

PRIOR AUTHORIZATION REQUEST FORM

EOC ID:

Adempas (riociguat)

Phone: 800-728-7947

Fax back to: 866-880-4532

The Scott & White Health Plan Pharmacy Department manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above. **Please note any information left blank or illegible may delay the review process.**

Patient Name:	Prescriber Name: Supervising Physician:	
Member/Subscriber Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Group Number:	NPI:	State Lic ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Primary Phone:	Specialty/facility name (if applicable):	

Drug Name and Strength:

Directions / SIG:

Please attach any pertinent medical history or information for this patient that may support approval. Please answer the following questions and sign.

Q1. What are the quantity and days sup period.	oly requested? Note: qu	uantity will be limited to a 14-day supply during titration	
Q2. For what diagnosis is this drug being	g prescribed (pick one)	?	
 Chronic thromboembolic pulmonary hypertension (CTEPH) WHO group 4 Pulmonary arterial hypertension (PAH) WHO Group 1 Other (please specify) 			
Q3. Please provide the ICD diagnosis code for the condition listed above.			
Q4. Is the patient a new start to therapy	?		
☐ Yes	No - please specify start date		
Q5. Specify the prescriber's specialty			
Cardiologist	Pulmonologist	Other	
Q6. If for PAH, was it confirmed by right heart catheterization?			



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☐ Yes	□ No	
 Q7. If for CTEPH, does the patient have the following? Please select all that apply: Recurrent or persistent CTEPH after pulmonary endartectomy (PEA) Inoperable CTEPH confirmed by perfusion scanning or pulmonary angiography Inoperable CTEPH confirmed by pretreatment right heart catheterization with mean pulmonary artery pressure (mPAP) > 25 mmHg, pulmonary capillary wedge pressure (PCWP) < 15 mmHg, and pulmonary vascular resistance (PVR) > 3 Wood units 		
Q8. Will the patient be taking any of the following concomitantly with Adempas? Please select all that apply: Nitrates or nitric oxide donors Specific or non-specific phosphodiesterase-5 (PDE5) inhibitors Theophylline derivatives None of the above 		
Q9. Is the patient a smoker?		
	□ No	
Q10. If the patient is female, is she currently enrolled in the Yes No Not applicable - patient is male	e Adempas REMS program?	
Q11. Please select all that apply regarding child-bearing po	otential:	
Pregnancy has been excluded prior to treatment initiation		
Pregnancy tests to exclude pregnancy will be conducted treatment discontinuation	cted monthly during treatment and for 1 month after	
 Patient will use effective forms of contraception to provide treatment discontinuation Patient is male 	revent pregnancy during treatment and for one month after	
Patient is female but does not have child-bearing po	tential	
Q12. Does the patient have failure of an adequate trial of, with the following? Please select all that apply:	contraindication, intolerance to, or persistence of symptoms	
Calcium channel blocker (if WHO Group 1 and posit	ive vasoreactivity test)	



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Letairis or Tracleer		
Other (please specify)		
Q13. If for continuation, please select all the following that apply:		
Patient is tolerating treatment		
Patient has evidence of continued disease stabilization or improvement		
Patient has continued medical need for Adempas		
Q14. Additional Comments:		

Prescriber Signature

Date

□ Expedited/Urgent - By checking this box and signing above, I certify that applying the standard review timeframe may seriously jeopardize the life or health of the enrollee or the enrollee's ability to regain maximum function

Lack of the necessary documentation may result in a medical necessity denial. Requesting providers may speak to a SWHP pharmacist or medical director at 1-800-728-7947 regarding the case to have an opportunity to help impact the decision on a request before coverage has been decided.

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