

## Prior Authorization

## JOHNS HOPKINS HEALTH PLANS (MEDICAID) Opsumit - Priority Partners MCO

This fax machine is located in a secure location as required by HIPAA regulations. Complete/review information, sign and date. Fax signed forms to Johns Hopkins Health Plans at **1-410-424-4607**. Please contact Johns Hopkins Health Plans at **1-888-819-1043** with questions regarding the Prior Authorization process. When conditions are met, we will authorize the coverage of Opsumit - Priority Partners MCO.

Drug Name (select from list of drugs shown) Opsumit (macitentan)

Quantity	Frequency		Strength	
Route of Administration		Expected Length o	f Therapy	
Patient Information Patient Name:				
Patient ID: _			-	
Patient DOB:			-	
Patient Phone:				
Prescribing Physician Physician Name: Physician Phone: Physician Fax: Physician Address:			-	
City, State, Zip: _				

Diagnosis:

ICD Code:

Comments:

Please circle the appropriate answer for each question.				
1.	Has the plan authorized this medication in the past for this Y N patient (i.e., previous authorization is on file under this plan)?			
	NOTE: The use of physician samples, or manufacturer product discounts, does not guarantee coverage under the provisions of the medical and/or pharmacy benefit. All pertinent criteria must be met in order to be eligible for benefit coverage.			
	[If yes, skip to question 12.]			

2.	Does the patient have a definitive diagnosis of pulmonary Y N arterial hypertension?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
3.	Was the diagnosis of pulmonary arterial hypertension Y N confirmed by a cardiologist or pulmonologist by right heart catheterization?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
4.	Does the patient have World Health Organization (WHO) Y N Functional Class II or Class III symptoms?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
5.	Has the pulmonary arterial hypertension progressed  Y N    despite surgical treatment and/or maximal medical  Y N    treatment?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
6.	Has the patient had treatment failure with oral calcium Y N channel blockers?
	NOTE: Patients who have substantial reductions in pulmonary arterial pressure from short acting vasodilators at the time of catheterization may require high doses (e.g., nifedipine 240 milligrams per day (mg/day) or amlodipine 20 mg/day) \ NOTE: Submission of medical records is required.
	[If yes, skip to question 8.]
7.	Is the patient unable to take oral calcium channel blockers, Y N or is their use inappropriate in this patient?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
8.	Has the patient had a documented trial and inadequate Y N response to generic sildenafil or tadalafil?
	NOTE: Submission of medical records is required.
	[If yes, skip to question 10.]
9.	Does the patient have a documented contraindication to Y N generic sildenafil or tadalafil?
	NOTE: Submission of medical records is required.
	[If no, no further questions.]
10.	Has the patient had a documented trial and inadequate Y N response to Letairis AND Tracleer?
	NOTE: Submission of medical records is required.
	[If yes, skip to question 13.]
11.	Does the patient have a documented contraindication to Y N Letairis AND Tracleer?

NOTE: Submission of medical records is required.
[If yes, skip to question 13.]
[If no, no further questions.]
12. Is there clinical documentation showing the patient has experienced improved exercise capacity or a delay in symptom worsening?
NOTE: Submission of medical records is required.
[If no, no further questions.]
13. Will the requested drug be used in combination with one or Y N more drugs with the same pharmacology when the patient has not adequately responded to monotherapy?
NOTE: Submission of medical records is required.

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.

Prescriber (Or Authorized) Signature and Date