AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-305-671-0200</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

<u>Drug Requested</u>: Prevymis[®] (letermovir) tablets (Pharmacy)

MEMBER & PRESCRIBER INFO	RMATION: Authorization may be delayed if incomplete.
Member Name:	
Member AvMed #:	
Prescriber Name:	
Prescriber Signature:	
Office Contact Name:	
	Fax Number:
NPI #:	
DRUG INFORMATION: Authorizati	on may be delayed if incomplete.
Drug Name/Form/Strength:	
Dosing Schedule:	
Diagnosis:	ICD Code, if applicable:
Weight (if applicable):	Date weight obtained:
Quantity Limit:	

- 480 mg tablets 1 tablet per day
- 240 mg tablets 1 tablet per day
- 120 mg oral pellets 2 packets per day
- 20 mg oral pellets 4 packets per day

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.

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		Diagnosis:	Cytomeg	galovirus,	prophyla	xis in hem	atopoietic co	ell transplant	t recipients
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Initiate therapy between Day 0 and Day 28 post-HSCT (before or after engraftment) and continue through Day 100 post-HSCT. In patients at risk for late CMV infection and disease, Prevymis® may be continued through Day 200 post-HSCT.

Recommended Dosage:

• Adult and Pediatric Patients 12 Years of Age and Older and Weighing at least 30 kg: 480 mg administered orally once daily

Recommended Dosage:

• Pediatric Patients 6 Months to Less than 12 Years of Age or 12 Years of Age and Older and Weighing Less than 30 kg:

Body Weight	Daily Oral Dose	Tablets	Oral Pellets
15 kg to less than 30 kg	240 mg	One 240 mg tablet	Two 120 mg packets
7.5 kg to less than 15 kg	120 mg	Not Recommended	One 120 mg packet
6 kg to less than 7.5 kg	80 mg	Not Recommended	Four 20 mg packets

Length of Authorization: 200 days of therapy

Member is 6 months of age or older and weighs at least 6 kg
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

□ Diagnosis: Cytomegalovirus, prophylaxis in kidney transplant recipients

Initiate therapy between Day 0 and Day 7 post-transplant and continue through Day 200 post-transplant.

Recommended Dosage:

• Adult and Pediatric Patients 12 Years of Age and Older and Weighing at least 40 kg: 480 mg administered orally once daily

Length of Authorization: 200 days of therapy

- ☐ Member is 12 years of age or older and weighs at least 40 kg
- ☐ Member will be receiving a kidney transplant
- ☐ Member will be receiving Prevymis® for the prophylaxis of cytomegalovirus (CMV) disease

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Medication being provided by Specialty Pharmacy – Proprium Rx		
	Member is <u>NOT</u> receiving the medication beyond 200 days post-transplantation	
	Enter date transplant was performed:	
	Medication will be initiated between day 0 and day 7, before or after engraftment	
	Member is at high-risk for CMV disease [documentation recording kidney donor is CMV-seropositive, and the recipient (member) is CMV-seronegative (D+/R-)]	

^{**}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.*