## Pharmacy Prior Authorization

## AETNA BETTER HEALTH ILLINOIS FAMILY HEALTH PLAN (MEDICAID)

PH Agents (Medicaid)

This fax machine is located in a secure location as required by HIPAA regulations. Complete/review information, sign and date. Fax signed forms to Aetna Better Health Illinois at **1-844-242-0908**. When conditions are met, we will authorize the coverage of PH Agents (Medicaid). Please note that all authorization requests will be reviewed as the AB rated generic (when available) unless states otherwise.

Drug Name			
Please specify			
Quantity		ngth	
Route of Administration	Expected Length of therapy		
Patient Information			
Patient Name			
Patient ID:			
Patient Group No ·			
Patient DOB:			
Patient Phone:			
Prescribing Physician			
Physician Name:			
Specialty:	NPI Number:		
Physician Fax:	Physician Phone:		
Physician Address:	City, State, Zip:		
Diagnosis:	ICD Code:		
Please circle the appropriate answe	r for each question.		
•	this medication in the past for this patient on is on file under this plan)?	Y	Ν
[If no, skip to question 3.]			
responding to therapy to	ab results support that the patient is maintain or achieve a low risk profile (e.g., lk distance, functional class, or reducing time	Y	Ν
[No further questions.]			
3. Is the requested drug bein pulmonologist or cardiolo	ng prescribed by or in consultation with a gist?	Y	Ν
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[If no, then no further questions.]

4.	Does the patient have a diagnosis of pulmonary hypertension with a mean pulmonary artery pressure (MPAP) of at least 25 mmHg at rest confirmed by right-heart catheterization (RHC)?	Y	Ν
	Submit test results or provide here:		
	[If no, then no further questions.]		
5.	Is the request for generic sildenafil, Revatio suspension, Adcirca, Letairis, Tracleer, Opsumit, generic epoprostenol, Remodulin, Orenitram, Uptravi or Adempas?	Y	Ν
	[If no, skip to question 7.]		
6.	Does the patient have World Health Organization (WHO) Class II, III, or IV symptoms (fatigue, dizziness, and fainting with normal physical activity or at rest)?	Y	Ν
	[If no, then no further questions.]		
	[If yes, skip to question 8.]		
7.	Does the patient have World Health Organization (WHO) Class III or IV symptoms (fatigue, dizziness, and fainting with less than normal physical activity or at rest)?	Y	Ν
	[If no, then no further questions.]		
8.	Does the patient have a diagnosis of Pulmonary Arterial Hypertension (PAH) WHO Group I?	Y	Ν
	[If no, skip to question 35.]		
9.	Does the patient meet ONE of the following: A) Inadequate response, intolerance or contraindication to, a calcium channel blocker, or B) Had a negative vasoreactivity test	Y	N
	Indicate which applies:		
	[If no, then no further questions.]		
10	Is the request for generic sildenafil tablets?	Y	Ν
	[If no, skip to question 12.]		

11. Does the patient have any of the following exclusions to therapy? A) Concurrent use of organic nitrates (ie, isosorbide mononitrate, isosorbide dinitrate, nitroglycerin), or B) Pulmonary veno-occlusive disease (PVOD)	Y	Ν
[No further questions.]		
12. Is the request for Adcirca?	Y	Ν
[If no, skip to question 15.]		
13. Has the patient had a documented trial and failure of, intolerance or contraindication to, sildenafil?	Y	Ν
[If no, then no further questions.]		
14. Does the patient have any of the following exclusions to therapy? A) Concurrent use of organic nitrates (ie, isosorbide mononitrate, isosorbide dinitrate, nitroglycerin), or B) Pulmonary veno-occlusive disease (PVOD)	Y	Ν
[No further questions.]		
15. Is the request for Tracleer or Letairis?	Y	Ν
[If no, skip to question 17.]		
16. Does the patient have any of the following exclusions to therapy? A) Pregnancy, or B) Pulmonary veno-occlusive disease (PVOD)	Y	Ν
[No further questions.]		
17. Is the request for generic epoprostenol?	Y	Ν
[If no, skip to question 19.]		
18. Does the patient have any of the following exclusions to therapy? A) Pulmonary veno-occlusive disease (PVOD), or B) Heart failure with left ventricular dysfunction	Y	Ν
[No further questions.]		
19. Is the request for Revatio suspension?	Y	Ν
[If no, skip to question 22.]		
20. Is there documentation to support the patient's difficulty or inability to swallow and the necessity of the brand suspension formulation?	Y	Ν

[If no, then no further questions.]

