UTAH MEDICAID PHARMACY PRIOR AUTHORIZATION REQUEST FORM

Yeztugo (lenacapavir)

Member and Medication Information			
	* indicates required field		
*Member ID:	*Member Name:		
*DOB:	*Weight:		
*Medication Name/ Strength:	<u> </u>		
Do Not Substitute. Authorizations w	ill be processed for the preferred Generic/Brand equivalent unless specified.		
*Directions for use:			
	Provider Information		
	* indicates required field		
*Requesting Provider Name:	*Requesting Prescriber NPI:		
Address:			
*Contact Person:	*Office Phone:		
*Office Fax:	*Office Email:		
	Medically Billed Information tes required field for all medically billed products		
*Diagnosis Code:	*HCPCS Code:		
*Dosing Frequency:	*HCPCS Units per Dose:		
Servicing Provider Name:	NPI:		
Servicing Provider Address:	NI I.		
Facility/Clinic Name:	NPI:		
•	INPI.		
Facility/Clinic Address:			
	ation including: laboratory results, chart notes and/or updated acy PA at 855-828-4992 , to prevent processing delays.		
'	ation, see Sunlenca (lenacapavir) PA form		
Criteria for Approval: (All of the following	g criteria must be met)		
1. Does the patient weigh at least 35kg? □ Yes □ No			
2. Is the patient being prescribed Yes	ztugo for pre-exposure prophylaxis (PrEP) to reduce the risk of HIV-1?		
	□ Yes □ No		
	d by or in consultation with an HIV specialist or a provider specializing in		
the treatment of infectious diseas			
4. Has the patient had a confirmed r	negative HIV-1 test within two weeks prior to treatment initiation?		
5. Has the patient tried and failed a	oreferred oral PrEP regimen AND Apretude (cabotegravir), or has the		
provider given rationale for the la	•		
· -			
	thereof:		
	of Apretude (cabotegravir):		
	thereof:		
	managing planned and unplanned missed doses per the prescribing		
information? Chart note page#:	□ Yes □ No		

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7.	Does the provider attest that the patient agrees to the required testing and every 6-month in schedule, and that the provider has counseled the patient about the importance of adhering Yeztugo dosing visits to help reduce the risk of acquiring HIV-1 infection and development or	g to scheduled
8.	Does the provider attest to follow supplemental dosing recommendations per the prescribir	
	the patient is initiated on strong or moderate CYP3A4 inducers?	□ Yes □ No
Reaut	horization Criteria:	
1.	Has the provider submitted an updated letter with medical justification or updated chart no	tes
	demonstrating the need for PREP treatment?	☐ Yes ☐ No
2.	Has the provider submitted a confirmed negative HIV-1 test taken within the past 2 weeks?	
		□ Yes □ No
Initial	Authorization: Up to six (6) months	
Reaut	chorization: Up to one (1) year	
Note:		
*	Risk of drug resistance with use of lenacapavir for HIV-1 preexposure prophylaxis in undiagrinfection	nosed HIV-1

PROVIDER CERTIFICATION			
I hereby certify this treatment is indicated, necessary and meets the guidelines for use.			
Prescriber's Signature	 Date		