

# SmartPA Criteria Proposal

<b>Drug/Drug Class:</b>	Pulmonary Arterial Hypertension (PAH) Agents: Endothelin Receptor Antagonists (ETRA) PDL Edit
<b>First Implementation Date:</b>	June 25, 2008
<b>Proposed Date:</b>	September 17, 2020
<b>Prepared For:</b>	MO HealthNet
<b>Prepared By:</b>	MO HealthNet/Conduent
<b>Criteria Status:</b>	<input type="checkbox"/> Existing Criteria <input checked="" type="checkbox"/> Revision of Existing Criteria <input type="checkbox"/> New Criteria

## Executive Summary

**Purpose:** The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Pulmonary arterial hypertension (PAH) is a severe, progressive, and often fatal condition that occurs idiopathically as well as in association with pulmonary, cardiac, and other thoracic conditions. Reliable estimates of the total prevalence of this condition are difficult to obtain because of the diversity of identifiable causes. PAH, formerly known as primary pulmonary hypertension, is characterized by elevations in pulmonary arterial pressure (PAP) to greater than 25 mmHg at rest and greater than 30 mmHg with exercise. The disease occurs when the PAP is abnormally elevated and forces the right side of the heart to progressively work harder when it pumps blood to the lungs. Symptoms include dyspnea, fatigue, chest pain, palpitations, syncope, and edema. Prognosis varies based on the severity of disease, whether right heart failure is present, and response to vasodilator therapy. If left untreated, the disease produces increases in PAP that may lead to right ventricular failure and death. Despite recent developments in the symptomatic treatment of PAH, there is still no cure.

PAH should be differentiated from pulmonary hypertension secondary to diseases of the heart and lung based on both pathology of the underlying disease and accepted treatments.

**Why Issue Selected:**

Total program savings for the PDL classes will be regularly reviewed.

**Program-Specific Information:**

Preferred Agents	Non-Preferred Agents
<ul style="list-style-type: none"> <li>Ambrisentan</li> <li>Tracleer®</li> </ul>	<ul style="list-style-type: none"> <li>Bosentan</li> <li>Letairis®</li> <li>Opsumit®</li> </ul>

**Type of Criteria:**

- |   |   |
|---|---|
| <input type="checkbox"/> Increased risk of ADE              | <input checked="" type="checkbox"/> Preferred Drug List |
| <input checked="" type="checkbox"/> Appropriate Indications | <input type="checkbox"/> Clinical Edit                  |

**Data Sources:**

- |  |   |
|--|---|
| <input type="checkbox"/> Only Administrative Databases | <input checked="" type="checkbox"/> Databases + Prescriber-Supplied |
|--|---|

## Setting & Population

- Drug class for review: Pulmonary Arterial Hypertension (PAH) Agents: Endothelin Receptor Antagonists (ETRAAs)
- Age range: All appropriate MO HealthNet participants

## Approval Criteria

- Documented diagnosis of pulmonary hypertension **AND**
- Failure to achieve desired therapeutic outcomes with trial on 2 or more preferred agents
  - Documented trial period for preferred agents **OR**
  - Documented ADE/ADR to preferred agents **OR**
- Documented compliance on current therapy regimen **OR**
- **For Opsumit: Clinical Consultant Review**

## Denial Criteria

- Participant is currently pregnant
- Lack of adequate trial on required preferred agents
- Therapy will be denied if all approval criteria are not met

## Required Documentation

Laboratory Results:

Progress Notes:

MedWatch Form:

Other:

## Disposition of Edit

Denial: Exception Code "0160" (Preferred Drug List)  
Rule Type: PDL

## Default Approval Period

1 year

## References

1. Evidence-Based Medicine and Fiscal Analysis: "Pulmonary Hypertension Agents, Oral – Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; July 2020.
2. Evidence-Based Medicine Analysis: "Primary Pulmonary Arterial Hypertension (PAH) Agents", UMKC-DIC; June 2020.
3. Lajoie A, Bonnet S, Provencher S. Review Article: Combination Therapy in Pulmonary Arterial Hypertension: Recent accomplishments and future challenges. *Pulmonary Circulation* 2017; 7(2) 312–325.
4. Lippincott, Williams, Wilkins. PDR Electronic Library, Montvale NJ; 2020.
5. USPDI, Micromedex; 2020.
6. Facts and Comparisons eAnswers (online); 2020 Clinical Drug Information, LLC.
7. Klinger JR, Elliott CG, Levine DJ, et al. Therapy for pulmonary arterial hypertension in adults: update of the CHEST Guideline and Expert Panel Report. *Chest*. 2016; 155(3): 565-586.
8. Tracleer [package insert]. South San Francisco, CA: Actelion Pharmaceuticals US Inc; 2019
9. Letairis [package insert]. Foster City, CA: Gilead; 2019
10. Opsumit [package insert]. South San Francisco, CA: Actelion Pharmaceuticals US Inc; 2019

*SmartPA PDL Proposal Form*

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