Pickwickian) Syndrome

#### **Qualifying Testing**

1. PaCO2 > 45 mmHg 2. BMI ≥ 30 Kg/m 2

#### **Progress Note Documentation**

- 1. Obstructive sleep apnea ruled out as the primary cause of hypercapnia (examples):
  - PSG/HST AHI < 5</li>
- Increase in PaCO2 during sleep of > 10 mmHg
- 2. Reasons BIPAP has been considered and ruled out (examples):
  - BIPAP tried and failed due to nocturnal
    - desaturations or failure to correct hypercarbia Need for audible alarms Expected use > 8 hours per day
    - Need for pressures greater than those provided by BIPAP





## Chronic Obstructive Pulmonary Disease (COPD)

#### **Qualifying Testing**

1.ABG with PaCO2 ≥ 52 mmHg (on prescribed oxygen level)

#### **Progress Note Documentation**

- 1.Sleep apnea ruled out as the primary cause of hypercapnia
- 2. For hospital discharges patient must have used NIV within 24-hour period prior to discharge and is at risk of symptom exacerbation or PCO2 rise
- 3. One of the following:
  - Requires oxygen therapy at an FiO2 ≥36% or ≥ 4L nasally, OR
  - Requires ventilatory support for more than 8 hours per 24-hour period, OR
  - Requires the alarms and internal battery of a HMV, because the patient is unable to effectively breathe on their own for more than a few hours and the unrecognized interruption of ventilatory support is likely to cause a lifethreatening condition if the patient or caregiver
  - cannot be otherwise alerted as determined, OR
     Per the treating clinician, none of the below are likely to be achieved with consistent use of a RAD with backup rate feature for at least 4 hours per 24-hour period on at least 70% of days because the patient's needs exceed the capabilities of a RAD as justified by the patient's medical condition:

## Chronic Obstructive Pulmonary Disease (COPD) continued

- Normalization (< 46 mmHg) of PaCO2, OR</li>
- Stabilization of a rising PaCO2, OR
   20% reduction in PaCO2 from baseline value. OR
- Zow reduction in Pack 2 from dastelline Value, of Improvement of at least one of the patient symptoms associated with chronic hypercapnia Valid RX including supplies, hours of use, and settings

#### **Prescription Requirements**

- 1. Set tidal volume
- 2. Backup respiratory rate
- 3. EPAP minimum
- 4. FPAP maximum
- 5. Pressure Support (PS) minimum
- 6. Pressure Support (PS) maximum
- 7. Hours of use
- 8. Oxygen bleed-in (LPM) if required
- 9. Supplies: Mask Tubing Filters10. Ordering provider signature may be electronically signed
- 11. Date of signature may be electronically dated

Page 9

#### **Initial Setup**

#### **Dispensing Order Requirements**

- 1. Patient name
- 2. Order date
- 3. Specific brand of CGM/Accessories/Supplies
- 4. Frequency, if applicable, and quantity of supplies
- 5. Treating practitioner's name
- 6. Treating practitioner's signature 7. Treating practitioner's signature date

#### Progress Note Documentation

- 1. Diagnosis of diabetes mellitus AND
- 2. Within the prior 6 months, either an in-person or telehealth visit to evaluate diabetes control AND
- 3. One of the below:
  - a. Patient is on insulin OR
  - b. Patient has a history of problematic hypoglycemia with at least one of the following:
    - i. Patient a history of recurrent (more than one) level 2 hypoglycemic events documented by:
      - 1. Glucose value < 54 mg/dl (3.0 mmol/L) OR
      - 2. Classification of hypoglycemic events as level 2 OR
      - 3. A copy of the BGM testing log showing the specific qualifying events (glucose < 54 mg/dl (3.0 mmol/L)) AND
      - 4. Documentation of at least two previous medication adjustments and/or modification to the treatment plan prior to the most recent level 2 event
  - ii. Patient with at least one level 3 hypoglycemic event documented by:
    - Glucose value for the qualifying event (glucose < 54 mg/dl (3.0 mmol/L)) OR</li>
    - 2. Classification of the hypoglycemic event as level 3 OR
    - 3. A copy of the BGM log showing the specific qualifying event (glucose < 54 mg/dl (3.0 mmol/L)) AND 4. An indication in the medical record reflecting the patient required third party assistance for freatment

#### Resupply Requirements

Every six months following the initial prescription the patient must have an in-person or approved telehealth visit with the treating practitioner (pharmacists are not classified as treating practitioners)

#### **Progress Note Documentation** Adherence to the CGM regimen and diabetes treatment plan.

\*A new prescription is required every 12 months



#### Helpful Links Page 10

# Information Updated as of 9.25.25 <a href="https://www.cms.gov">www.cms.gov</a>

Please always refer to Centers for Medicare & Medicaid Services (CMS) for the most current information and guidelines.

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