SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-844-305-2331</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization can be delayed</u>.

<u>Drug Requested</u>: Prevymis[®] (letermovir) Injection for IV Infusion (J3490/C9399) (Medical)

MEMBER & PRESCRIBER	INFORMATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	
DEA OR NPI #:	
DRUG INFORMATION: Aut	chorization may be delayed if incomplete.
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight:	Date:
☐ Standard Review. In checking this	s box, the timeframe does not jeopardize the life or health of the member naximum function and would not subject the member to severe pain.

Quantity Limit (max daily dose) [NDC/HCPCS Unit]:

- 480 mg/24 mL solution per 1 vial; NDC 00006-5004-02; 1 billable unit per day
- 240 mg/12 mL solution per 1 vial; NDC 00006-5003-02; 1 billable unit per day

Recommended Dose: 480 mg IV once daily

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CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

o I	Diagnosis: Cytomegalovirus, prophylaxis in hematopoietic cell transplant recipients
	Recommended Dose: 480 mg orally once daily. Initiate therapy between Day 0 and Day 28 post ransplantation (before or after engraftment), and continue through Day 200 post-transplantation
Ī	Length of Authorization: 200 days of therapy
	Member is ≥ 18 years of age
	Member will be receiving Prevymis® for the prophylaxis of cytomegalovirus (CMV) disease
	Member is a CMV-seropositive recipient [R+] of an allogeneic hematopoietic stem cell transplant (HSCT
	Medication will be initiated between day 0 and day 28, before or after engraftment
	Enter date transplant was performed:
	Member is NOT receiving the requested medication beyond 200 days post-transplantation
	Provider must submit chart notes to document contraindication to therapy with oral Prevymis® tablets and rationale for medical necessity to continue IV Prevymis® therapy
o I	Diagnosis: Cytomegalovirus, prophylaxis in kidney transplant recipients
	Recommended Dose: 480 mg orally once daily. Initiate therapy between Day 0 and Day 7 post ransplantation (before or after engraftment), and continue through Day 200 post-transplantation
I	Length of Authorization: 200 days of therapy
	Member is ≥ 18 years of age
	Member will be receiving a kidney transplant
	Member will be receiving Prevymis® for the prophylaxis of cytomegalovirus (CMV) disease
	Member is at high-risk for CMV disease [documentation recording kidney donor is CMV-seropositive, and the recipient (member) is CMV-seronegative (D+/R-)]
	Medication will be initiated between day 0 and day 7, before or after engraftment
	Enter date transplant was performed:
	Member is NOT receiving the requested medication beyond 200 days post-transplantation
	Provider must submit chart notes to document contraindication to therapy with oral Prevymis® tablets and rationale for medical necessity to continue IV Prevymis® therapy

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Medication being provided by: Please check applicable box below.		
	Location/site of drug administration:	
	NPI or DEA # of administering location:	
	<u>OR</u>	
	Specialty Pharmacy – Proprium Rx	

For urgent reviews: Practitioner should call Sentara Health Pre-Authorization Department if they believe a standard review would subject the member to adverse health consequences. Sentara Health's definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member's ability to regain maximum function.

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *

REVISED/UPDATED: 6/21/2018; (Reformatted) 3/19/2019; 7/8/2019; 9/24/2019; 10/17/2023