

ANTIVIRALS, CMV PRIOR AUTHORIZATION FORM *(form effective 1/8/2024)*

Prior authorization guidelines for **Antivirals, CMV** and **Quantity Limits/Daily Dose Limits** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	# of pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:		NPI:	State license #:	
LTC facility contact/phone:			Street address:	
Beneficiary name:			City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:	
Beneficiary address:			Beneficiary phone number:	

CLINICAL INFORMATION

Drug requested:	Strength:	Dosage form:	
Directions:	Quantity:	Refills:	
Diagnosis (<i>submit documentation</i>):		Diagnosis code (<i>required</i>):	
Is the requested medication being prescribed by or in consultation with a hematologist/oncologist, infectious disease specialist, or transplant specialist?		<input type="checkbox"/> Yes <input type="checkbox"/> No <i>Submit documentation of consultation.</i>	

**Complete all sections that apply to the beneficiary and this request.
 Check all that apply and submit documentation for each item.**

1. For Livtency (maribavir):

The beneficiary is/was taking ganciclovir or valganciclovir AND:

Ganciclovir/valganciclovir will be/was discontinued before starting Livtency (maribavir)

Is being treated for post-transplant CMV infection/disease AND:

Is continuing treatment with Livtency (maribavir) upon inpatient discharge

Tried and failed or has a reason not to try at least one of the following:

cidofovir foscarnet ganciclovir valganciclovir

Has culture and sensitivity results showing that only Livtency (maribavir) will be effective

Is receiving concomitant therapy with carbamazepine OR phenobarbital AND:

The dose of Livtency (maribavir) was adjusted according to FDA-approved package labeling

2. For Prevmis (letermovir):

- Is using Prevmis (letermovir) for post-transplant CMV prophylaxis AND:
 - Is CMV-seropositive
 - Is at high risk for CMV reactivation (eg, cord blood transplant, CMV-seropositive donor)
- Is NOT receiving concomitant therapy with a contraindicated drug/drug combination (eg, ergot alkaloids, pimozide, pitavastatin with cyclosporine, simvastatin with cyclosporine)

- Is or will be receiving concomitant therapy with cyclosporine AND:
 - The dose of Prevmis (letermovir) was adjusted according to FDA-approved package labeling
- Initiated or will initiate treatment with Prevmis (letermovir) in the post-transplant period in accordance with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature
- Is continuing treatment with Prevmis (letermovir) upon inpatient discharge

3. For all other NON-PREFERRED Antivirals, CMV:

- Has a history of trial and failure of or a contraindication or an intolerance to the preferred Antivirals, CMV approved or medically accepted for the beneficiary's diagnosis or condition (*Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.*)
- Has culture and sensitivity results showing BOTH of the following:
 - The beneficiary's infection is NOT susceptible to the preferred Antivirals, CMV
 - The beneficiary's infection IS susceptible to the requested non-preferred Antivirals, CMV

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

Prescriber Signature:

Date:

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