

# Pulmonary Arterial Hypertension

<b>Member and Medication Information</b>	
* indicates required field	
*Member ID:	*Member Name:
*DOB:	*Weight:
*Medication Name/Strength: <span style="float: right;"><input type="checkbox"/> Do Not Substitute. Authorizations will be processed for the preferred Generic/Brand equivalent unless specified.</span>	
*Directions for use:	
<b>Provider Information</b>	
* indicates required field	
*Requesting Provider Name:	*NPI:
*Address:	
*Contact Person:	*Phone #:
*Fax #:	Email:
<b>Medically Billed Information</b>	
* indicates required field for all medically billed products	
*Diagnosis Code:	*HCPCS Code:
*Dosing Frequency:	*HCPCS Units per dose:
Servicing Provider Name:	NPI:
Servicing Provider Address:	
Facility/Clinic Name:	NPI:
Facility/Clinic Address:	
Fax form and relevant documentation including: laboratory results, chart notes and/or updated provider letter to Pharmacy PA at <b>855-828-4992</b> , to prevent processing delays.	

**Select requested medication(s):**

**Preferred products are bold.** Non-Preferred Product Criteria also applies to (non-bolded) products.

Adempas (riociguat)    Adcirca (**tadalafil**)    Alyq (**tadalafil**)    Flolan (**epoprostenol**)    **Letairis** (ambrisentan)  
 Opsumit (macitentan)    Orenitram (treprostинil)    Remodulin (treprostинil)    Revatio (**sildenafil**)    **Tracleer** (bosentan)  
 Tyvaso (treprostинil)    Uptravi (selexipag)    Veletri (**epoprostenol**)    Ventavis (iloprost)    Other: \_\_\_\_\_

**Criteria for Approval: (All criteria must be met)**

Medication prescribed by, or in consultation with a pulmonologist or cardiologist.  
 Diagnosis of pulmonary arterial hypertension, confirmed in adults by right heart catheterization.  
 Indicate mean PAP: \_\_\_\_\_  
 Patient had vasoreactivity testing and failed maximum tolerated doses of calcium channel blockers.  
 Chart Note Page#: \_\_\_\_\_  
 Patient has WHO functional class of:  II    III    IV  
 Indicate all of the following medications that the patient has tried and failed:

Nitric Oxide-cGMP Enhancers	Endothelin Receptor Antagonists	Prostacyclin Pathway Agonists	
<input type="checkbox"/> Adcirca ( <b>tadalafil</b> )	<input type="checkbox"/> <b>Letairis</b> (ambrisentan)	<input type="checkbox"/> Flolan ( <b>epoprostenol</b> )	<input type="checkbox"/> Uptravi (selexipag)
<input type="checkbox"/> Adempas (riociguat)	<input type="checkbox"/> Opsumit (macitentan)	<input type="checkbox"/> Orenitram (treprostинil)	<input type="checkbox"/> Veletri (epoprostenol)
<input type="checkbox"/> Alyq ( <b>tadalafil</b> )	<input type="checkbox"/> <b>Tracleer</b> (bosentan)	<input type="checkbox"/> Remodulin (treprostинil)	<input type="checkbox"/> Ventavis (iloprost)
<input type="checkbox"/> Revatio ( <b>sildenafil</b> )		<input type="checkbox"/> Tyvaso (treprostинil)	

# UTAH MEDICAID PHARMACY PRIOR AUTHORIZATION REQUEST FORM

**Non-Preferred Product:** *(Criteria above must also be met)*

Trial and failure of preferred product, per Utah Medicaid's PDL, or prescriber must demonstrate medical necessity for non-preferred product. Details: \_\_\_\_\_ Chart Note Page #: \_\_\_\_\_

**NOTE:**

- ❖ Per federal regulation, Medicaid does not reimburse for drugs used for the treatment of sexual dysfunction or erectile dysfunction. Pharmacies should dispense only those products with pulmonary hypertension NDCs.

**Re-authorization Criteria:**

Updated letter with medical justification or updated chart notes demonstrating positive clinical response with six-minute walk test or FEV1.

**Authorization:**

28 days for titration dosing (up to three (3) months for Uptravi), or maintenance dosing = six (6) months

**Re-authorization:**

Up to six (6) months

**PROVIDER CERTIFICATION**

I hereby certify this treatment is indicated, necessary and meets the guidelines for use.

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Prescriber's Signature

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Date