

MEDICARE FORM

Pulmonary Hypertension (Inhalation or Injectable Medication) Precertification Request

Virginia (HMO D-SNP) FAX: 1-833-280-5224 PHONE: 1-855-463-0933

For other lines of business:

Please use other form.

Page 1 of 2
(All fields must be completed and legible for precertification review.)

Please indicate:	Start of treatmen	·	of last treatment	,	,			
Precertification F		• •				Fax:		
A. PATIENT INFORMATION								
First Name:			Last Name:			DOB:		
Address:				City:		State:	ZIP:	
Home Phone:		Work Phone:		Cell Phone:		Email:	L	
Patient Current We	eight: lbs or	kgs Patie	nt Height: inche	s or cms Al	llergies:			
B. INSURANCE II		3	<u> </u>		3			
Aetna Member ID #:			Does patient have other coverage? ☐ Yes ☐ No					
Group #:			If yes, provide ID#: Carrier Name:					
Insured:			Insured:					
C. PRESCRIBER	INFORMATION							
First Name:			Last Name:		(Check C	ne): 🔲 M.D. 🗀] D.O. 🗌 N.P. 🗌 P.A.	
Address:				City:		State:	ZIP:	
Phone:	Fax:		St Lic #:	NPI #:	DEA #:		UPIN:	
Provider Email:		Offi	ice Contact Name:		Phone:			
D. DISPENSING	PROVIDER/ADMINIS	TRATION INFO	RMATION					
Center Na Home Infusion Agency N	red Physic Presion Center Presion Center President Presi	none:		Dispensing Prov Physician's (Specialty Ph Name: Address: Phone: TIN:	Office	tetail Pharmacy Other:		
Request is for: epoprostenol injection Flolan (epoprostenol injection) Remodulin (treprostinil injection) Revatio (sildenafil injection) Tyvaso (treprostinil inhalation solution) Veletri (epoprostenol injection) Ventavis (iloprost inhalation solution) Dose: Frequency: Implantable infusion pump External infusion pump IV SC F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.								
Primary ICD Code		e indicate prima	Other:	ly any other where a	ipplicable.			
•		ad clinical inform		ad in its entirety for a	all precertification	n reguests		
G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests. For All Requests (clinical documentation required): Please indicate the severity of the patient's symptoms using the World Health Organization (WHO) functional classification system: Select one:								
Please identify the type of pulmonary hypertension: Chronic thromboembolic pulmonary hypertension (CTEPH) Hereditary PAH due to activin receptor-like kinase type 1 (ALK1), endoglin, mothers against decapentaplegic 9 (SMAD9), caveolin-1 (CAV1), or potassium channel subfamily K member-3 (KCNK3) Hereditary PAH due to bone morphogenetic protein receptor type 2 (BMPR2) Hereditary PAH due to unknown causes Idiopathic PAH (formerly primary pulmonary hypertension) PAH due to diseases that localize to small pulmonary arterioles, including drug and toxin-induced (e.g., anorectic agents (diet drugs)) PAH associated with congenital heart disease PAH associated with connective tissue diseases PAH associated with HIV infection PAH associated with portal hypertension PAH associated with schistosomiasis Persistent pulmonary hypertension of the newborn (PPHN) (such as associated with congenital diaphragmatic hernia) Pulmonary hypertension associated with pulmonary veno-occlusive disease (PVOD) or pulmonary capillary hemangiomatosis (PCH) Sarcoidosis associated with pulmonary hypertension Other:								



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Page 2 of 2

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB						
G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.									
Yes No N/A Has the patient undergone an acute vasoreactivity test prior to initiation of therapy?									
Yes ☐ No Is an acute vasoreactivity test contraindicated due to right heart failure, low systemic blood pressure, low cardiac index, or presence of severe (functional class IV) symptoms?									
Please select: Low cardiac index Low systemic blood pressure Right heart failure Severe functional class IV symptoms									
→ ☐ Yes ☐ No Did the patient have a positive acute vasoreactivity test result (defined as a decrease in mPAP (mean									
pulmonary artery pressure) at least 10 mmHg to an absolute level of less than 40 mgHg without a decrease in cardiac output)?									
Yes No Does the patient have a documented trial and failure of a calcium channel blocker (dihydropyridine or diltiazem)?									
		the patient have a contraindicaright heart failure, hemodynam	ation to a calcium channel blocker nic instability)?						
For Initiation Requests (clinical documentation required):									
Revatio (sildenafil injection)									
☐ Yes ☐ No Is the patient concurrently on organic nitrates (e.g., isosorbide mononitrate, isosorbide dinitrate, nitroglycerin)?									
☐ Yes ☐ No Is the patient concurrently on guanylate cyclase (GC) stimulators (e.g., Adempas (riociguat))?									
For Continuation of Therapy Requests (clinical documentation required):									
☐ Yes ☐ No Is this continuation request a result of the patient receiving samples?									
☐ Yes ☐ N Is there clinical documentation indicating disease stability or improvement?									
Please select: Disease stability Disease improvement									
For Revatio (sildenafil injection) only:									
☐ Yes ☐ No Is the patient concurrently on organic nitrates (e.g., isosorbide mononitrate, isosorbide dinitrate, nitroglycerin)?									
☐ Yes ☐ No Is the patient concurrently on guanylate cyclase (GC) stimulators (e.g., Adempas (riociguat))?									
H. ACKNOWLEDGEMENT									
Request Completed By (Signature Required): Date:/									
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.									

The plan may request additional information or clarification, if needed, to evaluate requests.