



MEDICAL POLICY STATEMENT

Marketplace

Policy Name & Number	Date Effective
Positive Airway Pressure Devices for Pulmonary Disorders-MP-MM-1323	09/01/2023
Policy Type	
MEDICAL	

Medical Policy Statement prepared by CareSource and its affiliates are derived from literature based on and supported by clinical guidelines, nationally recognized utilization and technology assessment guidelines, other medical management industry standards, and published MCO clinical policy guidelines. Medically necessary services include, but are not limited to, those health care services or supplies that are proper and necessary for the diagnosis or treatment of disease, illness, or injury and without which the patient can be expected to suffer prolonged, increased or new morbidity, impairment of function, dysfunction of a body organ or part, or significant pain and discomfort. These services meet the standards of good medical practice in the local area, are the lowest cost alternative, and are not provided mainly for the convenience of the member or provider. Medically necessary services also include those services defined in any Evidence of Coverage documents, Medical Policy Statements, Provider Manuals, Member Handbooks, and/or other policies and procedures.

Medical Policy Statements prepared by CareSource and its affiliates do not ensure an authorization or payment of services. Please refer to the plan contract (often referred to as the Evidence of Coverage) for the service(s) referenced in the Medical Policy Statement. If there is a conflict between the Medical Policy Statement and the plan contract (i.e., Evidence of Coverage), then the plan contract (i.e., Evidence of Coverage) will be the controlling document used to make the determination. According to the rules of Mental Health Parity Addiction Equity Act (MHPAEA), coverage for the diagnosis and treatment of a behavioral health disorder will not be subject to any limitations that are less favorable than the limitations that apply to medical conditions as covered under this policy.

This policy applies to the following Marketplace(s):				
<input checked="" type="checkbox"/> Georgia	<input checked="" type="checkbox"/> Indiana	<input checked="" type="checkbox"/> Kentucky	<input checked="" type="checkbox"/> Ohio	<input checked="" type="checkbox"/> West Virginia

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A. Subject

Positive Airway Pressure Devices for Pulmonary Disorders

B. Background

Positive airway pressure (PAP) devices involve using a machine that includes a mask or other device that fits over the nose and/or mouth to provide positive pressure to keep breathing airways open. Continuous positive airway pressure, or CPAP, is used to treat sleep-related breathing disorders, including sleep apnea. It also may be used to treat preterm infants who have underdeveloped lungs. Bi-level or two-level positive airway pressure, or BiPAP, is used to treat lung disorders, such as chronic obstructive pulmonary disease (COPD). While CPAP delivers a single pressure, BiPAP delivers positive pressure both on inhalation and exhalation. PAP can provide better sleep quality, reduction or elimination of snoring, and less daytime sleepiness. The PAP machines should always be used according to the physician's order, as well as every time during sleep at home, while traveling, and during naps in order to produce the most effective outcome.

C. Definitions

- **Adherence** – Use of the device regularly as prescribed by the ordering physician for 4 or more hours per night for 70% of the nights during the most recent consecutive 30-day period during the first initial usage.
- **Bi-level Positive Airway Pressure (BiPAP) Device** – A device that uses mild bi-level or two levels of air pressure to keep breathing airways open.
- **Continuous Positive Airway Pressure (CPAP) Device** – A device that uses mild continuous air pressure to keep breathing airways open.
- **Positive Airway Pressure (PAP) Device** – A device that uses air pressure to keep breathing airways open, including both continuous positive airway pressure (CPAP) devices and bi-level positive airway pressure (BiPAP) devices.

D. Policy

I. PAP devices addressed in this policy include:

- A. E0601 – CPAP, continuous pressure capability, used with noninvasive nasal or face mask.
- B. E0470 – BiPAP, Bi-level pressure capability, without backup rate feature, used with noninvasive nasal or face mask.
- C. E0471 – BiPAP, Bi-level pressure capability, with backup rate feature, used with noninvasive nasal or face mask.
- D. E0472 – BiPAP, Bi-level pressure capability, with backup rate feature, used with invasive tracheostomy tube.

II. CareSource uses MCG Health clinical criteria to determine medical necessity.

A. PAP devices, CPAP and BiPAP:

1. During the first 3 months rental for a CPAP or BiPAP device, CareSource considers the device medically necessary when the MCG Health clinical criteria are met.

The MEDICAL Policy Statement detailed above has received due consideration as defined in the MEDICAL Policy Statement Policy and is approved.

2. For months 4-13 rental, CareSource considers the device medically necessary when MCG Health clinical criteria are met and submitted documentation confirms adherence.

Due to supply chain disruptions and a nationwide shortage of semiconductor chips for positive airway pressure (PAP) devices, providers may submit attestation to number of days of usage and need for continued use.

E. State-Specific Information

1. Georgia-NA
2. Indiana -NA
3. Kentucky-NA
4. Ohio-NA
5. West Virginia-NA

F. Conditions of Coverage

NA

G. Related Policies/Rules

NA

H. Review/Revision History

	DATE	ACTION
Date Issued	05/25/2022	New policy
Date Revised	06/07/2023	Annual review. Updated references. Approved at Committee.
Date Effective	09/01/2023	
Date Archived		

I. References

1. A-0431: Continuous Positive Airway Pressure (CPAP) Device. 27th ed. MCG Health; 2023. Retrieved June 01, 2023. www.careweb.careguidelines.com.
2. A-0994: Bi-level Positive Airway Pressure (BPAP) Device. 27th ed. MCG Health; 2023. Retrieved June 01, 2023. www.careweb.careguidelines.com.
3. American Academy of Sleep Medicine. Practice Guidelines. 2023. Retrieved June 01, 2023. www.aasm.org.
4. CMS. Local Coverage Determination for Positive Airway Pressure (PAP) Devices for the Treatment of Obstructive Sleep Apnea (L33718). 2021. Retrieved June 01, 2023. www.cms.gov.
5. CPAP. U.S. Department of Health & Human Services. National Heart, Lung and Blood Institute. March 24, 2022. Retrieved June 01, 2023. www.nhlbi.nih.gov.

This guideline contains custom content that has been modified from the standard care guidelines and has not been reviewed or approved by MCG Health, LLC.

The MEDICAL Policy Statement detailed above has received due consideration as defined in the MEDICAL Policy Statement Policy and is approved.