

## PRIOR AUTHORIZATION POLICY

- POLICY:** Pulmonary Arterial Hypertension – Orenitram Prior Authorization Policy
- Orenitram® (treprostinil extended-release tablets – United Therapeutics)

**REVIEW DATE:** 09/23/2020

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### OVERVIEW

Orenitram, prostacyclin mimetic, is indicated for the treatment of pulmonary arterial hypertension (PAH) World Health Organization (WHO) Group 1 to delay disease progression and to improve exercise capacity.<sup>1</sup>

### Disease Overview

PAH is a serious but rare condition impacting approximately fewer than 20,000 patients in the US. It is classified within Group 1 pulmonary hypertension among the five different groups that are recognized. In this progressive disorder the small arteries in the lungs become narrowed, restricted, or blocked causing the heart to work harder to pump blood, leading to activity impairment.<sup>2,3</sup> In time, right-sided heart failure and/or death may occur. Common PAH symptoms include shortness of breath, fatigue, chest pain, dizziness and fainting, along with impairment in activity tolerance. It is more prevalent in women. Patients of all ages may develop the disease; however, the mean age of diagnosis typically happens between 36 to 50 years. Children may also have PAH. The condition may occur due to various underlying medical conditions or as a disease that uniquely impacts the pulmonary circulation; both genetic and environmental factors may be involved. PAH is defined as a mean pulmonary artery pressure (mPAP)  $\geq 25$  mmHg with a pulmonary capillary wedge pressure (PCWP)  $\leq 15$  mmHg measured by cardiac catheterization. The prognosis in PAH has been described as poor, with the median survival being approximately 3 years. However, primarily due to advances in pharmacological therapies, the long-term prognosis has improved. Lung transplantation may be recommended if pharmacologic or medical therapies fail, based upon patient status. The WHO categorizes PAH into stages, which is also referred to as the functional class (Class I to IV) and is an adaptation of the New York Heart Association (NYHA) system to evaluate activity tolerance.

### Guidelines

In 2019, an updated CHEST guideline and Expert Panel Report regarding therapy for pulmonary arterial hypertension in adults was released.<sup>3</sup> Many other agents other than Orenitram are recommended as initial and subsequent therapy such as endothelin receptor antagonists (Letairis® [ambrisentan tablets], Tracleer® [bosentan tablets], Opsumit® [macitentan tablets]), phosphodiesterase type 5 [PDE 5] inhibitors [tadalafil, sildenafil], and Adempas® (riociguat tablets). The addition of an oral prostanoid product is recommended in patients with PAH who are in Functional Class III without evidence of rapid disease progression or a poor prognosis among those not willing or able to manage parenteral prostanoids.

### Safety

Abrupt discontinuation or sudden large reductions in the dosage of Orenitram may cause PAH symptoms to worsen.<sup>1</sup> In the event of a planned short-term treatment interruption for patients unable to take oral medication, consider a temporary infusion of subcutaneous or intravenous treprostinil.

## POLICY STATEMENT

Prior Authorization is recommended for prescription benefit coverage of Orenitram. All approvals are provided for the duration noted below. Because of the specialized skills required for evaluation and diagnosis of patients treated with Orenitram as well as the monitoring required for adverse events and long-term efficacy, approval requires Orenitram to be prescribed by or in consultation with a physician who specializes in the condition being treated.

**Documentation:** In the *Pulmonary Arterial Hypertension – Orenitram Prior Authorization Policy*, documentation is required for initiation of therapy where noted in the criteria as **[documentation required]**. Documentation may include, but is not limited to, chart notes and catheterization laboratory reports. For a patient case in which the documentation requirement of the right heart catheterization upon prior authorization coverage review for a different medication indicated for WHO Group 1 PAH has been previously provided, the documentation requirement in this *Pulmonary Arterial Hypertension – Orenitram Prior Authorization Policy* is considered to be met.

**Automation:** None.

## RECOMMENDED AUTHORIZATION CRITERIA

Coverage of Orenitram is recommended in those who meet the following criteria:

### FDA-Approved Indication

1. **Pulmonary Arterial Hypertension (World Health Organization [WHO] Group 1).** Approve for the duration noted if the patient meets ONE of the following (A or B):
  - A) **Initial Therapy.** Approve for 3 years if the patient meets all of the following criteria (i, ii, iii, and iv):
    - i. Patient has a diagnosis of World Health Organization (WHO) Group 1 pulmonary arterial hypertension (PAH); AND
    - ii. Patient meets the following criteria (a and b):
      - a) Patient has had a right heart catheterization **[documentation required]** (see documentation section above); AND
      - b) Results of the right heart catheterization confirm the diagnosis of WHO Group 1 PAH; AND
    - iii. Patient meets one of the following conditions (a or b):
      - a) Patient has tried two oral therapies for PAH (or is currently receiving them) from two of the three following different categories (either alone or in combination) each for  $\geq 60$  days: one phosphodiesterase type 5 (PDE5) inhibitor, one endothelin receptor antagonist (ERA), or Adempas (riociguat tablets); OR  
Note: Examples of PDE5 inhibitors include Revatio® (sildenafil tablets and suspension [generic]), Adcirca® (tadalafil tablets [generic]) and Alyq™ (tadalafil tablets) and examples of ERAs include Tracleer® (bosentan tablets), Letairis® (ambrisentan tablets [generic]), and Opsumit® (macitentan tablets).
      - b) Patient is receiving or has received in the past one PAH prostacyclin therapy or a prostacyclin receptor agonist (i.e., Uptravi® [selexipag tablets]) for PAH; AND  
Note: Examples of prostacyclin therapies for PAH include Tyvaso® (treprostinil inhalation solution), Ventavis® (iloprost inhalation solution), Remodulin® (treprostinil injection [generic]), and epoprostenol injection [Flolan, Veletri, generics]); AND
    - iv. Medication is prescribed by or in consultation with a cardiologist or a pulmonologist.

- B) Patient is Currently Receiving Orenitram.** Approve for 3 years if the patient meets all of the following criteria (i, ii, and iii):
- i.** Patient has a diagnosis of World Health Organization (WHO) Group 1 pulmonary arterial hypertension (PAH); AND
  - ii.** Patient meets the following criteria (a and b):
    - a)** Patient has had a right heart catheterization; AND
    - b)** Results of the right heart catheterization confirm the diagnosis of WHO Group 1 PAH); AND
  - iii.** The medication is prescribed by, or in consultation with, a cardiologist or a pulmonologist.

#### **CONDITIONS NOT RECOMMENDED FOR APPROVAL**

Coverage of Orenitram is not recommended in the following situations:

- 1.** Coverage is not recommended for circumstances not listed in the Recommended Authorization Criteria. Criteria will be updated as new published data are available.

#### **REFERENCES**

1. Orenitram<sup>®</sup> extended-release tablets [prescribing information]. Research Triangle Park, NC: United Therapeutics Corporation; October 2019.
2. McLaughlin VV, Archer SL, Badesch DB, et al; Writing committee members. ACCF/AHA 2009 Expert consensus document on pulmonary hypertension: A report of the American College of Cardiology Foundation Task Force on Expert Consensus Documents and the American Heart Association: Developed in collaboration with the American College of Chest Physicians, American Thoracic Society, Inc., and the Pulmonary Hypertension Association. *Circulation*. 2009;119:2250-2294.
3. 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*. 2019;155(3):565-586.