

<b>HEALTH PLAN POLICY</b>	
<b>Policy Title:</b> Oscillatory Devices for Airway Clearance	<b>Policy Number:</b> MUM36 <b>Revision:</b> B
<b>Department:</b> Medical Management	<b>Sub-Department:</b> Utilization Management
<b>Applies to Product Lines:</b> <input type="checkbox"/> Medicaid <input checked="" type="checkbox"/> USFHP <input type="checkbox"/> Children’s Health Insurance Plan <input checked="" type="checkbox"/> Commercial Insured <input checked="" type="checkbox"/> Health Insurance Exchange <input type="checkbox"/> Non Insured Business <input checked="" type="checkbox"/> Medicare Advantage	
<b>Origination/Effective Date:</b> 02/22/2018	
<b>Reviewed Date(s):</b>	<b>Revision Date(s):</b> 04/24/2019, 05/15/2020

**SCOPE:**

The focus of this policy is the use of oscillatory devices as opposed to manual chest physiotherapy for purposes of airway clearance of pulmonary secretions in selected disease states. There are two broad types of oscillatory devices – chest compression devices & intrapulmonary percussive ventilation devices; both will be discussed. Manual chest physiotherapy has long been the standard of care where it is possible at least 67% of the time. Where appropriate airway clearance devices are adjunctive to chest physiotherapy.

**DEFINITIONS AND ACRONYMS:**

- **Airway Clearance Therapy (ACT)**
- **American College of Chest Physicians (ACCP)**
- **Amyotrophic Lateral Sclerosis (ALS)**
- **Bronchiectasis** – Destruction of bronchial tissue as opposed bronchial irritation (bronchitis)
- **Chest Physiotherapy (CPT)**
- **Cystic Fibrosis Foundation (CFF)**
- **High Frequency Chest Compression (HFCC)**
- **Intrapulmonary Percussive Ventilation (IPV)**

**POLICY:**

- A. Medically necessary conditions for HFCC
  1. Cystic fibrosis – evidence is weak, but sufficient
  2. Bronchiectasis – evidence is weak, but sufficient
  3. Chronic neuromuscular disease such as ALS – evidence is weak, but sufficient
- B. No Medical Necessity for ACT
  4. COPD/Chronic Bronchitis – smoking related
  5. Upgrades to newer technology when current device is functional
  6. Contraindications such as chest trauma
  7. Any form of IPV- very limited data

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**REFERENCES:**

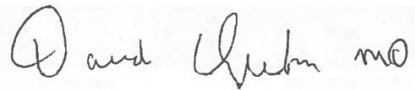
- ALS 2006 (Chaisson, et al) – 7: 107-111
- ACCP 2006 (McCool)– Clinical Practice Guidelines
- CFF 2009 – (Flume) – Metanalysis of 13 trials/7 reviews
- Respiratory Care 2012 -57: 221-228

**RELATED DOCUMENTS:**

None



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**REVISION HISTORY:**

<b>Revision</b>	<b>Date</b>	<b>Description of Change</b>	<b>Committee</b>
New	02/22/2018	Initial release.	Executive Leadership
A	04/24/2019	Annual review. Removed Medicaid and CHIP from lines of business.	Executive Leadership
B	05/15/2020	Annual review. No change to policy content.	Executive Leadership