

# HEALTHY U CHIP

## PRIOR AUTHORIZATION REQUEST FORM

### PREVYMIS™

**For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U CHIP Prior Authorization Department.**

- For **Medical Pharmacy** please fax requests to: 801-213-1547
- For **Retail Pharmacy** please fax requests to: 385-425-4052

**Failure to submit clinical documentation to support this request will result in a dismissal of the request.**

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 385-425-5094

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:

Height/Weight:

**Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.**

**Product being requested:**  Previmis™ (letermovir)

Dosing/Frequency: \_\_\_\_\_

**If the request is for reauthorization, proceed to reauthorization section.**

Questions	Yes	No	Comments/Notes
<b>Prophylaxis of CMV infection and disease in allogeneic hematopoietic stem cell transplant recipients</b>			
1. Is the requesting provider an infectious disease specialist, hematologist, oncologist, or transplant specialist?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does documentation show the member is cytomegalovirus (CMV)-seropositive [R+]?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
3. Is the member an allogeneic hematopoietic stem cell transplant recipient?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
4. Is the therapy initiated between day 0 and day 28 post-transplant?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
5. Does the member have severe (Child-Pugh C) hepatic impairment?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
<b>Prophylaxis of CMV infection and disease in kidney transplant recipients at high risk</b>			
1. Is the requesting provider an infectious disease specialist, hematologist, oncologist, or transplant specialist?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does documentation show the donor is cytomegalovirus (CMV) seropositive [D+]?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
3. Does documentation show the member (recipient) is CMV seronegative [R-]?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
4. Is the member a kidney transplant recipient?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>

5. Does the patient have valganciclovir and ganciclovir?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
6. Is the therapy initiated between day 0 and day 7 post-transplant?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
7. Does the member have severe (Child-Pugh C) hepatic impairment?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
<b>INJECTABLE PREVYMIS™</b>			
1. Is the member unable to swallow or has severe dysphagia preventing the use of solid oral medication?	<input type="checkbox"/>	<input type="checkbox"/>	<b>Please provide documentation</b>
<b>What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.</b>			
Additional information:			
Physician Signature:			

**\*\* Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\***

Policy: PHARM-CHIP-100  
 Origination Date: 07/01/2024  
 Reviewed/Revised Date:  
 Next Review Date:  
 Current Effective Date: 07/01/2024

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