21. Does the patient have any of the following exclusions to therapy? A) Concurrent use of organic nitrates (ie, isosorbide mononitrate, isosorbide dinitrate, nitroglycerin), or B) Pulmonary veno-occlusive disease (PVOD)	Y	Ν
[No further questions.]		
22. Is the request for Opsumit?	Y	Ν
[If no, skip to question 25.]		
23. Has the patient had a documented trial and failure of, intolerance or contraindication to, 2 preferred oral agents: one PDE-5 inhibitor (e.g., sildenafil or Adcirca) and one endothelin receptor antagonist (e.g., Tracleer or Letairis)?	Y	Ν
[If no, then no further questions.]		
24. Does the patient have any of the following exclusions to therapy? A) Pregnancy, or B) Pulmonary veno-occlusive disease (PVOD)	Y	Ν
[No further questions.]		
25. Is the request for Adempas?	Y	Ν
[If no, skip to question 28.]		
26. Has the patient had a documented trial and failure of, intolerance or contraindication to, 2 preferred oral agents: one PDE-5 inhibitor (e.g., sildenafil or Adcirca) and one endothelin receptor antagonist (e.g., Tracleer or Letairis)?	Y	Ν
[If no, then no further questions.]		
27. Does the patient have any of the following exclusions to therapy? A) Concurrent use of organic nitrates (ie, isosorbide mononitrate, isosorbide dinitrate, nitroglycerin), or B) Pregnancy	Y	Ν
[No further questions.]		
28. Is the request for Uptravi?	Y	Ν
[If no, skip to question 30.]		
29. Has the patient had a documented trial and failure of, intolerance or contraindication to, 2 preferred oral agents: one PDE-5 inhibitor (e.g., sildenafil or Adcirca) and one endothelin receptor antagonist (e.g.,	Y	Ν

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Tracleer or Letairis)?

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[No further questions.]		
30. Is the request for Orenitram?	Y	Ν
[If no, skip to question 33.]		
31. Has the patient had a documented trial and failure of, intolerance or contraindication to, 2 preferred oral agents: one PDE-5 inhibitor (e.g., sildenafil or Adcirca) and one endothelin receptor antagonist (e.g., Tracleer or Letairis)?	Y	Ν
[If no, then no further questions.]		
32. Does the patient have Child Pugh Class C hepatic impairment?	Y	Ν
[No further questions.]		
33. Is the request for Tyvaso or Ventavis or Remodulin?	Y	Ν
[If no, then no further questions.]		
34. Has the patient had a documented trial and failure of, intolerance or contraindication to, 2 preferred oral agents: one PDE-5 inhibitor (e.g., sildenafil or Adcirca) and one endothelin receptor antagonist (e.g., Tracleer or Letairis)	Y	Ν
[No further questions.]		
35. Does the patient have WHO Group IV Pulmonary Hypertension (Chronic thromboembolic pulmonary hypertension, CTEPH)?	Y	Ν
[If no, then no further questions.]		
36. Is the request for Adempas?	Y	Ν
[If no, then no further questions.]		
37. Does patient meet ONE of the following: A) Has recurrent or persistent CTEPH after surgical treatment, or B) Patient is NOT a candidate for surgery.	Y	Ν
Provide reason:		
[If no, then no further questions ]		
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[If no, then no further questions.]

38. Does the patient have any of the following exclusions to therapy? A) Y N
Concurrent use of organic nitrates (ie, isosorbide mononitrate,
isosorbide dinitrate, nitroglycerin), or B) Pregnancy
[No further questions.]
 Comments:

I affirm that the information given on this form is true and accurate as of this date.

Prescriber (Or Authorized) Signature

Date