

## Prior Authorization

## JOHNS HOPKINS HEALTH PLANS (MEDICAID)

Rukobia - 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 Rukobia - Priority Partners MCO.

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Drug Name (select from list of drugs shown) Rukobia (fostemsavir)					
Quantity	Frequency	Strength			
Route of Administration	Expected Length	of Therapy			
Patient Information					
Patient Name:		-			
Patient Group No.:		-			
Patient DOB:		-			
Patient Phone:		_			
Prescribing Physician					
Physician Name:		_			
Physician Phone:		_			
Physician Fax: Physician Address:		-			
City, State, Zip:		_			
<u> </u>					
Diagnosis:	ICD Code:				
Comments:					
Please circle the appropriate at 1. Does the patient have	-				
<ol> <li>Does the patient have a human immunodeficiency virus-1 Y N (HIV-1) infection?</li> </ol>					
NOTE: Submission of medical records is required.					
[If no, no further que	estions.]				
2. Is the patient 18 years	of age or older?	Y N			
[If no, no further que	[If no, no further questions.]				
3. 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 11.]		
4.	Does the patient have significant antiretroviral treatment experience with documented historical or baseline resistance, intolerability, and/or contraindications to antiretrovirals in at least three classes?	Y N	]
	NOTE: Submission of medical records is required.		
	[If no, no further questions.]		
5.	Has the patient had an inadequate response to current antiretroviral regimens evidenced by HIV RNA viral load greater than or equal to 400 copies per milliliter?	ΥN	]
	NOTE: Submission of medical records is required.		
	[If no, no further questions.]		
6.	Does the patient have at least one responsive antiretroviral (but no more than two antiretrovirals) that can be used concurrently with the requested drug to create an effective treatment regimen?	Y N	]
	NOTE: Submission of medical records is required.		
	[If no, no further questions.]		
7.	Is or has the prescriber consulted with an infectious disease specialist, or a certified HIV provider?	ΥN	]
	[If no, no further questions.]		
8.	Is the patient using strong cytochrome P450 3A inducers (carbamazepine, phenytoin, rifampin, enzalutamide, mitotane, St. John's wort, etc.)?	YN	]
	[If yes, no further questions.]		
9.	Does the patient have a history of hypersensitivity reactions to fostemsavir or any other component of the requested drug?	ΥN	]
	[If yes, no further questions.]		
10.	Is the patient breastfeeding?	ΥN	
	[No further questions.]		
11.	Is the patient using strong cytochrome P450 3A inducers (carbamazepine, phenytoin, rifampin, enzalutamide, mitotane, St. John's wort, etc.)?	ΥN	
	[If yes, no further questions.]		
12.	Does the patient have a history of hypersensitivity reactions to fostemsavir or any other component of the requested drug?	Y N	]
	[If yes, no further questions.]		
13.	Is the patient breastfeeding?	ΥN	]
	[If yes, no further questions.]		

14. Is there documentation showing beneficial response to	Y N
treatment?	
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